

CrossMark  
click for updatesCite this: *RSC Adv.*, 2017, 7, 14537

# Confinement of hydrogen and hydroxyl radicals in water cages: a density functional theory study†

Liuxie Liu,<sup>a</sup> Shuang Mao,<sup>a</sup> Quan Li,<sup>\*a</sup> Xiaolan Wang,<sup>a</sup> Mingli Yang<sup>b</sup> and Laicai Li<sup>\*a</sup>

Density functional theory calculations with D3 empirical dispersion correction reveal that hydrogen and hydroxyl radicals encapsulated in typical water cages found in clathrate hydrates exhibit similar structures and properties in their confined states to those in their corresponding free states, including atomic charges, spin densities, and electronic configurations. Diffusion studies reveal that energy barriers exist for these radicals to approach or exit these water cages. Energy decomposition analyses further reveal that coulombic repulsion between the radical and water cage is responsible for these energy barriers and the inability of these species to react with cage water molecules. This study provides insight into mechanisms for the storage of free radicals, which is normally extremely difficult because of their high reactivities toward many substances.

Received 29th December 2016  
Accepted 26th February 2017

DOI: 10.1039/c6ra28804a

rsc.li/rsc-advances

## Introduction

Clathrate hydrates are crystalline solid compounds that exist widely in ocean sediments, permafrost regions, and comets and on certain planets.<sup>1–9</sup> Interest in clathrate hydrates arose due to of two aspects. First, their ability to store methane on a large scale, both underground and in the ocean, means that they might become a major source of energy in the future. It has been shown that the methane reserves in these hydrates are perhaps twice as abundant as those in other proven fossil fuel sources, including oil, gas, and coal.<sup>10–16</sup> The other important aspect is that clathrate hydrates can be used as nano-containers for confining radicals, which are of great significance in chemical reactions,<sup>17–21</sup> biological systems,<sup>22–25</sup> and ageing processes.<sup>26,27</sup> Free radicals, which contain one or more unpaired electrons, are reactive and usually exist as transient reaction intermediates.<sup>28</sup> They are key constituents that initiate many chemical reactions, but are difficult to contain for subsequent use because of their high reactivities. The containment of free radicals is of particular interest in synthetic and catalytic chemistry.

Over the past few years, a number of experiments have revealed that radicals can exist in clathrate hydrates with extended lifetimes.<sup>29–31</sup> Yeon *et al.*<sup>30</sup> observed stably entrapped

hydrogen radical (H·) in clathrate hydrates by electron spin resonance (ESR) and nuclear magnetic resonance (NMR) spectroscopies. Cha *et al.*<sup>31</sup> studied O<sub>2</sub> + Me<sub>4</sub>NOH in these hydrates at a 60 kGy irradiation dose and various temperatures and demonstrated the stable existence of superoxide radical anions in clathrate hydrates by magnetic property measurement system (MPMS) studies and ESR spectroscopy. Clathrate hydrates have cage-like structures that can encapsulate small molecules such as CH<sub>4</sub> and CO<sub>2</sub>.<sup>10–16</sup> These species, however, are chemically inert compared to reactive radicals. Encapsulating radicals in water cages and moderating their reactivities are therefore interesting issues for the design and development of nano-containers and nano-reactors. In this article, we present our theoretical study on the status of H· and hydroxyl radicals (OH·) in the water cages and reveal that coulombic interactions between these radicals and the water cages prevent them from combining. Moreover, the curvature of these water cages contributes to the enhancement of coulombic interactions.

The hydrates are thermodynamically stable when guest molecules occupy the host cages to a certain minimum level, and the interactions between the host and guest involve mainly van der Waals forces (noncovalent interactions).<sup>32,33</sup> To study the stability, Lebsir *et al.* and Khan computed the stabilization energy (SE<sub>k</sub>) of one cage occupied by different gases using Hartree–Fock (HF) and second-order Møller–Plesset perturbation theory (MP2). Their results showed that *ab initio* quantum chemical methods provide independent and accurate means of reaching a good understanding of the nature of hydrates and their host–guest interactions.<sup>34–36</sup> The HF method is somewhat less accurate, while MP2 is more expensive for larger molecules and tends to overestimate noncovalent interactions. More accurate and economical methods that correctly describe non-covalent interactions are therefore required. For a long time,

<sup>a</sup>College of Chemistry and Materials Science, Sichuan Normal University, Chengdu 610068, China. E-mail: liquan6688@163.com; lilcmail@163.com

<sup>b</sup>Institute of Atomic and Molecular Physics, Sichuan University, Chengdu 610065, China

† Electronic supplementary information (ESI) available: Bond lengths (Å) and binding energies of the optimized water cluster W<sub>6</sub> computed at different levels; the coordination energies of water cages s1 and s2; energies for water approach by and H· or OH·; and LMOEDA energy decomposition analyses for H· and OH· radical attack at waters. See DOI: 10.1039/c6ra28804a

density functional theory (DFT) was disregarded as a reliable method for the description of noncovalent interactions. In particular, for dispersion-dominated complexes that require a non-localized energy description, GGA functionals are inadequate as they do not explicitly describe these dispersion phenomena.<sup>37</sup> Numerous ways for overcoming these disadvantages of DFT currently exist: the use of DFT-D methods, which have empirical dispersion corrections added to their functionals, is one example. A significant amount of work has been directed toward validating the performance of the DFT-D3 method,<sup>38</sup> which has been shown to be reliable for describing noncovalent interactions.<sup>38–41</sup>

In this work, we employ DFT-D3 to study whether or not H· and OH· radical are stable in typical clathrate hydrate structures I and/or II and focus on variations in the  $SE_k$  of the hydrates, charges and spin densities of the encapsulated free radicals, and energy barriers for radical transfer between cages.

## Computational methods

There are three common clathrate hydrate structures: structure I, structure II, and structure H, which respectively contain  $5^{12}-5^{12}6^2$ ,  $5^{12}-5^{12}6^4$ , and  $5^{12}-4^35^66^3-5^{12}6^8$  cages.<sup>3,42,43</sup> The third cage, structure H, has seldom been studied. Therefore, this work focused on the first two structures. Two types of water cages, s1 and s2, with the structural motifs of structures I and II, respectively, were constructed. The first, s1, consists of one  $5^{12}$  cage linked to another  $5^{12}6^2$  cage, while s2 contains one  $5^{12}$  and one  $5^{12}6^4$  cage, as shown in Fig. 1. The initial coordinates of the O atoms were taken from previous studies,<sup>36,44–46</sup> however, some of the H atoms are oriented differently.

The structures of empty s1 and s2 were optimized within the DFT framework using the B97D,<sup>47</sup> B3LYP,<sup>48</sup> BLYP,<sup>48</sup> and PBE<sup>49</sup> exchange-correlation functionals with Grimme's D3 dispersion correction.<sup>38</sup> The triple- $\zeta$  basis set def-TZVP,<sup>50</sup> which includes polarized function, as implemented in the Turbomole suite,<sup>51</sup> was used. These four methods produce similar geometries for the empty s1 and s2 cages. To further validate our computational strategy, we also optimized the cage structure of the water hexamer,  $W_6$ , at the MP2/def-TZVP<sup>52</sup> level, and compared the results obtained with DFT-D3 method. The calculated bond lengths are given in Table S1,<sup>†</sup> while the binding energies computed at the four DFT-D3 levels are listed in Table S2.<sup>†</sup> It is clear that the four DFT-D3 methods provide results that are in reasonable agreement with those of MP2. In particular, B97D-D3 provides the best agreement with MP2. The largest structural discrepancy, 0.01 Å,

was found for an H···O bond. Hence, B97D-D3 was mainly used in further calculations.

The two free radicals, H· and OH·, were separately placed at the centers of these cages to study the encapsulation of free radicals in clathrate hydrates, and the resulting structures were fully optimized without any constraints. The H· and OH· radicals were placed in either cage (s1 or s2) and all possible variations were optimized for comparison. Tetrahydrofuran (THF) is commonly used during the preparation of many clathrate hydrates to stabilize their structures.<sup>4,30</sup> The influence of THF was studied by placing each radical in the small cage and THF in the large one. In addition, we studied the stability of H· and OH· at different locations, which is useful for the understanding of their movements, and diffusion inside, and between cages. Harmonic vibrational frequencies were computed to ensure the located stationary structures are true minima on their respective potential energy surfaces. In the self-consistent field (SCF) calculations, the density tolerance was set to  $10^{-6}$  a.u. during geometry optimization. The resolution-of-identity (RI) approximation<sup>53,54</sup> and multipole accelerated RI (MARI) approach<sup>55</sup> were employed to speed up the computations.

## Results and discussion

### Structures of H· and OH· hydrates

The structures of the s1 and s2 water cages were optimized at the DFT-D3/def-TZVP level. The maximum and minimum values of the O–H bond lengths and hydrogen-bond lengths are listed in Table S5.<sup>†</sup> Using the B97D-D3 functional, for example, the O–H bond length, which is linked to other O atom by hydrogen bonding, is in the range 0.974–1.035 Å for s1 and 0.973–1.039 Å for s2. The O–H bond length, without hydrogen bonding, is about 0.966 Å for s1 and s2. In addition, the hydrogen-bond length is in the range 1.560–1.971 Å for s1 and 1.509–1.987 Å for s2. These results are similar to those of Chattaraj *et al.*,<sup>56</sup> and are listed for comparison in Table S5.<sup>†</sup>

The structures of the eight radical-included clathrate hydrates, with the radical either in the small or large cage for s1 or s2, were optimized at the unrestricted DFT-D3/def-TZVP level. Furthermore, to check for the spin contamination<sup>57</sup> in the open-shell species, the  $\langle S^2 \rangle$  values were determined and are listed in Table S6.<sup>†</sup> None of the clathrate hydrates are significantly affected by spin contamination (after allowing 10% variation<sup>58</sup>). Optimized structures are depicted in Fig. S1.<sup>†</sup> For the four H-included clathrate hydrates, the structures of the optimized water-cage frameworks are essentially the same as those of the cages devoid of the encapsulant. The inter- and intramolecular distances and angles are almost unchanged for the water molecules. H· always sits at around center of the cage, with distances of about 3.70–4.80 Å to the O atoms of the (cage) water molecules. Alavi *et al.* examined the movement of H· within  $5^{12}$  and  $5^{12}6^4$  water cages, and found that structures in which the H· was located at the center of the small or large cage were energetically more favorable than other conformers.<sup>59</sup> The structures of the cages of the four OH-included clathrate hydrates are similar to those of their H-included counterparts.

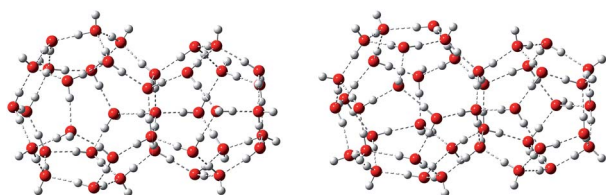


Fig. 1 Prototypical hydrate structures s1 (left) and s2 (right). O atoms are red, and H atoms are gray.



The encapsulation of OH· has little effect on the water cages, with the exception that one of the water molecules moves inward to form a hydrogen bond with OH·. Consequently, OH· is displaced about 1.5 Å from the center of the cage. The shortest distance between OH· and the surrounding water molecules is about 2.02–2.44 Å in the four hydrates. Only small changes in the O–H bond length are noted, about –0.005–0.006 Å, compared to free OH· in the gas phase.

Table 1 lists the stabilization energies (SEs) of the H· and OH·-included hydrates. For these systems, two kinds of SE can be defined. The first takes the stabilization energy among the water molecules into account, and it is computed by  $SE_a = E(\text{hydrate}) - E(\text{guest}) - N \times E(\text{water})$ , where  $E(X)$  is the total energy of the encapsulated hydrate, free guest radical, or free water molecule, respectively, and  $N$  is the number of water molecules. The second emphasizes the interaction between the guest and the water cage, and is computed by  $SE_b = E(\text{hydrate}) - E(\text{guest}) - E(\text{water cage})$ . All SE values have been subjected to basis set superposition error (BSSE) correction, using the counterpoise method of Boys and Bernardi.<sup>60</sup> In addition, all SE values have been zero-point energy (ZPE) corrected.

The  $SE_b$  values for H· in s1 and s2 are very small in magnitude. Positive values are predicted with UB97D-D3, while UB3LYP-D3, UBLYP-D3, and UPBE-D3 produce small negative  $SE_b$  values. For OH·, the computed  $SE_b$  values are negative using all four functionals. In addition, the  $SE_b$  values of OH· are larger than those of H·. However, for both H· and OH· radicals, there are no differences evident in the  $SE_b$  values between the s1 and s2 structures, or between the small and large cages. In contrast, the  $SE_a$  of H· and OH· in s1 and s2 are large and negative, and this can be attributed mainly to the stabilization energy of the water cage. The stabilities of the H· and OH·-included hydrates are reflected by their large  $SE_a$ , although their encapsulation leads to very small  $SE_b$  values. For comparison, the  $SE_a$  of empty s1 and s2 are also given in Table 1. It is apparent that more than 93% of the  $SE_a$  of H· and OH·, in s1 or s2, comes from the  $SE_a$  of the water molecules. Vikas

*et al.*<sup>61</sup> investigated the effect of D3, so the UB97D functional is further used, the results of which are listed in Table S7.† The UB97D results and UB97D-D3 are similar. Therefore, we can conclude that the water cages themselves help stabilize the encapsulation of H· and OH·, indicating that these water cages serve as stabilizers for these active radicals.

### Status of H· and OH· inside the water cages

We further studied the status of H· and OH· radicals in these water cages. Table 2 lists the net charges and spin densities of H· and OH·, which were evaluated by natural bond orbital (NBO) analyses.<sup>62</sup> The net charge on H· is very small, less than –0.014, regardless of the hydrate structure and cage size, indicating that little charge is exchanged between H· and the water cage. However, the net charge on OH· varies in the range of –0.123 to –0.165. Compared to the isolated OH·, the negative charge on the O atom increases, and the positive charge on the H atom also increases. The encapsulated OH· is clearly more polar than its isolated, gas-phase form. It is clear that very small amounts of charge flow from the water cage toward the OH·. The weak hydrogen bond between the OH· and the cage serves as a bridge for this flow of charge. Further analysis reveals that the transferred charge comes not only from the directly connected water molecule, but also from some of the other neighboring molecules, through the hydrogen bonding network, however, this contribution is rather small. In other words, the encapsulated H· and OH· retain almost the same charge states as their isolated forms.

The H· and OH· hydrate systems are found to be stable as doublet states. For the four H-containing hydrates, almost all (close to unity) of the spin density is located on the central H·, indicating that it maintains the spin state of its gas-phase form. On the other hand, the spin density of OH· lies in the range of 0.800–0.858, and a small part of the spin is distributed to the nearby water molecules. These spin distributions are also reflected in their electronic configurations, as shown in Table 3. The H· keeps its  $1s^{1.00}$  configuration in these cages, while the H atom in OH· is configured to be  $1s^{0.57-0.58}$ , close to that of free OH· ( $1s^{0.61}$ ). The O atom in OH· is configured to be  $2s^{1.88}2p^{4.64-4.69}3p^{0.00-0.01}$ , also close to that observed for free OH· ( $2s^{1.89}2p^{4.49}$ ). These spin density and electronic configuration analyses confirm that, when encapsulated, both H· and OH· retain most of the radical characteristics of their (free) gas-phase forms.

In this study, the influence of THF on the status of H· and OH· in these water cages was examined. THF normally occupies

**Table 1** Stabilization energies (SE, kcal mol<sup>–1</sup>) calculated for the s1 and s2 structures with H· or OH· occupied in the large (L,  $5^{12}6^2$  or  $5^{12}6^4$ ) or small (S,  $5^{12}$ ) cage

	$SE_a$		$SE_b$		
	UB97D-D3	UB97D-D3	UB3LYP-D3	UBLYP-D3	UPBE-D3
<b>H·</b>					
s1-L	–326.3	2.2	–1.1	–1.0	–0.9
s1-S	–325.2	3.8	–1.2	–1.3	–1.2
s2-L	–342.6	2.7	–0.9	–0.9	–0.9
s2-S	–341.5	4.3	–1.5	–1.4	–1.6
<b>OH·</b>					
s1-L	–332.0	–4.8	–5.1	–7.8	–10.7
s1-S	–333.5	–6.2	–7.9	–10.0	–10.4
s2-L	–349.4	–4.8	–6.0	–7.8	–8.0
s2-S	–352.4	–7.1	–7.1	–10.4	–10.6
s1	–328.6				
s2	–346.6				

**Table 2** Net charges and spin densities for H· and OH· in the large (L) or small (S) cage computed with the UB97D-D3 functional

	H·		OH·	
	Net charge	Spin density	Net charge	Spin density
s1-L	–0.006	0.987	–0.135	0.836
s1-S	–0.014	0.970	–0.132	0.857
s2-L	0	1.000	–0.123	0.858
s2-S	–0.013	0.971	–0.165	0.800



**Table 3** Configuration of H· and OH· in the large (L) or small (S) cage computed with the UB97D-D3 functional

	H·	OH·			
		H	O		
	1s	1s	2s	2p	3p
s1-L	1.00	0.59	1.88	4.65	0.01
s1-S	1.01	0.57	1.88	4.67	
s2-L	1.00	0.58	1.88	4.64	
s2-S	1.01	0.58	1.88	4.69	0.01
Gaseous	1.00	0.61	1.89	4.49	

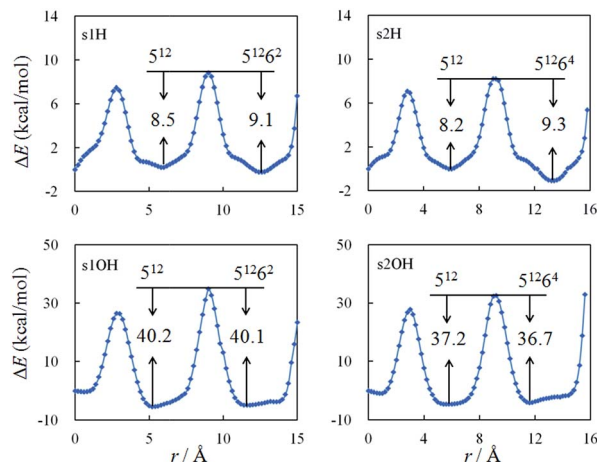
the larger cage of s1 and s2 because of its volume, while only the small cages are available for radicals. Using similar methods to those described above, we optimized structures in which H· or OH· occupies the small cage, while the large cage is occupied by THF, the results are given in Table 4. In all optimized structures THF is located at the center of the large cage with the shortest distance between its H atoms and the surrounding O atoms, in the s1 or s2 structure, calculated to be about 2.20 Å. In the presence of THF, very small variations in positions are noted for the H· and OH· radicals, compared to the structures devoid of THF. The H· is still located the center of the small cage for s1 and s2, and the average distance between H· and the surrounding water molecules is shorter than that without THF, by about 0.06 Å, while the shortest distance between OH· and the surrounding water molecules is shorter by about 0.1 Å. The changes in stabilization are less than 2.1 kcal mol<sup>-1</sup> for SE<sub>a</sub> and 22.2 kcal mol<sup>-1</sup> for SE<sub>b</sub>, regardless of the nature of the hydrate (s1 or s2). The corresponding changes in spin densities are less than 0.029 au for H· and 0.045 au for OH·. The net charges on H· and OH· remain almost the same, with corresponding changes of less than 0.001 for H· and 0.002 for OH·. In other words, although THF plays an important role in stabilizing these clathrate structures, it has little effect on the status of H· or OH· in the adjacent cage. Both H· and OH· are found to retain their radical characteristics in the presence of THF.

### Diffusion of H· and OH· radical between the water cages

When the radicals are confined in the clathrate structures, their movements are localized within the cages. We further studied

**Table 4** Stabilization energies (SE, kcal mol<sup>-1</sup>), net charges and spin densities of H· and OH· in the presence of THF encapsulated in the large cage of s1 and s2, calculated with the UB97D-D3 functional

	SE <sub>a</sub>	SE <sub>b</sub>	Net charge	Spin density
<b>H·</b>				
s1-S	-334.7	2.6	-0.013	0.972
s2-S	-364.8	2.4	-0.013	0.971
<b>OH·</b>				
s1-S	-341.5	-4.9	-0.131	0.858
s2-S	-368.6	-6.9	-0.163	0.813

**Fig. 2** Energy profiles for H· and OH· diffusion between cages in the s1 and s2 structures calculated with the UB97D-D3 functional.

the diffusion of H· and OH· radical between the water cages, as shown in Fig. S2.† In the s1 and s2 structures, the small (5<sup>12</sup>) and large (5<sup>12</sup>6<sup>2</sup> or 5<sup>12</sup>6<sup>4</sup>) cages share a pentagonal face. The energy profiles for radical diffusion between the cages are shown in Fig. 2, at different distances from the outer pentagonal face of the 5<sup>12</sup> cage, to the center of the 5<sup>12</sup> cage, to the center of the 5<sup>12</sup>6<sup>2</sup> or 5<sup>12</sup>6<sup>4</sup> cage, and finally to near the edge of the 5<sup>12</sup>6<sup>2</sup> or 5<sup>12</sup>6<sup>4</sup> cage.

The barriers for the diffusion of the H· radical from outside these hydrates through the pentagonal face of the 5<sup>12</sup> cage are calculated to be 8.5 and 8.3 kcal mol<sup>-1</sup> for the s1 and s2 structures, respectively, and they are essentially the same as the barriers for the reverse process, which are 8.5 and 8.2 kcal mol<sup>-1</sup> for the s1 and s2 structures, respectively. Alavi *et al.* studied the migration of H· through one 5<sup>12</sup> cage at the MP2 and B3LYP levels with the 6-311++G(*d,p*) basis set. One-dimensional barriers of 14.80 and 10.98 kcal mol<sup>-1</sup>, respectively, were calculated.<sup>59</sup> The barriers for OH· radical diffusion are calculated to be larger than those for H·, at 25.4 and 26.0 kcal mol<sup>-1</sup> for movement from outside the pentagonal face of each of the s1 and s2 structures, respectively.

From the center of the small cage to the center of the large cage in the s1 and s2 structures, the radicals migrate through another pentagonal face. Calculations predict that the barriers for diffusion from the center of the small cage, through the pentagonal face, to the center of the large cage are always similar to those for the reverse pathway. The differences are less than 1.1 kcal mol<sup>-1</sup>. Finally, as the radicals approach the furthest edges of these hydrates, the barriers increase enormously. This means that if a radical is confined by a water cage, it is very difficult for it to escape from the cage, or approach the cage edges.

### Stabilization mechanism of H· and OH· inside the water cages

Several authors have studied the reactions of H<sub>2</sub>O with H· or OH·.<sup>63–65</sup> These studies show that H· attacks the O atom in H<sub>2</sub>O





to form  $\text{H}_3\text{O}^+$ , with  $C_{3v}$  symmetry, while the O atom in  $\text{OH}^\cdot$  bonds to one of the H atoms in  $\text{H}_2\text{O}$ . In contrast to free  $\text{H}_2\text{O}$ , the water molecules in these cage structures are inactive toward the encapsulated  $\text{H}^\cdot$  or  $\text{OH}^\cdot$  radicals.

Fig. 3 depicts the interactions of  $\text{H}^\cdot$  and  $\text{OH}^\cdot$  radicals with the water cage. Firstly, one needs to note the specific structure that comes from water cage. The water molecules constitute a network in which the positively charged H atoms and the negatively charged O atoms are arranged alternatively with  $\angle\text{H}-\text{O}-\text{H} \approx 106^\circ$ ,  $\angle\text{O}-\text{H}\cdots\text{O} \approx 175^\circ$ , and  $\angle\text{H}-\text{O}\cdots\text{H} \approx 111^\circ$  along the surface, to simulate the arrangements of water molecules in cages. For  $\text{H}^\cdot$ , the angles  $\theta$  between the radical and O atom in water molecules 1, 2, and 3 are  $67^\circ$  without optimization, as shown in Fig. S3a.† The positions of water molecules 1, 2 and 3 are then adjusted, and angles  $\theta$  between  $\text{H}^\cdot$  radical and the O atoms of water molecules are respectively  $67^\circ$ ,  $72^\circ$ ,  $77^\circ$ ,  $82^\circ$ , and  $90^\circ$ . Obviously, if  $r$  and  $\theta$  are reduced, the  $\text{H}^\cdot$  becomes positioned closer to the water molecules. When  $\text{H}^\cdot$  approaches any of the atoms in an  $\text{H}_2\text{O}$  molecule, it is polarized by the charge of the interacting O or H atom, and an opposing charge is developed on the interacting side, which is however of the same sign as those of the atoms near the interacting atom. Coulombic repulsion from these atoms results in an increase in energy, and consequently, the  $\text{H}^\cdot$  is unable to approach the cage structure any closer. To further study this effect, energies were computed without optimization at CCSD(T)<sup>66</sup>/6-311++G(*d,p*)<sup>67</sup> and UB97D-D3/6-311++G(*d,p*) levels of theory, and the results of which are shown in Fig. S4,† both methods show the same

tendency in behavior. At the various angles, the energies are all calculated to increase as the distances,  $r$ , between the radical and the water molecules decrease. On the other hand, when  $r$  is fixed, energies are observed to increase with decreasing  $\theta$ . This means that if the  $\text{H}^\cdot$  is close to the water molecules, the overall energy increases.

Through localized molecular orbital energy decomposition analysis (LMOEDA),<sup>68</sup> implemented in GAMESS,<sup>69</sup> energy decomposition analyses was performed at CCSD(T)/6-311++G(*d,p*) level, in order to better understand why these energies increase, the results of this study are listed in Fig. S5.† The interaction energies were decomposed into electrostatic energy, exchange energy, polarization energy and repulsion energy. Except for repulsion energy, which increases with reduced  $r$ , the other three energy terms are all reduced with decreasing  $r$ . Moreover, the changes observed in the repulsion and exchange energies with  $r$  are larger than the changes in the electrostatic and polarization energies. Fig. 4a shows the total energy and energy decomposition analyses for  $\text{H}^\cdot$  at an angle of  $\theta = 67^\circ$ . When  $r$  is reduced, in other words  $\text{H}^\cdot$  is close to the water molecules, repulsion energies clearly become larger, while the exchange energies are decreased more than the electrostatic and polarization energies. The changes in interaction energy are led by the coulombic repulsion and exchange energies, with coulombic repulsion energy being the major factor for the observed increases in interaction energy.

Similar analyses can be made for the  $\text{OH}^\cdot$  radical, as depicted in Fig. 3c and S3c,† at  $\theta = 155^\circ$ ,  $160^\circ$ ,  $165^\circ$ ,  $170^\circ$ . Energies were also calculated at the CCSD(T)/6-311++G(*d,p*) and UB97D-D3/6-311++G(*d,p*) levels, the results of which are shown in Fig. S4.† At both levels, the approaching of  $\text{OH}^\cdot$  toward the cage results in increases in energy. LMOEDA analyses were also performed, as shown in Fig. S6† and 4b. Once again, the coulombic repulsion energy is a major factor for the observed increases in interaction energy.

As shown in Fig. S5 and S6,† repulsion energies increase with decreasing  $\theta$ . In the water cage, there are specific angles between the water molecules. As a result, the curvatures of the s1 and s2 water cages contribute to the increases in repulsion energies. Coulombic repulsion makes  $\text{H}^\cdot$  and  $\text{OH}^\cdot$  unable to approach the

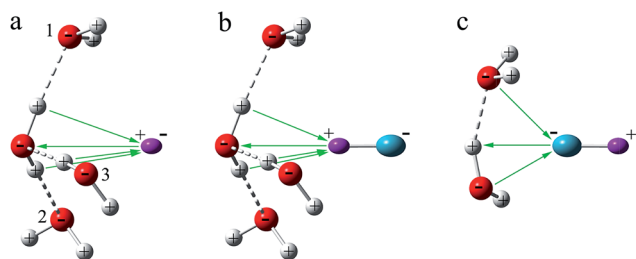


Fig. 3 Schematic diagram showing the interaction of  $\text{H}^\cdot$  and  $\text{OH}^\cdot$  with the water cage.  $\text{H}^\cdot$  and the H atom in the  $\text{OH}^\cdot$  radical are purple, the O atom in  $\text{OH}^\cdot$  is blue-green.

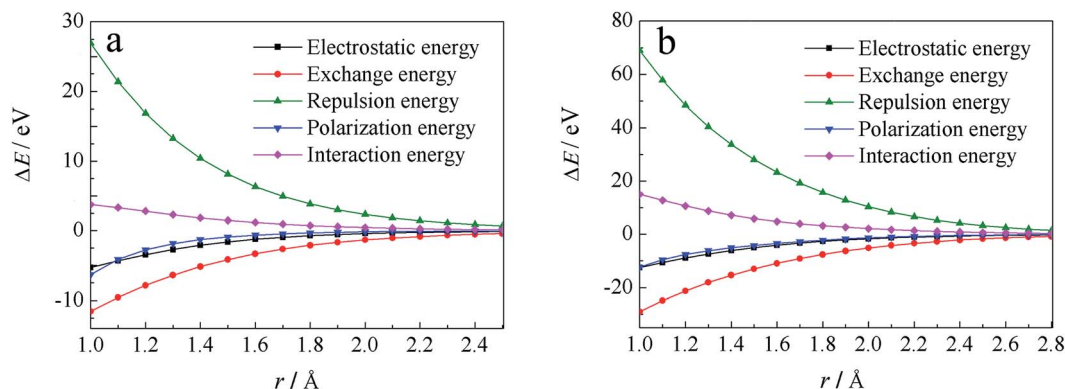


Fig. 4 Energy decomposition analysis for  $\text{H}^\cdot$  radical attack at  $\theta = 67^\circ$  (a) and  $\text{OH}^\cdot$  radical at  $\theta = 155^\circ$  (b) using LMOEDA analysis.



water cage edges, or react with water molecules, and this results in these radicals having relatively long survival times.

## Conclusions

By combining a D3 empirical dispersion correction to DFT functionals, we have modeled the confinement of hydrogen and hydroxyl radicals in typical water cages, s1 and s2, found in clathrate hydrates. After optimization, H $\cdot$  and OH $\cdot$  were found to be located near the center of the water cages, and these cages assist their encapsulation. Both radicals exhibit similar structures and properties to their corresponding free forms, including atomic charges, spin densities, and electronic configurations. Inter-cage diffusion studies reveal that energy barriers exist for these radicals to leave or approach the cage edges. LMOEDA energy decomposition analyses reveal that the changes in energy are mainly led by coulombic repulsion and exchange energies, with the coulombic repulsion energy between the radical and the water cage being the major factor for the observed increases in energy. The curvatures of the s1 and s2 water cages contribute to the increases in repulsion energies. Therefore, radicals are unable to approach or leave the water cage, and they cannot react with the water molecules of the cage. These studies provide insight into mechanisms for the prolonged capture of free radicals, with applications to synthetic and catalytic chemistry.

## Acknowledgements

We acknowledge the Education Committee of Sichuan Province (16ZB0067) for financial support.

## References

- 1 E. D. Sloan, *Energy Fuels*, 1998, **12**, 191–196.
- 2 W. L. Mao, H. Mao, A. F. Goncharov, V. V. Struzhkin, Q. Guo, J. Hu, J. Shu, R. J. Hemley, M. Somayazulu and Y. Zhao, *Science*, 2002, **297**, 2247–2249.
- 3 E. D. Sloan, *Nature*, 2003, **426**, 353–359.
- 4 L. J. Florusse, C. J. Peters, J. Schoonman, K. C. Hester, C. A. Koh, S. F. Dec, K. N. Marsh and E. D. Sloan, *Science*, 2004, **306**, 469–471.
- 5 H. Lu, Y. Seo, J. Lee, I. Moudrakovski, J. A. Ripmeester, N. R. Chapman, R. B. Coffin, G. Gardner and J. Pohlman, *Nature*, 2007, **445**, 303–306.
- 6 M. R. Walsh, C. A. Koh, E. D. Sloan, A. K. Sum and D. T. Wu, *Science*, 2009, **326**, 1095–1098.
- 7 A. Falenty, T. C. Hansen and W. F. Kuhs, *Nature*, 2014, **516**, 231–233.
- 8 Y. Huang, C. Zhu, L. Wang, X. Cao, Y. Su, X. Jiang, S. Meng, J. Zhao and X. C. Zeng, *Sci. Adv.*, 2016, **2**, e1501010.
- 9 M. Muraoka, N. Susuki and Y. Yamamoto, *RSC Adv.*, 2016, **6**, 63880–63885.
- 10 P. G. Debenedetti and S. Sarupria, *Science*, 2009, **326**, 1070–1071.
- 11 A. Boetius and E. Suess, *Chem. Geol.*, 2004, **205**, 291–310.
- 12 R. Boswell, *Science*, 2009, **325**, 957–958.
- 13 J. Zhu, S. Du, X. Yu, J. Zhang, H. Xu, S. C. Vogel, T. C. Gremann, J. S. Francisco, F. Izumi, K. Momma, Y. Kawamura, C. Jin and Y. Zhao, *Nat. Commun.*, 2014, **5**, 4128–4137.
- 14 C.-G. Xu and X.-S. Li, *RSC Adv.*, 2015, **5**, 54672–54699.
- 15 J. R. Cendagorta, A. Powers, T. J. H. Hele, O. Marsalek, Z. Bačić and M. E. Tukiman, *Phys. Chem. Chem. Phys.*, 2016, **18**, 32169–32177.
- 16 H. D. Nagashima, S. Takeya, T. Uchida and R. Ohmura, *Sci. Rep.*, 2016, **6**, 19345–19354.
- 17 S. Matsuishi, K. Hayashi, M. Hirano, I. Tanaka and H. Hosono, *J. Phys. Chem. B*, 2004, **108**, 18557–18568.
- 18 J. Yu, J. Chen, C. Li, X. Wang, B. Zhang and H. Ding, *J. Phys. Chem. B*, 2004, **108**, 2781–2783.
- 19 E. Carter, A. F. Carley and D. M. Murphy, *J. Phys. Chem. C*, 2007, **111**, 10630–10638.
- 20 C. R. Becer, R. Hoogenboom and U. S. Schubert, *Angew. Chem., Int. Ed.*, 2009, **48**, 4900–4908.
- 21 J. K. Nørskov, T. Bligaard, J. Rossmeisl and C. H. Christensen, *Nat. Chem.*, 2009, **1**, 37–46.
- 22 E. Cadenas and K. J. Davies, *Free Radical Biol. Med.*, 2000, **29**, 222–230.
- 23 S. Leon, *Nature*, 2009, **461**, 358–359.
- 24 M. Diehn, R. W. Cho, N. A. Lobo, T. Kalisky, M. J. Dorie, A. N. Kulp, D. Qian, J. S. Lam, L. E. Ailles, M. Wong, B. Joshua, M. J. Kaplan, I. Wapnir, F. M. Dirbas, G. Somlo, C. Garberoglio, B. Paz, J. Shen, S. K. Lau, S. R. Quake, J. M. Brown, I. L. Weissman and M. F. Clarke, *Nature*, 2009, **458**, 780–783.
- 25 A. T. Hoye, J. E. Davoren, P. Wipf, M. P. Fink and V. E. Kagan, *Acc. Chem. Res.*, 2008, **41**, 87–97.
- 26 W. Dröge, *Physiol. Rev.*, 2002, **82**, 47–95.
- 27 D. P. Jones, *Am. J. Physiol.: Cell Physiol.*, 2008, **295**, C849–C868.
- 28 M. Valko, D. Leibfritz, J. Moncol, M. T. D. Cronin, M. Mazur and J. Telser, *Int. J. Biochem. Cell Biol.*, 2007, **39**, 44–48.
- 29 K. Ohgaki, K. Nakatsuji, K. Takeya, A. Tani and T. Sugahara, *Phys. Chem. Chem. Phys.*, 2008, **10**, 80–82.
- 30 S.-H. Yeon, J. Seol, Y. Park, D.-Y. Koh, Y. S. Kang and H. Lee, *J. Am. Chem. Soc.*, 2008, **130**, 9208–9209.
- 31 M. Cha, K. Shin, M. Kwon, D.-Y. Koh, B. Sung and H. Lee, *J. Am. Chem. Soc.*, 2010, **132**, 3694–3696.
- 32 K. Yasuda, S. Takeya, M. Sakashita, H. Yamawaki and R. Ohmura, *J. Phys. Chem. C*, 2009, **113**, 12598–12601.
- 33 L. C. Jacobson, W. Hujo and V. Molinero, *J. Phys. Chem. B*, 2010, **114**, 13796–13807.
- 34 A. Khan, *J. Phys. Chem. A*, 2001, **105**, 7429–7434.
- 35 A. Khan, *J. Chem. Phys.*, 2002, **116**, 6628.
- 36 F. Lebsir, A. Bouyacoub, D. Bormann and A. Krallafa, *J. Mol. Struct.: THEOCHEM*, 2008, **864**, 42–49.
- 37 K. Riley, M. Pitoňák, P. Jurečka and P. Hobza, *Chem. Rev.*, 2010, **110**, 5023–5063.
- 38 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.
- 39 D. G. A. Smith, L. A. Burns, K. Patkowski and C. D. Sherrill, *J. Phys. Chem. Lett.*, 2016, **7**, 2197–2203.
- 40 P. Ghosh and P. Banerjee, *Phys. Chem. Chem. Phys.*, 2016, **18**, 22805–22815.



- 41 Z. Jian, X. Yang, D. Chen and Y. Qian, *RSC Adv.*, 2016, **6**, 114560–114565.
- 42 J. Vatamanu and P. G. Kusalik, *J. Am. Chem. Soc.*, 2006, **128**, 15588–15589.
- 43 S. Takeya, K. A. Udachin, I. L. Moudrakovski, R. Susilo and J. A. Ripmeester, *J. Am. Chem. Soc.*, 2010, **132**, 524–531.
- 44 J. Vatamanu and P. G. Kusalik, *J. Phys. Chem. B*, 2006, **110**, 15896–15904.
- 45 D. A. G. Gualdrón and P. B. Balbuena, *J. Phys. Chem. C*, 2007, **111**, 15554–15564.
- 46 N. I. Papadimitriou, I. N. Tsimpanogiannis, A. T. Papaioannou and A. K. Stubos, *J. Phys. Chem. C*, 2008, **112**, 10294–10302.
- 47 S. Grimme, S. Ehrlich and L. Goerigk, *J. Comput. Chem.*, 2011, **32**, 1456–1465.
- 48 T. Schwabe and S. Grimme, *Phys. Chem. Chem. Phys.*, 2007, **9**, 3397–3406.
- 49 J. J. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1997, **78**, 1396.
- 50 F. Weigend, M. Haser, H. Patzelt and R. Ahlrichs, *Chem. Phys. Lett.*, 1998, **294**, 143–152.
- 51 *TURBOMOLE V5-9-1*, University of Karlsruhe, 2007.
- 52 M. Head-Gordon, J. A. Pople and M. J. Frisch, *Chem. Phys. Lett.*, 1988, **153**, 503–506.
- 53 K. Eichkorn, O. Treutler, H. Oehm, M. Haeser and R. Ahlrichs, *Chem. Phys. Lett.*, 1995, **242**, 652–660.
- 54 K. Eichkorn, F. Weigend, O. Treutler and R. Ahlrichs, *Theor. Chem. Acc.*, 1997, **97**, 119–124.
- 55 M. Sierka, A. Hogekamp and R. Ahlrichs, *J. Chem. Phys.*, 2003, **118**, 9136.
- 56 P. K. Chattaraj, S. Bandaru and S. Mondal, *J. Phys. Chem. A*, 2011, **115**, 187–193.
- 57 M. W. Wong and L. Radom, *J. Phys. Chem.*, 1995, **99**, 8582–8588.
- 58 R. Kaur and Vikas, *RSC Adv.*, 2016, **6**, 29080–29098.
- 59 S. Alavi and J. A. Ripmeester, *Chem. Phys. Lett.*, 2009, **479**, 234–237.
- 60 S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553–566.
- 61 P. Sangwan and Vikas, *J. Chem. Phys.*, 2016, **144**, 044305.
- 62 J. P. Foster and F. Weinhold, *J. Am. Chem. Soc.*, 1980, **102**, 7211–7218.
- 63 D. H. Zhang, M. A. Collins and S.-Y. Lee, *Science*, 2000, **290**, 961–963.
- 64 M. Brouard, P. O'Keeffe and C. Vallance, *J. Phys. Chem. A*, 2002, **106**, 3629–3641.
- 65 T. Uchimaru, A. K. Chandra, S. Tsuzuki, M. Sugie and A. Sekiya, *J. Comput. Chem.*, 2003, **24**, 1538–1548.
- 66 J. A. Pople, M. Head-Gordon and K. Raghavachari, *J. Chem. Phys.*, 1987, **87**, 5968–5975.
- 67 K. Raghavachari, J. S. Binkley, R. Seeger and J. A. Pople, *J. Chem. Phys.*, 1980, **72**, 650–654.
- 68 P. F. Su and H. Li, *J. Chem. Phys.*, 2009, **131**, 014102–014115.
- 69 M. W. Schmidt, K. K. Baldridge, J. A. Boatz, S. T. Elbert, M. S. Gordon, J. H. Jensen, S. Koseki, N. Matsunaga, K. A. Nguyen and S. Su, *J. Comput. Chem.*, 1993, **14**, 1347–1363.

