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Hydrogenated Si₁₂Au₂₀ cluster as a molecular sensor with high performance for NH₃ and NO detection: A first-principle study

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abstract

Using density functional theory (DFT) calculations, we investigated the adsorption of NH₃, NO, CH₄, CO₂, H₂, N₂, H₂O, and O₂ molecules on the (hydrogenated) Si₁₂Au₂₀ cluster with the aim of finding a promising molecular sensor for NH₃ and NO detection. The Si₁₂Au₂₀ cluster could be a disposable molecular sensor for NH₃ and NO because of its long recovery time (over 3 h). To improve the recovery time, we considered the hydrogenation of Si atoms in the Si₁₂Au₂₀ cluster to reduce the strength of the adsorption of NH₃ and NO. The vibrational frequency analysis, molecular dynamics simulations and electronic properties show that the H₁₂Si₁₂Au₂₀ (HSA) structure is highly stable. NH₃ and NO are chemisorbed on HSA with moderate adsorption energies and evident charge transfer, while the other molecules are all physisorbed on HSA. Our results show that the electrical conductivities of HSA will change dramatically due to the adsorption of NH₃ and NO molecules. The recovery time of HSA is predicted to be very short at 300 K. To have a comprehensive comparison with the above results, we also considered different coverages of NO or NH₃ molecules adsorbed on HSA, and the adsorption of NO and NH₃ on the Au₃₂ and Si₃₂ clusters. We found that Au₃₂ and Si₃₂ clusters are not suitable for NO and NH₃ detection. We predict that HSA should be a promising gas sensor with high performance for NH₃ and NO detection at low coverage for future experimental validation.

1. Introduction

Intense research has been dedicated to the development of gas sensors for detecting and monitoring ultralow concentrations of toxic and harmful gases, which are of concern for safety and sustainable development. Nitrogen-based gases, such as ammonia (NH₃) and nitrogen monoxide (NO), which are mainly produced by the burning of fossil fuels (including vehicle emissions), chemical production processes, and biomass combustion, have adverse effect on health and cause respiratory illness, cardiovascular and immune diseases [1,2]. For example, the Occupational Safety and Health Administration (OSHA) standard for NO gases is given as 25 ppm for 8 h of work shift and 50 ppm for NH₃. Levels over these limits result in severe breathing difficulties, irritation to the nose, eyes, skin and throat and even death. Furthermore, NH₃ and NO gases also lead to pollution of water and soil, which results in further harm to plants and aquatic organisms. Therefore, it is of great importance to develop gas sensors with high performance for NH₃ and NO detection.

Recently, research related to gas sensors based on nanoclusters has attracted growing attention owing to their excellent advantages, such as large surface-to-volume ratio, high carrier mobility, and quantum confinement, which directly affect the sensing performances (e.g. sensitivity and selectivity) [3–23]. For example, Suematsu et al. [8] have found that Pd-loaded SnO₂ nanoparticle clusters, which were synthesized by a hydrothermal method, showed excellent sensitivity to toluene which they were able to detect down to low ppb levels, while Liu et al. [9] have found that the sensor based on Pt activated SnO₂ nanoparticle clusters not only has ultrahigh sensitivity, but also possesses good response–recovery properties, linear dependence, repeatability, selectivity and long-term stability for NH₃ detection. In particular, metallic nanoclusters have been viewed as promising candidates for gas sensor applications, especially for gold-based nanoclusters [3–6,12,17,21]. However, the physicochemical properties of metallic nanoclusters are found to have strong dependency on the cluster size, geometry, and composition [3]. It is clearly important to understand the characteristics size, geometry, and composition needed for use in gas sensor applications. More importantly, enhancing the gas sensing performance of cluster-based gas sensors is one of the main challenges in the development of high performance gas sensors. In recent years, the hydrogenation strategy has been successfully employed to improve sensing performances of the gas sensors based on metal oxide sensors

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[24–27]. The sensing active sites can be modified by the hydrogenation process, which adjusts the adsorption strength and response of gases on the sensors.

Apart from the fundamental understanding of quantum effects such as the finite-size effect, the interest in Si/Au clusters is primarily based on their applications such as molecular electronic devices, catalyst, gas sensors, etc. [28–41]. There are three main approaches to constructing these clusters: (i) Au doped Si clusters [28–32]: the Au doping process can stabilize particular structures (such as a fullerene-like cage) of Si clusters and lead to unusual properties, such as size selectivity, different charge transfer, large HOMO–LUMO gaps. (ii) Si doped Au clusters [33–37]: the structures of Au clusters can be changed significantly, and the reactivity and catalytic activity of Au clusters affected by Si doping [37]. (iii) mixed Si/Au clusters [38–41], where very recently, Guo et al. have predicted a very stable configuration of Si₁₂Au₂₀ cluster [41] using electronic structure methods. The stability of Si₁₂Au₂₀ cluster is verified by vibrational frequency analysis, molecular dynamics simulations, and electronic properties such as HOMO–LUMO gap [41]. The hollow, cage-like Si₁₂Au₂₀ cluster with I_h symmetry can be viewed as a hollow catalan pentakis dodecahedron, similar to the case of Cu₂₀Si₁₂ cluster [42], and may have potential applications in the electronics industry if suitable fabrication process can be developed.

In this work, using computational electronic structure techniques, we investigated the adsorption behaviour and electronic properties of NH₃, NO, CH₄, CO₂, H₂, N₂, O₂, and H₂O molecules on the Si₁₂Au₂₀ cluster to study the potential of the Si₁₂Au₂₀ cluster in gas sensors for NH₃ and NO detection. Our results show that NH₃, NO, and O₂ molecules are chemisorbed on the Si₁₂Au₂₀ cluster with large adsorption energies and charge transfer; however, CH₄, CO₂, H₂, N₂, and H₂O molecules are all physically adsorbed on the cluster. The changes of electronic properties of the Si₁₂Au₂₀ cluster, especially for the electric conductivity, are obvious before and after adsorption of NH₃ and NO. However, the strong adsorption of NH₃ and NO on the Si₁₂Au₂₀ cluster inhibits desorption which would result in a very long recovery time (over hours). The pure Si₁₂Au₂₀ cluster would not therefore be suitable as renewable gas sensors for NH₃ and NO detection, though it has high performance of sensitivity and selectivity. To reduce the strong adsorption of Si₁₂Au₂₀ cluster, we then considered the surface modification of the Si₁₂Au₂₀ cluster by hydrogenation. Our calculations predict a stable hydrogenated Si₁₂Au₂₀ cluster (H₁₂Si₁₂Au₂₀), whose stability was confirmed by vibrational frequency analysis and molecular dynamics simulations. Then the adsorption of NH₃, NO, CH₄, CO₂, H₂, N₂, O₂, and H₂O molecules on the H₁₂Si₁₂Au₂₀ cluster was investigated. Our results highlight that the H₁₂Si₁₂Au₂₀ is a new promising candidate as a gas sensor with potential for high performance (high stability, sensitivity and selectivity, short recovery time) for NH₃ and NO detection. In particular, our results suggest that hydrogenated Si/Au nanoparticles could be effective in the detection of ultralow concentrations of NH₃ and NO.

2. Computational method and details

All calculations in this work have been carried out using spin-polarized density functional theory (DFT) implemented in the DMol³ package (Accelrys Inc.) [43,44]. The exchange–correlation energy functional was treated using the generalized gradient approximation (GGA) formulated by Perdew, Burke, and Ernzerhof (PBE) [45] with van der Waals (vdW) correction under the scheme of Tkatchenko and Scheffler [46]. The electron ion–core interactions were represented by density-functional semi-core pseudopotentials fitted to all-electron relativistic DFT results [47] and the standard DNP basis sets. The convergence criterion of the energy and electron density was set to 1×10^{-6} Hartree and 1×10^{-6} e/bohr³ in the self-consistent field calculations, respectively. Meanwhile, convergence criteria of 1×10^{-3} Hartree/Bohr on the gradient and displacement and 1×10^{-5} Hartree on the total energy were set in geometrical optimization. The geometric parameters were optimized without symmetry constraints. The charge distributions and

the charge transfer between cluster and molecules were analysed based on Hirshfeld method [48]. Normal-mode frequency vibrational analysis was applied to ensure the structures obtained are real minima; all the configurations presented in this work have no imaginary frequencies. Transition state (TS) geometries were firstly searched by using the complete linear/quadratic synchronous transit (LST/QST) method [49], and then fully optimized within the DFT–PBE level. The ab-initio Born–Oppenheimer molecular dynamics (BOMD) simulation was carried out in a NVT ensemble. A Nosé–Hoover chain of thermostats was used to control the temperature.

To evaluate the adsorption strength of molecules on (hydrogenated) Si₁₂Au₂₀ cluster, the adsorption energy (E_{ads}) per molecule is defined as

$$E_{\text{ads}} = \frac{1}{n} (E_{\text{system}} - E_{\text{cluster}} - nE_{\text{mol}}) \quad (1)$$

where E_{system} , E_{cluster} and E_{mol} represents the total energy of the molecules adsorbed on the (hydrogenated) Si₁₂Au₂₀ cluster, the corresponding (hydrogenated) Si₁₂Au₂₀ cluster, and the pristine molecule, respectively, and n is the number of molecules.

3. Results and discussion

3.1. Adsorption of molecules on the pure Si₁₂Au₂₀ cluster

The pure Si₁₂Au₂₀ cluster with I_h symmetry was systematically investigated by Guo et al. [41]. Its configuration is shown in Fig. 1a, and its Cartesian coordinates are shown in Table S1 of Supplementary Information. Our calculated cohesive energy of the Si₁₂Au₂₀ cluster is 3.119 eV per atom, much larger than that of Au₃₂ cluster with I_h symmetry (2.338 eV per atom) and Si₃₂ cluster (2.331 eV per atom), indicating the greater stability of Si₁₂Au₂₀ cluster than Au₃₂ and Si₃₂ clusters. The HOMO–LUMO gap in present work is 1.038 eV, a little smaller by 0.019 eV than that of Guo et al. [41]. Meanwhile, our calculated cohesive energy is slightly larger than that of 3.069 eV per atom obtained by Guo et al. [41]. These differences are mainly due to the van der Waals (vdW) correction, which is considered in our calculations. The inclusion of the vdW correction generally gives more accurate energies and electronic properties of materials [46]. However, the structure of the Si₁₂Au₂₀ cluster is the same as predicted in previous work [41], indicating that the vdW correction does not affect the structural properties of the Si₁₂Au₂₀ cluster. Although Guo et al. [41] have verified the stability of Si₁₂Au₂₀ cluster by vibrational frequency analysis, MD simulations, and electronic structures, to further examine the dynamical stability of Si₁₂Au₂₀ cluster, we calculated its phonon dispersion. As shown in Fig. S1a (Supplementary Information), the Si₁₂Au₂₀ cluster has no imaginary frequency, indicating its dynamical stability.

Next, we investigated the adsorption of molecules on the Si₁₂Au₂₀ cluster. To obtain the most stable configurations of each adsorbed molecule, we have considered all possible adsorbed configurations. Since the Si₁₂Au₂₀ cluster has a high symmetry (I_h), all Au or Si atoms are equivalent. We thus considered two top sites (Si and Au atom), two types of bridge sites (Si–Au bonds and Au–Au bonds), and one type of hollow sites (triangle formed by Au–Au–Si) for the adsorption sites. We also considered the orientations of molecules on the adsorption sites. Taking NO molecule as an example, we considered the orientations in which both O and N atoms point to the adsorption sites, and the N–O bond parallel or vertical to the adsorption sites. After full relaxation of the several configurations considered, we have obtained the most stable plus a number of low-lying energy structures for each molecule adsorbed on the Si₁₂Au₂₀, which are shown in Figs. 1 and 2, while the corresponding calculated results are listed in Tables 1 and 2. To assist the visualisation of the configurations, the cage space is filled by light grey as shown in all the figures presented in the work. It is noted that all the configurations discussed in the subsequent section are found to have no imaginary frequencies, indicating that they are real stable energy minima.

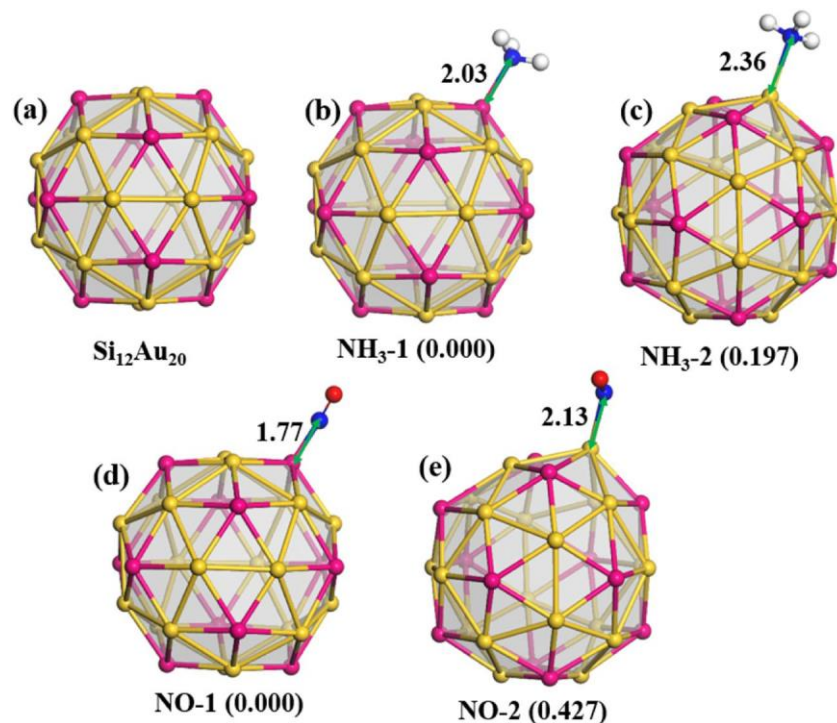


Fig. 1. The optimized structures of $\text{Si}_{12}\text{Au}_{20}$ cluster without and with gas molecules adsorption: (a) the pure $\text{Si}_{12}\text{Au}_{20}$ cluster; (b) and (c) NH_3 ; (d) and (e) NO . Values in parentheses are relative energies with respect to the most stable isomer for each adsorption in eV. The distances between molecules and cluster (shown in green arrowed line) are given in Å. Magenta, yellow, blue, white, and red balls are Si, Au, N, H and O atoms, respectively.

Firstly, we examined the adsorption of NH_3 on the $\text{Si}_{12}\text{Au}_{20}$ cluster. We found that the most stable configuration of NH_3 on the $\text{Si}_{12}\text{Au}_{20}$, which is shown in Fig. 1b, is formed by the N atom in NH_3 bonded with one Si atom. This configuration has an adsorption energy of

-0.967 eV. The $\text{Si}\backslash\text{N}$ bond lengths of 2.03 Å, is similar to the case of NH_3 adsorbed on silicene [50]. The adsorption energy, bond length, and charge transfer discussed below, indicate that the NH_3 molecule is chemisorbed on the $\text{Si}_{12}\text{Au}_{20}$ cluster. We also found chemisorption via

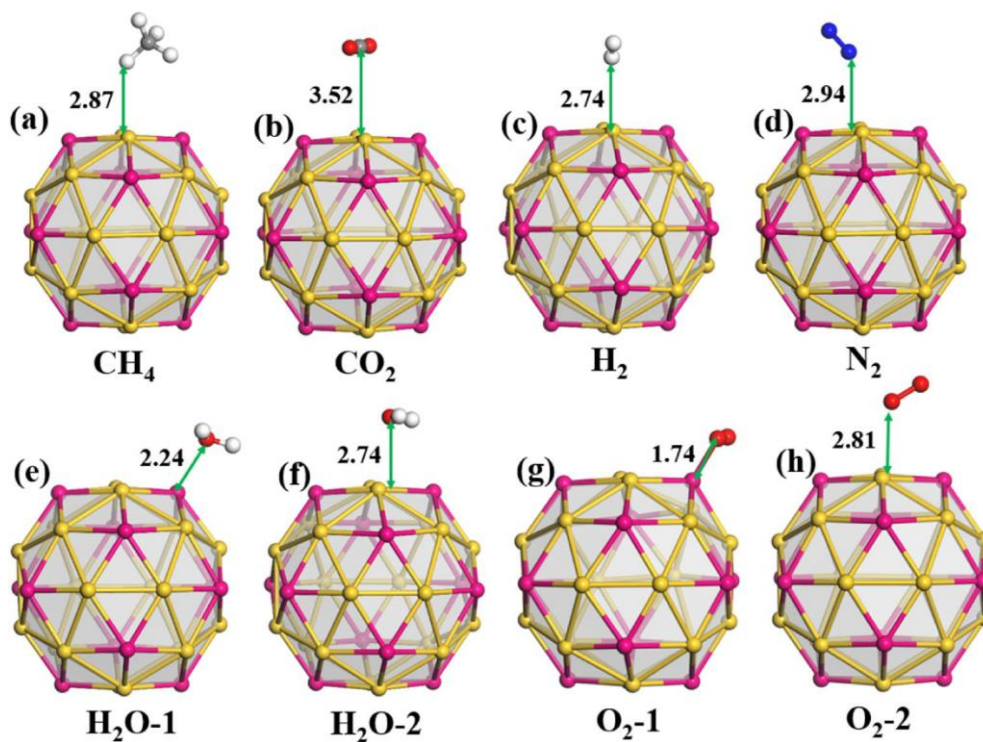


Fig. 2. The optimized structures of the $\text{Si}_{12}\text{Au}_{20}$ cluster with gas molecules adsorption: (a) CH_4 ; (b) CO_2 ; (c) H_2 ; (d) N_2 ; (e) and (f) H_2O ; (g) and (h) O_2 . The shortest distances between molecules and cluster (shown in green arrowed line) are given in Å. Grey balls are C atoms.

Table 1

Calculated adsorption energy (E_{ads}), charge transfer from the $\text{Si}_{12}\text{Au}_{20}$ cluster to molecules (E_{T}), the shortest distances between the $\text{Si}_{12}\text{Au}_{20}$ cluster and molecules (D), the HOMO-LUMO gap (E_{g}), and the recovery time (τ) for the configurations of NH_3 and NO adsorption on the $\text{Si}_{12}\text{Au}_{20}$ cluster.

System	E_{ads} (eV)	E_{T} (e)	D (Å)	E_{g} (eV)	τ (s)
NH3-1	-0.967	0.343	2.03	0.997	1.4×10^4
NH3-2	-0.770	0.195	2.36	0.863	7.3
NO-1	-1.058	-0.133	1.77	0.382	4.7×10^5
NO-2	-0.631	-0.064	2.13	0.459	3.5×10^{-2}

the covalent Au\N bond, which is shown in Fig. 1c, with an adsorption energy of -0.770 eV, which is higher by 0.197 eV than that of the most stable configuration. We note that the adsorption of NH_3 on the cluster via the N\Au bond leads to some deformation on the cluster, which occurs mainly in the region close to the Au atom bonded with the N atom (see Fig. 1c). The average Au\Au and Au\Si bond lengths around the adsorption site increase from 2.76 Å and 2.57 Å in pure cluster to 2.83

Å and 2.73 Å, respectively. The deformation electron density as shown in Fig. S2 (Supplementary Information) and the Hirshfeld analysis indicate there is an apparent charge transfer (of at least 0.195 e) from the $\text{Si}_{12}\text{Au}_{20}$ cluster to the NH_3 molecule (see Table 1), especially for the most stable configuration, indicating that NH_3 serves as an acceptor. Furthermore, the adsorption of NH_3 narrows the HOMO-LUMO gap of the cluster. We see from Table 1 that the HOMO-LUMO gap of isomer NH3-1 and NH3-2 changes from 1.038 eV in the pristine cluster to 0.997 and 0.863 eV, respectively, i.e. a change of 3.95% and 16.9% .

For NO adsorption on the $\text{Si}_{12}\text{Au}_{20}$ cluster, the NO molecule can chemisorb on the cluster only via the N atom bonding with the cluster, as shown in Fig. 1d–e, as is the case of the NO molecule adsorbed on other Au-based clusters [5,12]. As the most stable configuration as shown in Fig. 1d, the N atom in NO is bonded with Si to form an Si\N bond with a bond length of 1.77 Å. It has an adsorption energy of -1.058 eV, which is larger than that of NH_3 on the $\text{Si}_{12}\text{Au}_{20}$ cluster. This may be due to the shorter Si\N bond in the isomer NO-1 compared with that in isomer NH3-1, which leads to much stronger interaction in Si\N bond. Although there is a large adsorption energy, it does not result in any structural deformation for the cluster. When the N

atom in NO interacts with an Au atom, which forms the isomer NO-2 as shown in Fig. 1e, the adsorption energy decreases to -0.631 eV. However, this adsorption induces a substantial structural deformation of the cluster, which is similar to the case of NO molecule adsorbed on Au-based clusters [5,12]. Since the NO molecule can be viewed as an electron donor, some charge (about 0.064 – 0.133 e) is transferred to the $\text{Si}_{12}\text{Au}_{20}$ cluster for the two most stable configurations (see the deformation electron density as shown in Fig. S2 of Supplementary Information). Furthermore, the adsorption of the NO molecule results in a magnetic moment of approximately $1 \mu_{\text{B}}$, which is similar to the adsorption of NO on other nonmagnetic clusters [5,12–14]. The HOMO-LUMO

Table 2

Calculated adsorption energy (E_{ads}), charge transfer from the $\text{Si}_{12}\text{Au}_{20}$ cluster to molecules (E_{T}), the shortest distances between the $\text{Si}_{12}\text{Au}_{20}$ cluster and molecules (D), and the HOMO-LUMO gap (E_{g}) for the configurations of CH_4 , CO_2 , H_2 , N_2 , H_2O and O_2 adsorption on the $\text{Si}_{12}\text{Au}_{20}$ cluster.

System	E_{ads} (eV)	E_{T} (e)	D (Å)	E_{g} (eV)
CH_4	-0.169	0.051	2.87	1.028
CO_2	-0.204	0.027	3.52	1.035
H_2	-0.109	-0.041	2.74	1.029
N_2	-0.138	-0.002	2.94	1.036
H_2O -1	-0.378	0.189	2.24	1.022
H_2O -2	-0.345	0.036	2.74	1.000
O_2 -1	-0.895	-0.277	1.74	0.745
O_2 -2	-0.139	-0.052	2.81	1.036

gap of isomer NO-1 and NO-2 is 0.382 eV and 0.459 eV, respectively, indicating that there is an obvious change in the HOMO-LUMO gaps after the adsorption of NO , and the change is in the range of 55.8% to 63.2% . Comparing the HOMO-LUMO gaps of NO -cluster complexes with the pristine cluster clarifies the influence of NO adsorption on the electronic properties of the $\text{Si}_{12}\text{Au}_{20}$ cluster. We found that the adsorption of NH_3 and NO molecules can narrow the HOMO-LUMO gaps, which is because both of molecule adsorption can introduce some impurity states near Fermi level, and unoccupied local states in the unoccupied molecular states as shown in Fig. S3 (Supplementary Information).

We also considered other common gas molecules (CH_4 , CO_2 , H_2 , N_2 , H_2O , and O_2) adsorbed on the $\text{Si}_{12}\text{Au}_{20}$ cluster. Although we explored several initial structures for each molecule on the cluster, we found that they are all physically adsorbed on the $\text{Si}_{12}\text{Au}_{20}$ cluster with much small adsorption energy and no significant charge transfer except for the H_2O and O_2 molecules. The most stable configuration of each molecule on the cluster is shown in Fig. 2, and the relevant results are listed in Table 2. It can be seen from Table 2 that the physisorption features for CH_4 , CO_2 , H_2 , and N_2 on the $\text{Si}_{12}\text{Au}_{20}$ cluster are very evident, and their adsorption does not significantly affect the electronic properties of the $\text{Si}_{12}\text{Au}_{20}$ cluster. However, in the most stable configuration of H_2O on the $\text{Si}_{12}\text{Au}_{20}$ cluster, which has an adsorption energy of -0.378 eV, the distance between the O atom and the Si atom is 2.24 Å, which is much larger than the covalent Si\O bonds of ~ 1.65 Å in other Si\O systems [51–53]. There is, however, a significant charge transfer (0.189 e) between the H_2O molecule and the cluster. Therefore, the adsorption of H_2O on the $\text{Si}_{12}\text{Au}_{20}$ via O interacting has aspects that are intermediate between strong physisorption and chemisorption. However, when the H_2O molecule is close to an Au atom to form the configuration of isomer H_2O -2 as shown in Fig. 2f, the distance between O and Au atom is 2.74 Å, which is similar to the case of H_2O adsorbed on Gd@Au_n ($n = 14, 15$) clusters [12], and indicates a physisorption state for this configuration. The physisorption of CH_4 , CO_2 , H_2 , N_2 and H_2O molecules on $\text{Si}_{12}\text{Au}_{20}$ cluster just introduces some occupied states, which is located at ~ 2.0 eV below the Fermi level, resulting in little influence on the HOMO-LUMO gap (See the Fig. S3 of Supplementary Information, where we just showed the DOS of CO_2 and H_2O adsorption due to the similar features of CH_4 , H_2 , N_2 , CO_2 and H_2O). So the electronic properties of the cluster can be affected by the adsorption of these molecules. For the adsorption of O_2 on the $\text{Si}_{12}\text{Au}_{20}$ cluster, the singlet and triplet states are considered in the spin-polarized calculations. Here we just discussed the ground-state triplet O_2 on the $\text{Si}_{12}\text{Au}_{20}$ cluster. The most stable configuration as shown in Fig. 2g, in which one O atom in O_2 molecule is bonded with an Si atom to form an Si\O bond with the length of 1.74 Å, has an E_{ads} of -0.895 eV, accompanied by a charge transfer of 0.277 e from the cluster to O_2 molecule, indicating that the triplet O_2 chemisorbs on the cluster. More importantly, the HOMO-LUMO gap of the cluster narrows to 0.745 eV due to the adsorption of O_2 molecule. However, we found that triplet O_2 only physisorbs on the top of Au atoms (see Fig. 2h). The equilibrium distance of the O_2 -cluster is 2.81 Å, with small adsorption energy (-0.126 eV). This weak physisorption hardly affects the electronic properties of the cluster. These results indicate that, if we consider the $\text{Si}_{12}\text{Au}_{20}$ cluster as gas sensors for toxic gas detection, we must prevent the interaction between the H_2O and O_2 molecules and the Si sites in the cluster, as will be discussed below.

3.2. The possibility of the $\text{Si}_{12}\text{Au}_{20}$ cluster as gas sensors for NO and NH_3 detection

We then explored the possibility of the $\text{Si}_{12}\text{Au}_{20}$ cluster as a gas sensor for NO and NH_3 detection. As a good sensor, it should have excellent performance (sensitivity, selectivity, speed, and stability). The $\text{Si}_{12}\text{Au}_{20}$ cluster has been confirmed to have high thermodynamic stability by previous work [41]. The sensitivity can be estimated via calculating

the change of electric conductivity (σ) of the $\text{Si}_{12}\text{Au}_{20}$ cluster before and after adsorption according to the equation:

$$\sigma \propto \exp \left(\frac{-E_g}{2kT} \right); \quad \delta 2P$$

where E_g is the HOMO-LUMO gap value of the configurations, k is the Boltzmann's constant, and T is the thermodynamic temperature. According to this equation, the electric conductivity of the $\text{Si}_{12}\text{Au}_{20}$ cluster at a certain temperature rises by reducing the HOMO-LUMO gap. As discussed above, the HOMO-LUMO gap of the most stable configurations of NO (or NH_3) on the $\text{Si}_{12}\text{Au}_{20}$ cluster significantly decreases, which will lead to a substantial modification of the electric conductivity, indicating that the sensitivity of the $\text{Si}_{12}\text{Au}_{20}$ cluster for NO (or NH_3) detection would be much high.

As discussed above, the CH_4 , CO_2 , H_2 , N_2 , and H_2O are all physisorbed on the $\text{Si}_{12}\text{Au}_{20}$ cluster, and have negligible influence on the electronic properties (mainly conducting properties) of the $\text{Si}_{12}\text{Au}_{20}$ cluster, however, the adsorption of NO and NH_3 can remarkably change the electronic properties of the cluster (i.e. a large change in the HOMO-LUMO gap before and after adsorption is found). Therefore, the $\text{Si}_{12}\text{Au}_{20}$ -based gas sensor is predicted to show high selectivity for NH_3 and NO detection compared with CH_4 , CO_2 , H_2 , N_2 , and H_2O , which should be similar to the case of B_{40} -based gas sensor [15]. Using the non-equilibrium Greens function (NEGF) method, Lin et al. have predicted high sensitivity and selectivity for the B_{40} fullerene-based sensor for NH_3 detection in comparison with H_2 , N_2 and CH_4 [15]. Another important parameter of gas sensor is speed, i.e. the response and recovery rate. However, it is difficult to evaluate the response time due to the lack of knowledge of several parameters. In the present work, we attempt crudely to estimate the recovery time (τ) based on transition state theory, applied to the rate of desorption:

$$\tau \approx \nu_0^{-1} \exp \left(\frac{E_{\text{ads}}}{kT} \right); \quad \delta 3P$$

where ν_0 is the attempt frequency, E_{ads} is the adsorption energy as defined in Eq. (1). The adsorption energies ($-0.34 \sim -0.79$ eV) and recovery times ($0.5 \mu\text{s} \sim 16$ s) from previous theoretical calculations on NO_2 adsorbed on carbon nanotubes are remarkably smaller and shorter than the corresponding experimental results [54]. Peng et al. suggested that when NO_2 molecules experimentally interact with the carbon nanotubes, a reaction occurs to produce to NO and NO_3 molecules. NO and NO_2 have fast recovery time (≤ 1 s at room temperature), while NO_3 has longer ones (12 h), and the NO_3 concentration is major on the nanotube surface. So they deduce that NO_3 is responsible for the slow recovery [54]. Thus, we can successfully explain the reason why the experimental recovery time has been measured to be long or short by using the theory mentioned above. If we assume that the attempt frequencies of all the considered molecules have the same value as that for the NO_2 molecule (10^{12} s^{-1}) [54], the calculated recovery times at $T = 300$ K are listed in Table 1. We find that when the N atom is bonded with Si to form the most stable configuration of NH_3 or NO on the $\text{Si}_{12}\text{Au}_{20}$ cluster, the estimated recovery times for NH_3 and NO are 3.95 h and 130.7 h, respectively, which is too long to allow the reuse of $\text{Si}_{12}\text{Au}_{20}$ -based sensor. Furthermore, the interaction between O_2 and the $\text{Si}_{12}\text{Au}_{20}$ cluster, with important implications for the stability of the cluster upon air exposure, is so strong that it may hinder the applications of the cluster as chemical sensors. When these results (adsorption energy, charge transfer, thermodynamic stability, high sensitivity and selectivity, and recovery time) are considered together, the $\text{Si}_{12}\text{Au}_{20}$ sensor is predicted to have high sensitivity and selectivity, but poor performance as to recovery time; thus, the $\text{Si}_{12}\text{Au}_{20}$ sensor could only be used as a disposable molecular sensor for NH_3 or NO detection.

3.3. The structure and stability of the $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ cluster

It is clear that the strength of the adsorption of NH_3 and NO and the consequent slow recovery times is the main challenge for developing $\text{Si}_{12}\text{Au}_{20}$ -based sensors. As discussed above, we found that the most stable configurations of NH_3 and NO on the $\text{Si}_{12}\text{Au}_{20}$ cluster, which have the biggest adsorption energy, are always formed by the molecule (NH_3 and NO) interacting with the Si atoms; however, the configurations that are formed by the molecule interacting with the Au atoms have moderate adsorption energy, which corresponds to a much shorter recovery time (see Table 1). Moreover, the adsorption energy of H_2O and O_2 on the $\text{Si}_{12}\text{Au}_{20}$ cluster is a little stronger. Thus, preventing the interaction between H_2O and O_2 molecules and the Si sites would decrease the adsorption strength and hence the recovery time. Previous work has demonstrated that the hydrogenation strategy can be successfully employed to improve sensing performances of the gas sensors [24–27]. Moreover, hydrogenated Si-based clusters are stable because of the hydrogen termination of the dangling bonds of each Si, which hinders interaction with surrounding materials [55–59]. Therefore, we considered the hydrogenated $\text{Si}_{12}\text{Au}_{20}$ cluster, hydrogenating the Si atoms in the clusters, i.e. $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ (so-called HSA).

To investigate the hydrogenation of the cluster, we first adsorbed a single H atom on the $\text{Si}_{12}\text{Au}_{20}$ cluster. We found that the H atom can bond with Au and Si atom to form stable configurations, which are shown in Fig. 3A and B, respectively. The adsorption energies of the H atom on Si and Au are -2.526 eV and -1.938 eV, respectively, indicating the formation of Si\H bond is significantly more energetically favourable, as are the cases of NH_3 and NO on the cluster. To discuss further the adsorption behaviour of H on the $\text{Si}_{12}\text{Au}_{20}$ cluster, the transition state (TS) between structure A formed by H bonded to Au and structure B with H bonded to Si was searched by the LST/QST method. Fig. 3 shows the reaction path from structure A to B, for which there is a single TS with only one imaginary frequency of $327.3i \text{ cm}^{-1}$. In the TS, the Si\H and Au\H bond lengths are 1.69 \AA and 1.90 \AA , respectively. The energy barrier of 0.327 eV is relatively low and the reaction is exothermic with a large reaction energy of 0.588 eV. H migration from structure A to B will occur rapidly at moderate temperatures unlike the reverse reaction. Therefore, it should be possible to form the HSA structure by hydrogen termination of 12 Si atoms on the $\text{Si}_{12}\text{Au}_{20}$ cluster.

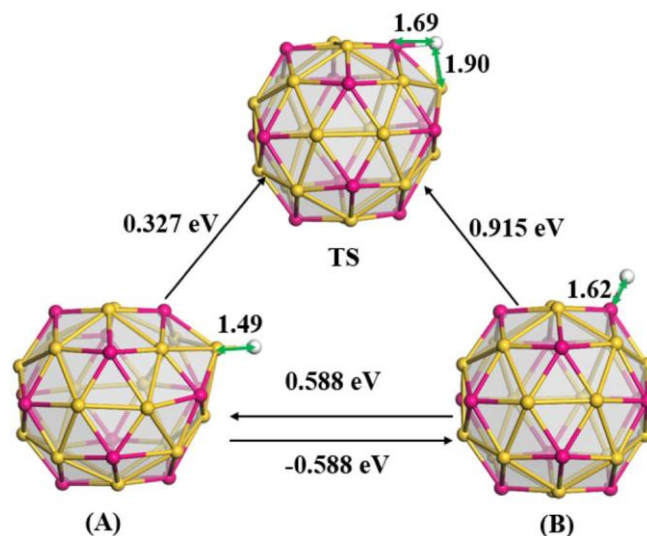


Fig. 3. Reaction path from the structure (A) formed by H atom bonded with Au atom to the structure (B) formed by H atom bonded with Si atom for H atom on the $\text{Si}_{12}\text{Au}_{20}$. The bond lengths (shown in green arrowed line) are given in Å .

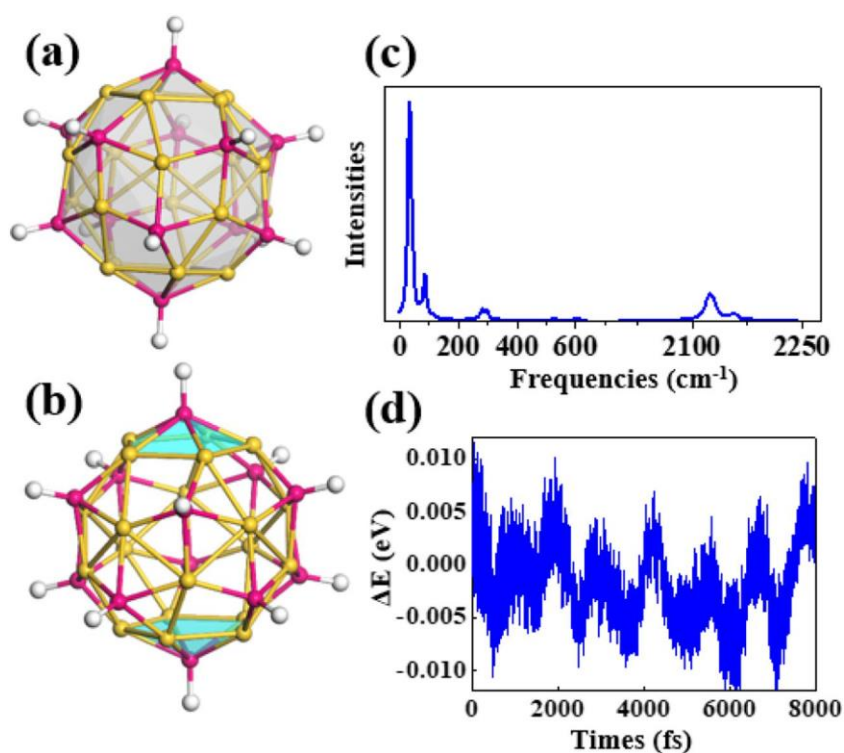


Fig. 4. (a) and (b) the optimized structure of the HSA. (c) Raman spectrum of the HSA. The temperature is 300 K and the incident light is 488.0 nm. The Lorentzian smearing is set to be 12.00 cm^{-1} . (d) Variation in the relative total energy (E) of the HSA structure as a function of time at $T = 300 \text{ K}$.

The optimized configuration of HSA is shown in Fig. 4a and b, and with cohesive energy, bond lengths, symmetry and HOMO-LUMO gap given in Table 3, meanwhile its Cartesian coordinates are shown in Table S1 of Supplementary Information. The structure of the pure $\text{Si}_{12}\text{Au}_{20}$ cluster as shown in Fig. 1a can be viewed as the composition of 12 Au pentagons with a Si atom at the hollow centre of each Au pentagon [41], and the average lengths of Au\Au bonds and Au\Si bonds are 2.76 \AA and 2.57 \AA , respectively. However, for the structure of the HSA, we found that only two Au pentagons with the Si atom at the centre of each pentagon (referred to as SiAu5 unit), as shown in green-filled areas in Fig. 4b, retain the same shape as in the pure $\text{Si}_{12}\text{Au}_{20}$ cluster, while the others have structural deformations. The average lengths of Au\Au bonds and Au\Si bonds in the SiAu5 unit are 2.74 \AA and 2.59 \AA , respectively, close to these of the pure $\text{Si}_{12}\text{Au}_{20}$ cluster. On the contrary, the average lengths of Au\Au bonds and Au\Si bonds in the other areas (i.e. not in the green filled areas as in Fig. 4b), are 2.83 \AA and 2.49 \AA , respectively. The increased Au\Au bond lengths and the decreased Au\Si bond lengths further show that the Si\Au bonds has significant polarity [41]. The average length of the Si\H bonds in the structure of HSA is 1.50 \AA , which is much smaller than the value of 1.62 \AA for one H atom on the $\text{Si}_{12}\text{Au}_{20}$ cluster, but consistent with previous work on hydrogenated Si-based clusters [56,57]. Because of these features, HSA has a high symmetry of S_{10} .

Next the stability of HSA was checked firstly by vibrational frequency analysis. We calculated the whole vibrational spectrum, but

Table 3
Cohesive energy (E_c), symmetry (S), the average bond lengths ($D_{\text{Au-Au}}$, $D_{\text{Si-Au}}$, and $D_{\text{Si-H}}$), and HOMO-LUMO gap of the $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ cluster (HSA).

System	E_c (eV)	S	$D_{\text{Au-Au}}$ (\AA) ^a	$D_{\text{Si-Au}}$ (\AA) ^a	$D_{\text{Si-H}}$ (\AA)	E_g (eV)
HSA	135.977	S_{10}	2.74/2.83	2.59/2.49	1.50	0.967

a The front values correspond to the bond lengths coming from the two SiAu5 units as green-filled areas as shown in Fig. 3b, while the back values correspond to the bond lengths coming from the rest areas.

just showed the Raman spectrum in Fig. 4c. It is noted that there is no frequency in the range of $650\text{--}2000 \text{ cm}^{-1}$. It can be seen from the Raman spectrum that there is no imaginary frequency, indicating the stability of the cluster. The highest frequency is 2159.79 cm^{-1} as shown in Fig. S4h (Supplementary Information), which relates to the vibration modes of the top H atom in the SiAu5 unit as shown in Fig. 4b. Moreover, the high frequencies in the range of over 2000 cm^{-1} all relate to the vibration modes of the H atoms (see Fig. S4). These results suggest that the structures of $\text{Si}_{12}\text{Au}_{20}$ and $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ can be easily characterized by the Raman spectra. To explore further the dynamical behaviour and thermal stability of HSA, the first-principles BOMD simulation and phonon dispersion were carried out. We set a simulation time of 8 ps with a time step of 1 fs, and the structural changes were calculated at a constant temperature of 300 K. Fig. 4d shows how the energies of the structure vary during the simulations. The total energies of HSA during the simulations remain close to constant at -209.903 eV within a very small range of variations (no 0.015 eV), strongly supporting the stability of the HSA structure at the temperature of 300 K. There is no imaginary frequency found for the HSA structure as shown in Fig. S2b (Supplementary Information), which is obtained by calculating the phonon dispersions, indicating its dynamical stability. Furthermore, the HOMO-LUMO gap of HSA is as large at 0.967 eV , just

a little smaller (by 0.071 eV) than that of the $\text{Si}_{12}\text{Au}_{20}$ cluster (1.038 eV), which indicates that the HSA structure is chemically stable. Therefore, the vibrational frequency analysis, MD simulation, phonon dispersions and electronic properties show that the HSA structure is mechanically, thermodynamically and chemically stable at the temperature of 300 K, which is the first step for its potential application in gas sensors.

3.4. Applications of HSA as a sensor for NH_3 and NO detection

Due to the high stability of the HSA structure, we next explored its application as a gas sensor for NH_3 and NO detection, mainly concerning the effect of hydrogenation on enhancing the gas sensing performance.

The aim of hydrogenation is to reduce the interaction between molecules and the Si atoms and to obtain the most stable configurations of molecules on the HSA structure, we considered all possible kinds of initial structure. The optimized structures of NH_3 and NO on HSA are shown in Fig. 5, and the corresponding results are summarized in Table 4. As we expected, we found that the stable configurations of NH_3 and NO molecules on the HSA structure are formed with the N atom bonded with Au atoms (see Fig. 5). More importantly, as shown in Fig. 5a and c, the most stable adsorption sites (i.e. the Au atoms) are located in the SiAu_5 units. For the most stable configuration of NH_3 on the HSA, we found that the length of the $\text{N}\backslash\text{Au}$ covalent bond is 2.47 Å, larger by 0.11 Å than that of NH_3 on the $\text{Si}_{12}\text{Au}_{20}$ cluster, which indicates a lower adsorption strength, which is consistent with the calculation of adsorption energy. The most stable configuration has an adsorption energy of -0.591 eV, much smaller than that of NH_3 on the $\text{Si}_{12}\text{Au}_{20}$ cluster. However, when the NH_3 molecule adsorbs on other Au atoms as shown in Fig. 5b, it can also form a low-lying energy configuration with an adsorption energy of -0.542 eV and with the $\text{N}\backslash\text{Au}$ bond length of 2.55 Å. From the deformation electron density as shown in Fig. S5a (Supplementary Information), we conclude that there is an apparent charge transfer (at least 0.15 e) from the HSA structure to the NH_3 molecule, and the HOMO-LUMO gap has been modified from the 0.967 eV in HSA to 0.937 eV and 0.925 eV for the isomer $\text{NH}_3\text{-HSA-1}$ and $\text{NH}_3\text{-HSA-2}$, respectively. These results show that the electronic properties of the HSA structure are influenced by the NH_3 adsorption, although the change in the gap is small.

For the NO adsorption on the HSA structure, the most stable configuration has an adsorption energy of -0.447 eV, much smaller than that of NO adsorption on $\text{Si}_{12}\text{Au}_{20}$ cluster, and the $\text{Au}\backslash\text{N}$ bond length is

Table 4

Calculated adsorption energy (E_{ads}), charge transfer from the HSA structure to molecules (E_{T}), the distances between the HSA and molecules (D), the HOMO-LUMO gap (E_{g}), and the recovery time (τ) for the configurations of NH_3 and NO adsorption on the HSA.

System	E_{ads} (eV)	E_{T} (e)	D (Å)	E_{g} (eV)	τ (s)
$\text{NH}_3\text{-HSA-1}$	-0.591	0.180	2.47	0.937	7.4×10^{-3}
$\text{NH}_3\text{-HSA-2}$	-0.542	0.152	2.55	0.925	1.1×10^{-3}
NO-HSA-1	-0.447	-0.022	2.18	0.342	2.9×10^{-5}
NO-HSA-2	-0.288	-0.017	2.53	0.237	6.5×10^{-8}

2.18 Å. The adsorption strength is further decreased when the NO molecule adsorbs on the Au atoms that are not in the SiAu_5 units (see Fig. 5d). In this case, the adsorption energy is just -0.288 eV, which is smaller by 0.159 eV than that of the most stable configuration. Appreciable charge (about 0.017–0.022 e) is found to be transferred from the NO molecule to the HSA structure, which can also be seen from the deformation electron density as shown in Fig. S5b (Supplementary Information). We note that the charges are obtained using the Hirshfeld method, which gives the smallest charge values compared with other schemes [60]. Although the HSA structure is nonmagnetic, after the adsorption of the NO molecule, there is about $1 \mu\text{B}$ magnetic moment generated from the configuration of NO-HSA systems, which is similar to the case of NO molecule on the $\text{Si}_{12}\text{Au}_{20}$ cluster. More importantly, the HOMO-LUMO gaps of NO-HSA systems are in the range of 0.237–0.342 eV, compared with 0.967 eV in HSA indicating that the electronic properties of the HSA structure are substantially changed by the adsorption of NO . Similar to the case of NH_3 and NO adsorbed on the pure $\text{Si}_{12}\text{Au}_{20}$ cluster, the adsorption of NH_3 and NO induces some impurity states near the Fermi level, which leads to the narrow of

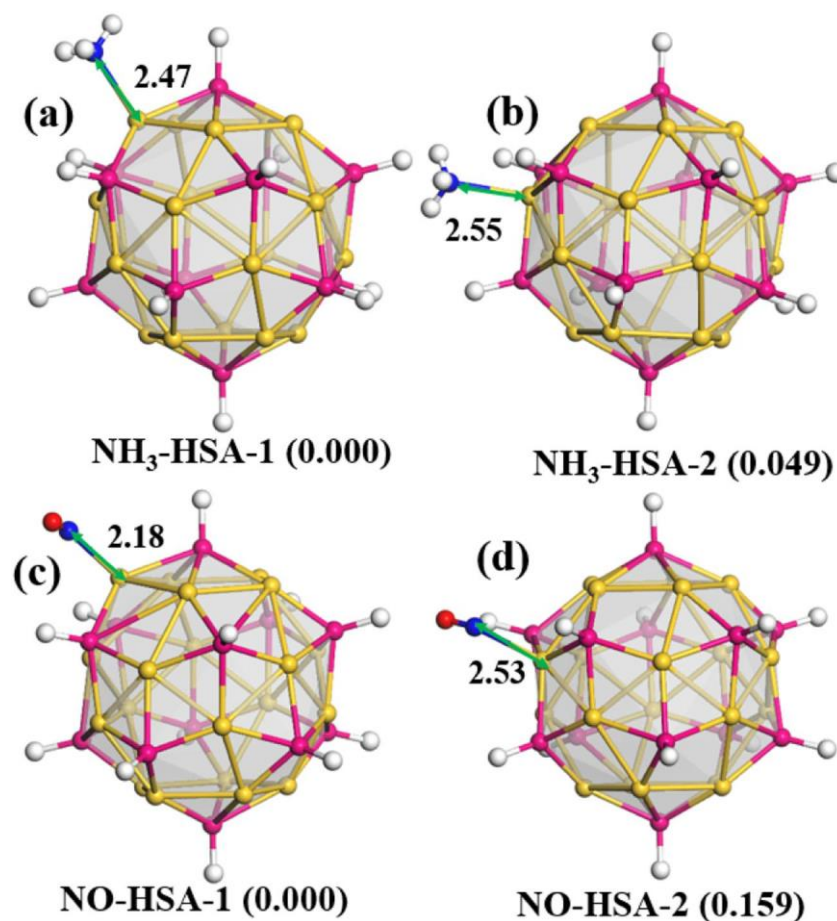


Fig. 5. The optimized structures of $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ cluster (HSA) with gas molecules adsorption: (a) and (b) NH_3 ; (c) and (d) NO . Values in parentheses are relative energies with respect to the most stable isomer for each adsorption in eV. The distances between molecules and cluster (shown in green arrowed line) are given in Å.

Table 5

Calculated adsorption energy per molecule (E_{ads}), charge transfer per molecule from the HSA to molecules (E_T), the average distances between the HSA and molecules (D), and the HOMO-LUMO gap (E_g) for the most stable configurations of more NO (NH_3) molecules adsorption on the HSA.

System		E_{ads} (eV)	E_T (e)	D (Å)	E_g (eV)
Species	Number of molecules				
NO	2	-0.411	-0.017	2.24	0.279
	3	-0.455	-0.027	2.19	0.183
	4	-0.463	-0.017	2.15	0.248
	5	-0.511	-0.026	2.38	0.156
	6	-0.365	-0.016	2.37	0.209
	NH_3	2	-0.573	0.170	2.47
3		-0.552	0.162	2.48	0.907
4		-0.533	0.149	2.52	0.864
5		-0.527	0.146	2.51	0.817
6		-0.504	0.132	2.56	0.783

HOMO-LUMO gap (See Fig. S6 of Supplementary Information). Overall, we can conclude that the hydrogenation of the $Si_{12}Au_{20}$ cluster makes the adsorption strength decrease, which will assist the development of $Si_{12}Au_{20}$ -based gas sensor with high performance.

We now discuss the gas sensing properties of the HSA-based sensor. The stability of the HSA structure has been shown above, and the adsorption of NO and NH_3 molecules on the HSA structure induce little deformation of the structure, further indicating its stability. The sensitivity of the HSA-based sensor is evaluated by calculating the change of electric conductivity via Eq. (2). The HOMO-LUMO gaps for the configurations of NH_3 (NO) molecule on the HSA structure are reduced in the range of 3.1%~4.4% (64.6%~75.5%) compared with that of the HSA structure, thus leading to an increase in the electric conductivity after the adsorption of the NH_3 (NO) molecule. We note that the change of HOMO-LUMO gap for NH_3 adsorption is much smaller compared with that due to NO adsorption, but we confirm that it should indeed result in the change of electric conductivity, which is similar to NH_3 adsorption on the B_{40} fullerene [15,16], monolayer GaN [61,62], atomic-layer MoS_2 [63], phosphorene [64], and monolayer Ti_2CO_2 [65]. Although the adsorption of NH_3 on these materials does not have a substantial effect on the electronic structures of these materials, the electric conductivity of these system changes obviously after the adsorption of NH_3 owing to the comparatively large charge transfer, that is to say, the adsorption-induced charge transfer affects the resistivity of the system. For example, Moradi et al. found that the change of HOMO-LUMO gap for the most stable configuration of NH_3 adsorption on B_{40} is about 7%, which

is small but would have a significant increase of electric conductivity [16]. This conclusion is demonstrated by Lin et al. [15] who have found that the adsorption of NH_3 distinctly affect the electric conductivity of the B_{40} using NEGF method. We would therefore expect a detectable change in conductivity due to NH_3 and a large change due to NO.

To demonstrate the high selectivity of HSA for NH_3 and NO detection, the adsorption behaviour of the HSA structure with CH_4 , CO_2 , H_2 , N_2 , H_2O , and O_2 are calculated, and the most stable configuration of each molecule on the HSA is shown in Fig. S7 (Supplementary Information), and the corresponding results are shown in Table S2 (Supplementary Information). We found that all these molecules are physisorbed on HSA, and there is virtually no influence on the electronic properties of HSA because of the molecular adsorption, which can also be found in the DOS as shown in Fig. S6 (Supplementary Information). These results conclude that these adsorptions have negligible effect on the electric conductivity of HSA. Therefore, it is expected that HSA show high selectivity to distinguish NO and NH_3 from CH_4 , CO_2 , H_2 , N_2 , O_2 , and H_2O gases. Furthermore, we can use the change of magnetic properties before and after adsorption to differentiate between NO and NH_3 . The pure HSA is nonmagnetic. The adsorption of NO introduces spin polarization in HSA with a magnetic moment of 1 μ_B , but the net spin polarization of HSA is not modified due to the adsorption of NH_3 . Finally, the recovery time can be calculated using Formula (3). Since the hydrogenated strategy has clearly decreased the adsorption strength of NH_3 and NO on the HSA structure, the recovery time is expected to be shorter. We have calculated the recovery times at the temperature of 300 K, which are summarized in Table 4. For most stable adsorption states of NH_3 and NO, the recovery time is estimated to be about 7.4 ms and 29 μs , respectively, which indicates high performance for the HSA-based gas sensor. Therefore, we conclude that HSA may show high performance as a gas sensor, including high stability, sensitivity, selectivity and a short recovery time for NH_3 and NO detection.

We further considered several NO and NH_3 molecules adsorbed on HSA to explore the coverage effect. In order to explore a range of different coverages, up to six molecules and several different starting configurations were investigated. The structural, energetic, and electronic properties of the most stable configurations for these systems are presented in Table 5 and Fig. 6. After the addition of each molecule, the system is geometry optimized prior to the addition of the subsequent molecule. Although interestingly, the most stable configuration for $5NH_3@HSA$ could only be generated from the second most stable configuration of $4NH_3@HSA$; however, we note that the energy difference between the first and second most stable configurations of $4NH_3@HSA$ is very small (only 0.008 eV). For NO adsorption we find that, the second

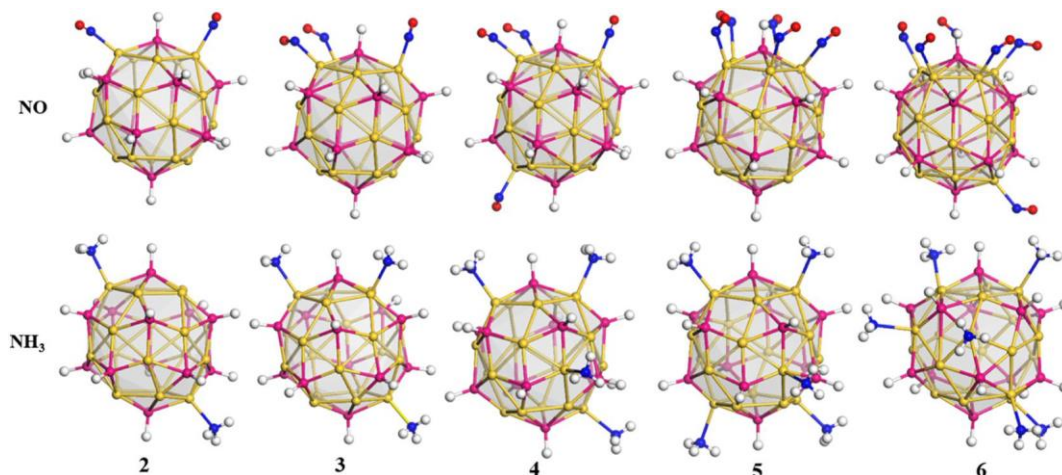


Fig. 6. The optimized structures of $H_{12}Si_{12}Au_{20}$ cluster (HSA) adsorbed by NO and NH_3 molecules of different coverage.

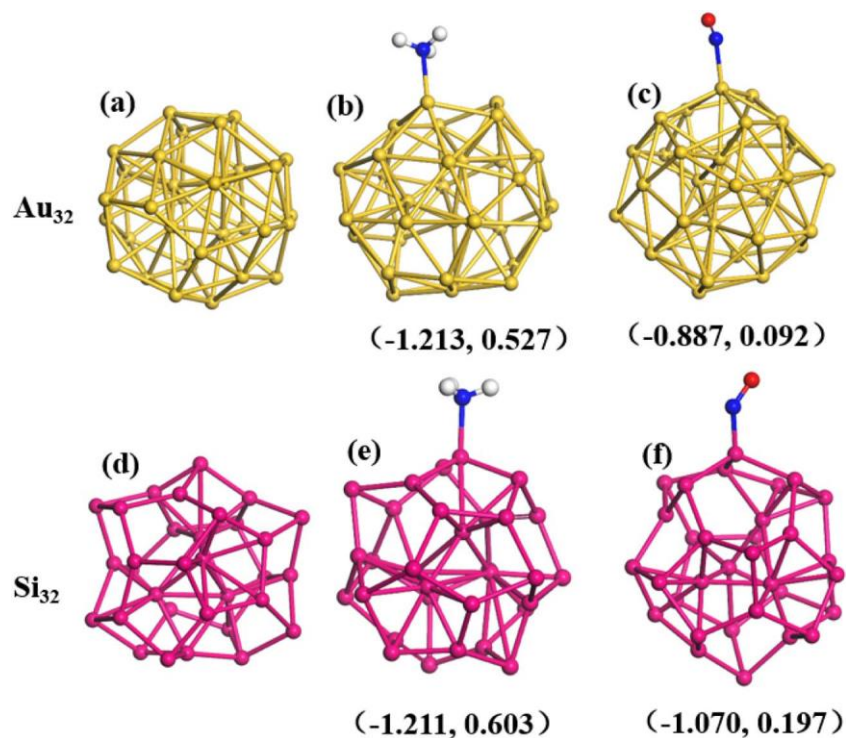


Fig. 7. The optimized structures of Au_{32} and Si_{32} clusters without and with NO and NH_3 adsorption. The values in parentheses, in turn, are adsorption energy and HOMO-LUMO gap.

configuration for four NO molecules are all located at the same SiAu_5 unit. Moreover, in general, for NO adsorption with different coverages, the changes of E_{ads} are very small, which is important for their use in gas sensors, while the HOMO-LUMO gaps basically are all smaller than that of one NO molecule adsorption. When the number of NO molecules reaches 6, the adsorption energy becomes much higher. For NH_3 adsorption, the E_{ads} and HOMO-LUMO gaps decrease as coverage increases but the variation of E_{ads} is still small. These results further indicate that HSA is a promising candidate for gas sensors with high performance for NH_3 and NO detection with low coverage.

Moreover, to have comprehensive and comparable studies on the gas sensing properties of NO and NH_3 molecules adsorbed on the (hydrogenated) $\text{Si}_{12}\text{Au}_{20}$ cluster, we also calculated the adsorption of NO and NH_3 molecules on the Au_{32} and Si_{32} clusters, which have the same size as $\text{Si}_{12}\text{Au}_{20}$ cluster. Although the Au_{32} cluster with I_h icosahedral symmetry is the most stable structure at zero temperature, experimental and theoretical studies have found that its amorphous isomer with C_1 symmetry becomes the most stable above 300 K [66–68]. In general, we aim for a gas sensor that can be used at room or higher temperatures, thus, in the present work, we only considered the most stable structure of Au_{32} cluster above 300 K. The most stable structure of the Si_{32} cluster can be viewed as endohedral fullerene (i.e. $\text{Si}_4@Si_{28}$) [69,70]. After the relaxation of NO and NH_3 molecules on the Au_{32} and Si_{32} clusters, the results are shown in Fig. 7. It is found that the adsorption energies of both NO and NH_3 on the Au_{32} and Si_{32} clusters are much bigger (at least -0.887 eV). These adsorption energies are so large that the recovery times of both clusters would be long. For example, the adsorption energies of NO on the Au_{32} cluster is -0.887 eV, which corresponds to a recovery time of about 10 min, while the others correspond too much longer recovery times (over hours). As discussed above, long recovery times will impede the reuse of gas sensors. Moreover, previous studies indicated that the adsorption energies of CO and O_2 molecules on the Au_{32} cluster are all larger than -1 eV [68], indicating that the selectivity of the Au_{32} cluster is not good. Therefore, from the viewpoint of stability, recovery time and selectivity, the pure Au_{32} and Si_{32} clusters are not suitable candidates for gas sensors for detecting NO and NH_3 .

4. Summary and conclusions

Using DFT techniques, we have investigated the adsorption behaviour and electronic properties of NH_3 , NO, CH_4 , CO_2 , H_2 , N_2 , O_2 , and H_2O molecules on the (hydrogenated) $\text{Si}_{12}\text{Au}_{20}$ cluster in order to find a promising molecular sensor for NH_3 and NO detection. NH_3 and NO are chemisorbed on the $\text{Si}_{12}\text{Au}_{20}$ cluster with a large adsorption energy and apparent charge transfer, while the other molecules are all physisorbed on the cluster. The electronic properties of the $\text{Si}_{12}\text{Au}_{20}$ cluster change dramatically due to the adsorption of NH_3 and NO molecules, particularly regarding the electric conductivity. Our results suggest that the $\text{Si}_{12}\text{Au}_{20}$ -based sensor could show high sensitivity and selectivity; however, because of the strong adsorption of NH_3 and NO molecules, the $\text{Si}_{12}\text{Au}_{20}$ -based sensor would suffer a longer recovery time (over 3 h), which directly prevents its application in gas sensors. To improve the recovery time, we then explored the hydrogenation of Si atoms in the $\text{Si}_{12}\text{Au}_{20}$ cluster due to the strong adsorption occurring in the interaction between the molecules and the Si atoms. The structure, stability and electronic properties of the hydrogenated $\text{Si}_{12}\text{Au}_{20}$ cluster ($\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$, i.e. HSA) were investigated. The vibrational frequency analysis, MD simulation and electronic properties showed that the HSA structure is mechanically, thermodynamically and chemically stable at $T = 300$ K. We found that the adsorption strength of NH_3 and NO on the HSA is moderate and much smaller than that of NH_3 and NO on the $\text{Si}_{12}\text{Au}_{20}$ cluster. The other molecules considered are still physisorbed on the HSA, and have a negligible effect on the electronic properties of material. For NH_3 and NO, the recovery time of the HSA is predicted to be 7.4 ms and 29 μs , respectively. We also considered two to six NO or NH_3 molecules adsorbed on HSA, corresponding to different coverage, and the adsorption of NO and NH_3 on the Au_{32} and Si_{32} clusters, respectively.

Overall, we conclude that due to the moderate adsorption energy, clear charge transfer, significant change of electric conductivity, and very short recovery time, HSA can be viewed as a promising gas sensor with high performance for NH_3 and NO detection at low coverage.

Declaration of Competing Interest

There are no conflicts of interest to declare.

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Appendix A. Supplementary data

Additional data and figures related to the deformation electronic densities of the $\text{Si}_{12}\text{Au}_{20}$ and $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ cluster with NH_3 and NO ; the frequency modes of $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ cluster; The stable configurations of common molecules on the $\text{H}_{12}\text{Si}_{12}\text{Au}_{20}$ and the corresponding results (PDF).

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