

The mechanism of hydrogen donation by bio-acids over metal supported on nitrogen-doped carbon nanotubes

Jiajun Zhang^{a,b}, Xiaolei Zhang^{a,*}, Amin Osatiashtiani^c, Kai Hong Luo^d, Dekui Shen^e, Jun Li^a, Anthony Bridgwater^c

^a Department of Chemical and Process Engineering, University of Strathclyde, Glasgow, United Kingdom

^b Center for Combustion Energy, Key Laboratory for Thermal Science and Power Engineering of Ministry of Education, Department of Energy and Power Engineering, Tsinghua University, Beijing 100084, China

^c Energy & Bioproducts Research Institute (EBRI), School of Engineering and Applied Science, Aston University, Aston Triangle, Birmingham, B4 7ET, United Kingdom

^d Department of Mechanical Engineering, University College London, Torrington Place, London, WC1E 7JE, UK

^e Key Laboratory of Energy Thermal Conversion and Control of Ministry of Education, Southeast University, Nanjing, China

ARTICLE INFO

Keywords:

Acetic acid
Formic acid
Carbon nanotubes
Molybdenum
Biomass

ABSTRACT

Biomass-derived carboxylic acids (e.g. acetic acid AcOH and formic acid FA) are a green and low-cost hydrogen source to replace hazardous H₂ gas in in-situ hydrogenation processes. To date, bio-acids dehydrogenation has been mainly conducted using noble metal catalysts which would negatively impact the process economy, thus development of efficient non-noble metal catalysts for this purpose is highly desirable. In this study, the performance of transition metals supported on nitrogen doped carbon nanotubes was thoroughly evaluated by computational modelling based on Density Functional Theory (DFT). Results revealed that, out of the 10 selected transition metal candidates, molybdenum (Mo) was most active for binding AcOH and a combination of Mo and nitrogen doping significantly enhanced binding to the carboxylic acid molecules compared to pristine carbon nanotubes (CNTs). The newly designed Mo/N-CNT catalysts considerably facilitated the bio-acids decomposition compared to the non-catalytic scenarios by lowering energy barriers. FA distinctly outperformed AcOH in hydrogen donation over Mo/N-CNT catalysts, through its spontaneous cleavage leading to facile hydrogen donation.

1. Introduction

The rapid economic growth and increase in the global population have accelerated the consumption of fossil resources. This has resulted in an unprecedented increase in the level of CO₂ and other greenhouse gases (GHG) emissions, threatening the future of our planet by contributing to global warming [1]. Therefore, alternative sustainable feedstocks are needed to meet the demands for organic chemicals and fuels.

Non-edible and waste biomass such as lignocellulose and triglycerides are considered a sustainable source of carbon that can be used to produce green and low-cost chemicals and fuels. However, lignocellulosic biomass has high oxygen content, thus itself or its derivative has to undergo some deoxygenation steps to be converted to the desired products [2]. Conventionally, deoxygenation is conducted via hydrogenation using precious metal-based catalysts and molecular H₂.

However, this approach can have negative impact on process economy and safety, owing to the high cost of precious metals (e.g. Pt, Pd, and Ru) and the need for high H₂ pressure (> 30 bar) [3]. Therefore, alternative sustainable hydrogen sources, as well as the efficient catalysts to maximise the energy efficiency of the hydrogen transfer process are sought after.

Various alternative organic hydrogen donors have been reviewed in detail in our previous work [4]. Among them, bio-acids are promising, not only as hydrogen donors, but also as potential chemicals for hydrogen storage [5]. The research on bio-acids decomposition for in-situ hydrogen generation has been focused mainly on acetic acid (AcOH) and formic acid (FA) [6–8], because they are abundant in bio-oil and more readily decomposed compared to other bio-based carboxylic acids [9,10]. Jiang et al. [6] predicted the pathway of FA decomposition over Cu (111) as HCOOH → HCO + OH → H + CO + OH. Li et al. [11] reported the decomposition mechanism of AcOH over Co (111) stepped

* Corresponding author.

E-mail address: xiaolei.zhang@strath.ac.uk (X. Zhang).

<https://doi.org/10.1016/j.mcat.2020.111289>

Received 6 July 2020; Received in revised form 15 October 2020; Accepted 21 October 2020

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surface, suggesting that the reaction pathway follows $\text{CH}_3\text{COOH} \rightarrow \text{CH}_3\text{CO} + \text{OH} \rightarrow \text{CH}_2\text{CO} + \text{H} + \text{OH} \rightarrow \text{CH}_2 + \text{CO} + \text{H} + \text{OH} \rightarrow \text{CH} + \text{CO} + 2\text{H} + \text{OH}$. There are other different reaction mechanisms over noble metals, producing CH_4 and CO from AcOH decomposition, and CO_2 and H_2 from FA decomposition, respectively [7,12,13]. Also, our previous DFT study demonstrated that Mo (110) exhibited a great potential as a catalyst for the dehydrogenation of FA and AcOH , due to their exothermic property and lower energy barriers during the decomposition [4]. Nevertheless, unsupported metal catalysts are rarely used in practice because metal clusters are usually needed to be evenly dispersed on a high surface area support to maximise the interface between the reactants and the catalytic sites. Furthermore, the interaction between the support and the metal can have significant impact on the activity and selectivity of the catalytic system, therefore selection of an appropriate support and understanding the metal-support interactions are crucial to the smart catalyst design [14].

Several studies have reported the decomposition of bio-acids over heterogeneous catalysts with a variety of supports including metal oxides [15,16], zeolites [17], as well as carbon based materials [18,19]. Among the various types of supports, carbon nanotubes (CNTs) are very interesting materials, owing to their excellent thermal stability (up to 3000 K), carbonaceous deposit resistance, as well as large specific surface area (over 1000 m^2/g) which allow a higher metal dispersion compared to other supports [20,21]. Additionally, CNTs can be prepared using residual biomass, contributing to the reduction in carbon footprint [22–24]. CNTs functionalised with transition metals [25], defects and heteroatoms [26,27] have demonstrated to be promising catalysts for redox reactions by achieving high conversion rates and selectivity to the desired products [28]. For example, N-doped CNTs have shown remarkable activities in redox reaction by tailoring the local electron property [29]. Metal doped CNTs have also shown promising results in the hydrodeoxygenation of anisole [30], Aerobic oxidation of 5-hydroxymethylfurfural to 2,5-furandicarboxylic acid [31], and transformation of furfural to cyclopentanone, as well as in-situ hydrogen generation via decomposition of bio-acids [32]. Ding et al. [33] reported the decomposition of FA into CO_2 and H_2 over Pd/CNTs at room temperature with a high turnover frequency (TOF) of 1135 h^{-1} . Nabid and co-workers [34] synthesized Ag core Pd shell nanoparticles supported on CNTs without any additives and achieved FA dehydrogenation with an overall activation energy of $28.28 \text{ kJ mol}^{-1}$. Bi et al. [35] modified CNTs by N doping, achieving $\text{TOF} > 5000 \text{ h}^{-1}$ and near complete conversion of FA .

In a N-CNT support, four types of nitrogen atoms could be present (as illustrated in Fig. 1); 1) graphitic N (N_g -CNTs), 2) (vacancy) pyridinic N (referred as pyridinic nitrogen in this work, N_p -CNTs), 3) pyrrolic N, and 4) pyridinic N-O. Previous investigations into N-CNTs indicate that N_g -CNTs and N_p -CNTs are the most common types of N dopants in CNTs [36,37]. Pyridinic N atoms (N_p) are normally located at the edges or in the vacancy of the CNT surface and are bonded to two carbon atoms, and

the three sp^2 orbitals are occupied by four electrons, where two electrons are used to form two δ bonds with carbon atoms, and the other two hybridized electrons form a lone pair. The remaining electron in N_p joins the adjacent carbon atoms to form a π bond. The lone electron pair would endow the N_p -CNTs with Lewis basicity, while the π electron acts as an electron acceptor [38–40]. N_g has similar sp^2 hybridisation as N_p ; four orbitals are occupied by only four electrons to form three δ bonds and one π , and the fifth electron is not accommodated in a bonding orbital, because it is in a higher energy state and delocalised from the N site, functioning as an electron donor [29,41,42].

Due to the different basicity and acidity, each of the various doping types of N in CNTs promotes different catalytic reactions [44,45]. Although experimental data suggest that functionalised CNTs are promising catalysts for in-situ hydrogen generation from the decomposition of bio-acids [33–35], the literature on the reaction mechanism is very limited. Particularly, the interactions between each functional component of the CNTs catalyst composite have rarely been explored.

In this study, the catalytic decomposition of AcOH and FA for hydrogen production over the metal supported on N-CNTs was investigated with a focus on the cleavage of hydrogen related bonds. Ten transition metals including Ni , Mo , Fe , Co , Pt , Rh , Ru , Zn , Cu , and Pd , as well as six types of CNT-based catalyst supports (5 functionalised CNT and the pristine CNT) were compared with respect to their binding energy in adsorbing AcOH . AcOH and FA were then evaluated in terms of their adsorption process and reactivity in dehydrogenation over the Metal/N-CNT catalysts, and the cleavage of the corresponding bonds was tracked throughout the decomposition reaction.

2. Methodology

The first-principle density functional theory with dispersion correction (DFT-D) calculations was implemented in CASTEP, and models were established in Materials Studio 2017 R2 from BIOVIA [46,47]. The generalised gradient corrected approximation (GGA) [48] treated by the Perdew–Burke–Ernzerhof (PBE) exchange-correlation potential with long-range dispersion correction via Grimme’s scheme was used to calculate the exchange-correlation energy [49]. The on-the-fly generated (OTFG) ultra-soft pseudopotential was employed as the scheme in the representation of reciprocal space for all the elements [50,51]. The plane-wave cut-off energy was set to 600 eV for all the calculations based on its independence test (Fig. S1a). The Brillouin zone was sampled using a $2 \times 1 \times 4$ Monkhorst-Pack k-point (spacing of 0.03 \AA^{-1}) with a smearing of 0.1 eV, based on its independence test (Fig. S1b). The self-consistent field (SCF) tolerance was set to 10^{-6} eV/atom. The entire calculation was performed with a convergence threshold of 10^{-5} eV/atom on energy, 0.03 eV/Å on maximum force, and 10^{-3} Å on the maximum displacement. No symmetry constraint was used for any modelling.

All the models were based on a two-unit supercell of achiral zigzag ($n = 12$, $m = 0$, $\Theta = 0^\circ$) form single wall CNT (SWCNT) with a diameter of 9.5 Å. A 15 Å vacuum region was created above the top of the CNT. Based on the tests with larger supercell models, the size of CNT model used in this study would be adequate to get reliable computation results (Fig. S2). Geometry optimisation was implemented to every model before energy was calculated. The transition state (TS) was completely determined by the LST/QST method, and the TSs for the dominant reaction steps were confirmed by the unique imaginary frequency (Table S2). Hirshfeld charge was calculated for atomic electron analysis. The adsorption energy (E_{ad}) was determined by Eq. 1, where E_{catalyst} , $E_{\text{adsorbate}}$ and $E_{\text{adsorbate/catalyst}}$ are the total energies of clean catalyst, free adsorbate molecule and catalyst with adsorbed molecule, respectively. The energy barriers of reactions (E_{barrier}) were determined by the difference between the energies of transition state and reactant, as shown in Eq. 2, where $E_{\text{transition state}}$ and E_{reactant} are the total energies of the transition state and reactant of a reaction, respectively.

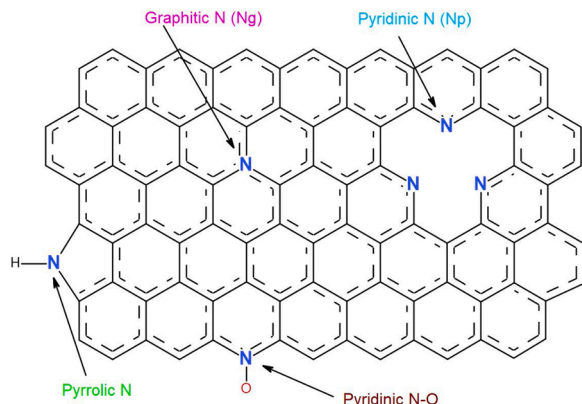


Fig. 1. Common types of N doping in carbon matrices [37,43].

$$E_{\text{ad}} = E_{\text{adsorbate/catalyst}} - (E_{\text{catalyst}} + E_{\text{adsorbate}}) \quad (1)$$

$$E_{\text{barrier}} = E_{\text{transition state}} - E_{\text{reactant}} \quad (2)$$

The bond dissociation energies (BDEs) were determined by Eq. 3, where E_{molecule} is the molecule energy, and $E_{\text{fragments}}$ is the energy summation of each decomposed fragments from bond cleavage.

$$\text{BDE} = E_{\text{molecule}} - E_{\text{fragments}} \quad (3)$$

The electron density difference (EDD) was determined by Eq. 4, where $\rho_{\text{adsorbate@cata}}$ is the electron density of the whole system of adsorbate and catalyst, and $\rho_{\text{adsorbate}}$ and ρ_{cata} are the unperturbed electron densities of the adsorbate and the catalyst structure, respectively.

$$\Delta\rho = \rho_{\text{adsorbate@cata}} - (\rho_{\text{adsorbate}} + \rho_{\text{cata}}) \quad (4)$$

3. Results and discussion

3.1. Impact of metal and nitrogen type on acetic acid adsorption energy

To evaluate the performance of various transition metals and N-CNT catalyst supports, the adsorption energy of AcOH on 10 N_p -CNT supported transition metals as well as the corresponding unsupported metals were compared, as shown in Figs. 2 and S3. The results reveal that the adsorption energy of AcOH on the supported metals is generally larger than that of the unsupported metal facets [4], indicating the enhanced interactions between AcOH and the CNT supported catalysts. Among the selected materials, Mo and Pt supported on N_p -CNT exhibited the strongest AcOH binding energy (1.76 eV), implying that they are probably more catalytically active than other transition metals selected in activating carboxylic acids. Since precious metal catalysts were avoided in this study, Mo was selected for further investigation.

The structure of the aforementioned two common types of N-CNT models (N_p -CNT and N_g -CNT) supported Mo catalysts was established and demonstrated in Fig. 3, along with the pristine CNT model. The impact of different N chemical environments on the adsorption energy of AcOH on the metal-doped and metal-free CNT was compared, and the results are summarised in Table 1.

Results showed that the pristine CNT as the catalyst led to the lowest adsorption energy (close to physical adsorption, distance larger than 3

Å) of AcOH. N-CNTs gave rise to slightly larger adsorption energy, however, no obvious difference was observed between N_p and N_g types; the distances between the molecule and catalysts for the above cases were all larger than 2.7 Å (Fig. S4). The modelling results reveal that the adsorption energy of AcOH significantly increased when Mo was present on the CNT surface, with simultaneous shorter distances (around 2 Å) between the molecule and catalysts, confirming the binding promotion effect of Mo on the AcOH adsorption, in agreement with the literature [4]. Over Mo/ N_p -CNT, AcOH exhibited the highest binding energy (-1.76 eV) than that of Mo/ N_g -CNT and Mo/CNT, ascribed to the unpaired electron of pyridinic N which acts as electron acceptor (Lewis acid), reinforcing the binding between the catalyst and the adsorbate molecule [52]. This also explains the slightly lower adsorption energy of AcOH over Mo/CNT. Compared to Mo/ N_p -CNT and Mo/CNT, the adsorption energy was obviously decreased by Mo/ N_g -CNT, indicating the weaker binding between the AcOH molecule and Mo/ N_g -CNT, resulting from the aforementioned delocalized electron of N_g [29]. It is also found that N_p -CNT and N_g -CNT exhibit different acid and base effects on AcOH adsorption only in the presence of Mo, which can be attributed to the synergistic effects between N and Mo atoms, in line with the reported results in the literature [40,53]. This difference in terms of acid and base effects may also endow the catalysts with different redox properties. It is thus hypothesised that the two types of N doping would lead to different catalytic performance of Mo/N-CNTs in the decomposition of bio-acids. Besides the adsorption performance, the stability of these two catalysts were tested (Fig. S5). Results show that Mo/ N_p -CNT had higher thermal stability, and thus could operate in high temperatures up to 1000 K with little distortion, whereas Mo/ N_g -CNT is suitable for low temperature operation scenarios.

3.2. Dehydrogenation of acetic acid over Mo/N-CNTs

Different adsorption energies of AcOH in various configurations (Table S1) over the two types of catalysts of Mo/ N_p -CNT and Mo/ N_g -CNT are compared in Table 2. The bond dissociation energies (BDEs) of each bond in free molecule are shown in Fig. 4 for comparison. In both cases, AcOH bonded onto Mo by chemical adsorption through C_β -Mo, O_α -Mo, and O_β -Mo, and by weaker binding through C_α -Mo, where Mo/ N_p -CNT resulted in higher binding energy than Mo/ N_g -CNT in all strong adsorption circumstances. Interestingly, dissociative adsorption was observed when AcOH bonded to the Mo/ N_p -CNT through O_β -Mo, where the molecule decomposed into hydroxyl and acetyl (with the C—O bond distance of 2.7 Å) during the adsorption. The dissociative adsorption is

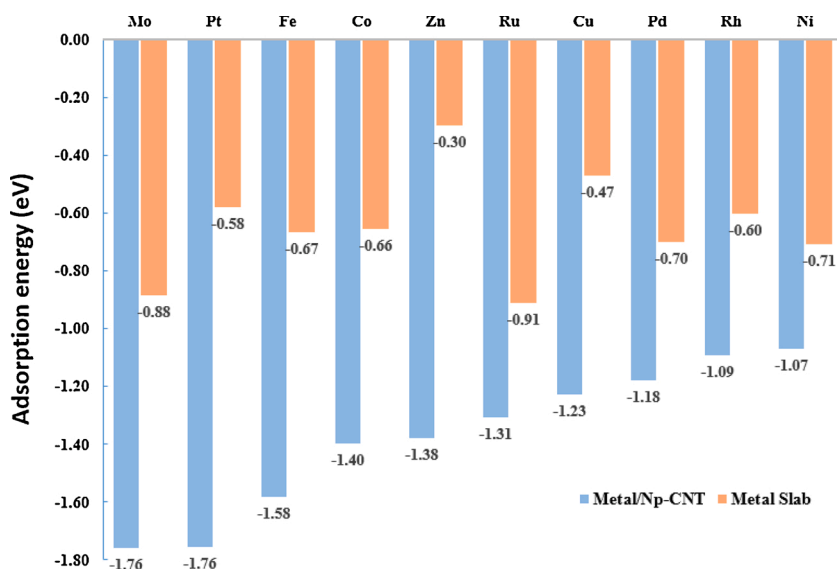


Fig. 2. Adsorption energy of acetic acid over metal/ N_p -CNTs (from this work) and pure metal facets (from our previous work [4]).

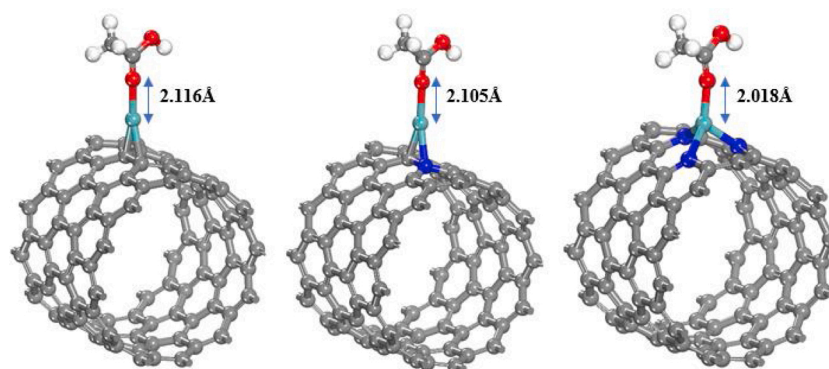


Fig. 3. Models of (a) Mo/CNT (b) Mo/N_g-CNT (c) Mo/N_p-CNT (grey denotes carbon, green denotes Mo, blue denotes nitrogen, red denotes oxygen and white denotes hydrogen).

Table 1

Adsorption energy of AcOH on (functionalised) CNTs.

Active site (catalyst)	Mo/N _p -CNT	Mo/N _g -CNT	Mo/CNT	N _p -CNT	N _g -CNT	CNT
E _{ad} (AcOH) (eV)	-1.76	-1.61	-1.72	-0.73*	-0.73*	-0.63*

* Only stable weak adsorption observed, and the corresponding configurations are shown in Fig. S4.

Table 2

Adsorption energy of AcOH on Mo/N-CNTs with different configurations.

E _{ad} (AcOH) (eV)	C _α -Mo	C _β -Mo	O _α -Mo	O _β -Mo
Mo/N _p -CNT	-0.58*	-2.41	-1.76	-2.61
Mo/N _g -CNT	-0.75*	-1.99	-1.61	-2.11

* Only stable weak adsorption observed.

common for molecular hydrogen adsorption over transition metals [54], however the dissociative AcOH adsorption is observed for the first time. The dissociative adsorption of AcOH simultaneously led to the largest adsorption energy of -2.61 eV over Mo/N_p-CNT through O_β-Mo bond. The same O_β-Mo bond also led to the largest adsorption energy of -2.11 eV when AcOH molecule adsorbed on Mo/N_g-CNT. The difference between the adsorption energies of AcOH on Mo doped N_g- and N_p-CNT catalysts is associated with their electron donation and acceptance properties respectively, where N_p-CNT with higher acidity shows stronger binding to AcOH.

The most favourable reaction pathways of AcOH decomposition over Mo/N_p-CNT are shown in Fig. 5, which were determined based on the cleavage energy calculation for each bond inside the acid molecule, as well as inside the lower fragment structures (the lowest energy barrier values among all parallel bonds cleavage are shown, and the details for other parallel reactions are shown in Table S3). The modelling results predict that the decomposition of AcOH started with the dissociative adsorption, producing hydroxyl and acetyl fragments. Subsequently, acetyl cracked into CO and methyl fragments with a small energy barrier of 0.32 eV and reaction energy of -0.68 eV. Further decomposition of methyl released three H atoms in sequence with the energy barriers of 1.74 eV, 1.47 eV and 2.54 eV, respectively. Another H cleaved from the hydroxyl with an energy barrier of 2.03 eV. It is found that over Mo/N_p-CNT, the majority of elementary steps amid the favourable reaction pathway of AcOH decomposition are endothermic, except the thermodynamically favoured C-C and initial C-O cleavage.

AcOH did not show dissociative adsorption over Mo/N_g-CNT (Fig. 6 and Table S1), instead the adsorbed structure underwent facile decomposition to produce acetyl and hydroxyl with a small energy barrier of 0.21 eV (Table S3), with an exothermic reaction energy of -0.77 eV. The

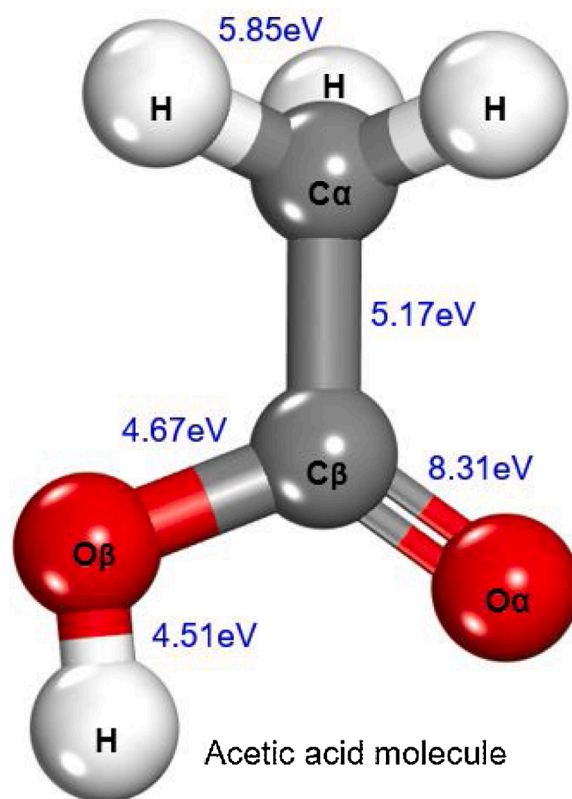


Fig. 4. Dissociation energy for each bond (blue font) in acetic acid free molecule [4].

modeling shows that the C-C bond strength in the adsorbed acetyl group was reinforced over Mo/N_g-CNT, with a shorter bond length of 1.49 Å compared to 1.53 Å over Mo/N_p-CNT and 1.51 Å in free molecule status [4,55], and this change would be analysed in detail in Section 3.4. As a result, the C=O bond cleaved firstly, retaining the structure of CH₃-C. CH₃-C would then decompose into methyl fragment and carbon atoms with a large energy barrier of 2.47 eV. The cleavage of C=O prior to C-C was also observed in the decomposition of AcOH over a Mo slab in our previous research [4]. The release of H atoms from methyl over Mo/N_g-CNT had the energy barriers of 0.36 eV, 0.72 eV and 3.74 eV, respectively. Besides, the cleavage of hydroxyl over Mo/N_g-CNT required additional energy of 1.58 eV. The cleaved H atoms from the decomposition process of acid molecules would stay on the metal temporarily before they continue to migrate or spill over in real hydrogenation circumstances [56].

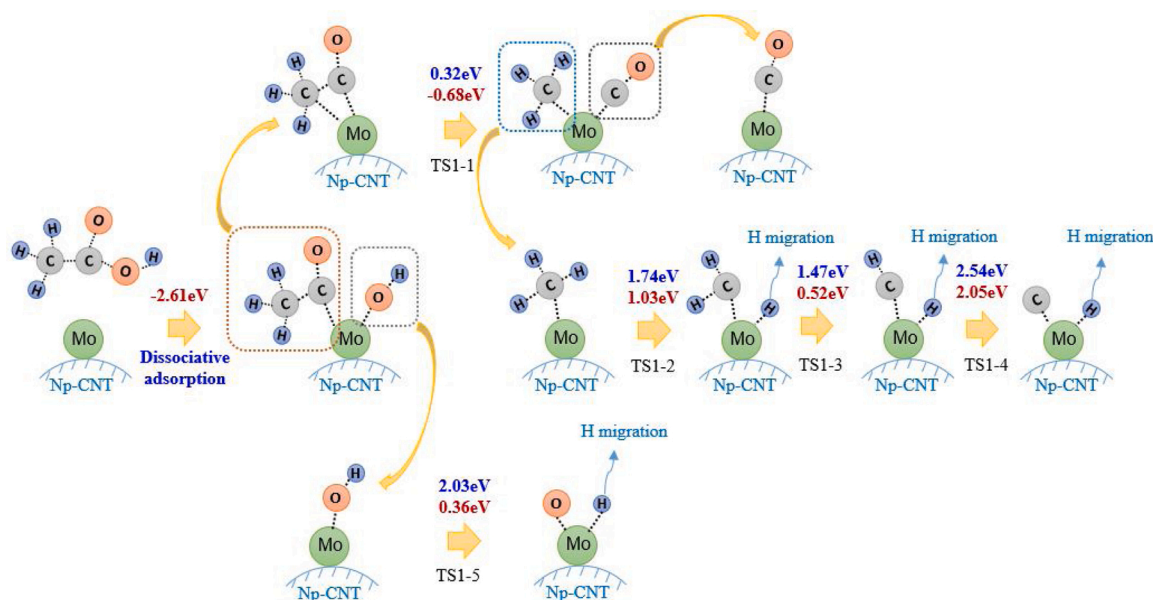


Fig. 5. Reaction flow of the decomposition of AcOH over Mo/N_p-CNT (Blue font denotes energy barrier and violet denotes reaction energy).

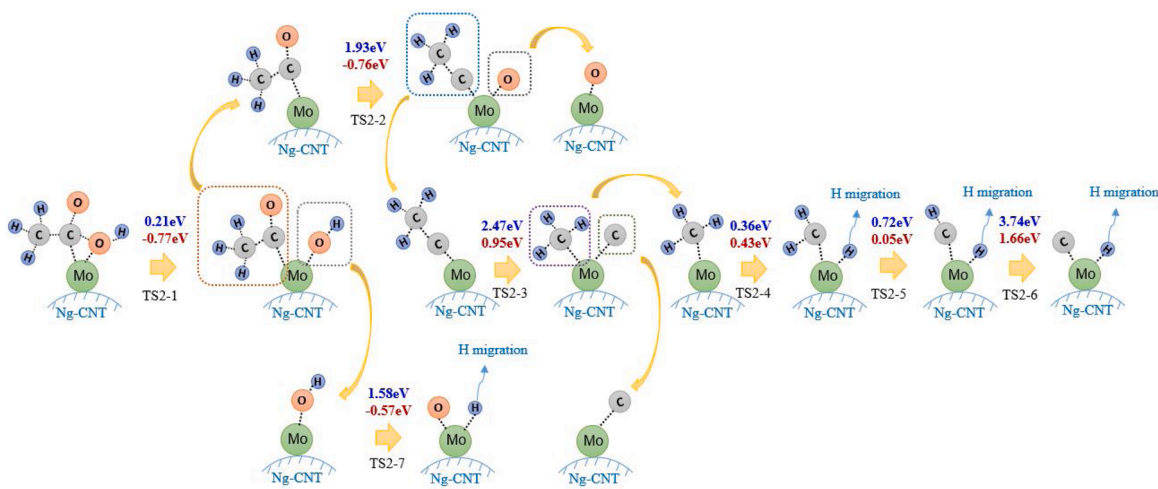


Fig. 6. Reaction flow of the decomposition of AcOH over Mo/N_g-CNT (Blue font denotes energy barrier and violet denotes reaction energy).

In comparison with the free molecular decomposition of AcOH [4], shown in Fig. 4, the cleavage energy of each bond in AcOH has been significantly decreased, suggesting that both Mo/N-CNT catalysts facilitate the decomposition of AcOH. The modelling results predict hydroxyl cleavage occurred at the beginning of the AcOH decomposition over both types of Mo/N-CNTs, which is different from the pathways over pure metal facets e.g. Mo and Rh [4,57]. Specifically, the formation of acetate is suppressed at the beginning of reaction, and the donation of methyl hydrogens occurs as final steps over both types of Mo/N-CNT.

Driven by its electrophilicity (electron acceptor) property, Mo/N_p-CNT gives rise to a barrier-free dissociation during the AcOH adsorption, whereas Mo/N_g-CNT leads to a small energy barrier for the same decomposition. The modelling predicts different favourable pathways for AcOH decomposition over the two Mo/N-CNTs catalysts. Mo/N_p-CNT favours the C—C cleavage with an energy barrier of 0.32 eV, which is significantly lower than that in the reaction over a pure Mo slab with an energy barrier of 4.22 eV. This is in perfect agreement with a commonly reported favourable reaction pathway for AcOH decomposition [10,14,58], with the merit of few hurdles before the hydrogen donation, while the release of the four hydrogens from the lower

fragments of methyl and hydroxyl are mostly endothermic processes with higher energy barriers. This may ease CH₄ and CO formation during the hydrogen donation process [10,59]. In contrast, the release of methyl H and the hydroxyl H show smaller energy barriers over Mo/N_g-CNT than over Mo/N_p-CNT, indicating the dedicated function of Mo/N_g-CNT in facilitating the dissociation of the C—H bond. However, the hydrogen donation has to first overcome the early stage decomposition steps with much higher energy barriers such as C—O_α and C—C cleavages.

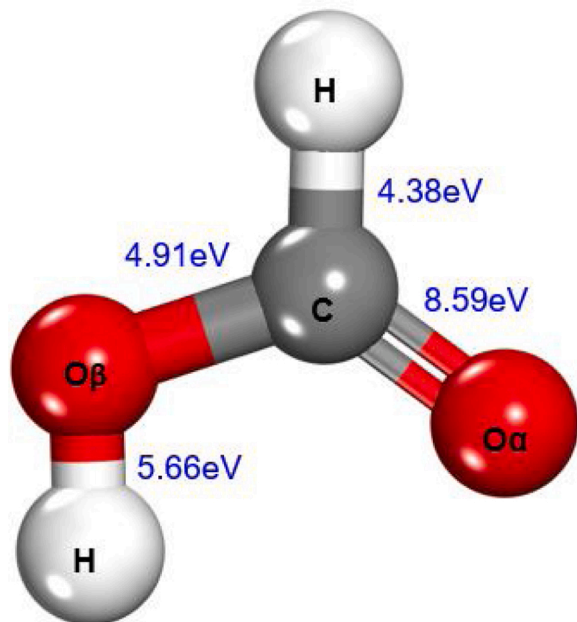
3.3. Dehydrogenation of formic acid over Mo/N-CNTs

The adsorption of FA over Mo/N_p-CNT and Mo/N_g-CNT in a variety of configurations was modelled and the adsorption energies are compared in Table 3. The corresponding configurations are shown in Table S1. The free molecular structure of FA and the associated BDEs of each bonds are shown in Fig. 7. The results reveal that the FA molecule binding to the catalyst via a C—Mo bond led to the largest adsorption energies, for -3.54 eV over Mo/N_p-CNT and -2.75 eV Mo/N_g-CNT, respectively, followed by the binding through the O_α—Mo bond over

Table 3
Adsorption energy of FA onto Mo/N-CNTs with different configurations.

E _{ad} (FA) (eV)	C-Mo	O _α -Mo	O _β -Mo
Mo/N _p -CNT	-3.54	-1.74	-0.81*
Mo/N _g -CNT	-2.75	-1.55	-1.15

* Only stable weak adsorption observed.



Formic acid molecule

Fig. 7. FA molecule with cleavage energy barriers (blue font) in free molecule [4].

Mo/N_p-CNT (-1.75 eV) and Mo/N_g-CNT (-1.55 eV). FA adsorption via the O_β-Mo bond resulted in the lowest adsorption energies compared to other configurations with energies of -0.81 eV over Mo/N_p-CNT and -1.15 eV over Mo/N_g-CNT. Interestingly, the dissociative adsorption was observed for FA adsorption over both catalysts, which happens to AcOH only over Mo/N_p-CNT. The most favourable reaction pathways for FA adsorption and decomposition in both cases are depicted in Figs. 8 and 9 respectively. More details regarding the parallel reactions are shown in Table S3.

Over Mo/N_p-CNT, the dissociation amid the adsorption of FA led to the simultaneous barrier-free cleavage of C-H and C-O_β with the adsorption energy of -3.54 eV, producing three lower fragments of bonded CO, hydroxyl and H atom. The other hydrogen release from hydroxyl showed a 1.98 eV intrinsic energy barrier with the reaction energy of 0.41 eV. Over Mo/N_g-CNT, the FA molecule underwent analogous dissociative adsorption without any energy barriers and produced aldehyde and hydroxyl fragments. The aldehyde H was then cleaved with a small energy barrier of 0.36 eV during an exothermic reaction with -1.34 eV energy, and the hydroxyl cleavage would release the other H atom in a -0.49 eV exothermic process, with an energy barrier of 1.57 eV.

The reactions involved in FA decomposition over both catalysts are facilitated through lower intrinsic energy barriers compared to the free molecule scenario (Fig. 7). The model also predicts that Mo/N-CNTs give rise to different favourable reaction pathways of FA decomposition in comparison with common catalysts [19,60,61]. Instead of the formation of carboxylic or formate intermediates, the dehydroxylation reaction occurs at the beginning, giving rise to facile hydrogen donation at the

early stage. The dehydroxylation is also reported to take place over the Cu (111) facet, but normally with a higher energy barrier [6]. Compared to the decomposition of FA over pure Mo [4], Mo/N-CNTs exhibit better performance in terms of facilitating hydrogen donation by lowering the energy barriers and by decreasing the number of reaction steps.

The two H atoms in the FA molecule show different readiness for donation over the catalysts. Modelling results predict that the H-C bond is more likely to cleave over Mo/N-CNTs compared to the hydroxyl H. Especially, N_p doped CNT induces barrier-free H release during the adsorption, which is comparable to H₂ dissociation over the transition metal surface. The C-H bond cleavage was unfavourable in some circumstances, e.g. over a Mo slab or a Pd/C catalyst [4,62], but it turns out to be much facile over both catalysts in this study. In comparison, the release of hydroxyl H is more difficult than aldehyde H, therefore leading to the peak energy barrier of FA decomposition over both Mo/N-CNTs. It is found also that Mo/N_g-CNT leads to a smaller energy barrier for the release of hydroxyl H than Mo/N_p-CNT does. But it is also noteworthy that the dissociative adsorption energy of FA over Mo/N_p-CNT is much larger than that for Mo/N_g-CNT, which could potentially contribute to the consequent cleavage of hydroxyl and the hydrogen donation (migration) by preventing the formation of H₂O [19].

The modelling has predicted different potentials of hydrogen donation for FA and AcOH. Despite the larger number of H atoms in AcOH, FA has a greater potential for in-situ hydrogen donation over both Mo/N-CNTs catalysts, especially in mild temperature conditions. This is mainly attributed to the facile cleavage of the C-H bond during the initial step of interaction between FA and Mo/N-CNT catalysts, as shown in Figs. 8 and 9.

3.4. Electron density difference (EDD) and atomic charge analyses for typical groups

The reaction modelling results indicate that Mo/N_g-CNT and Mo/N_p-CNT show different performances in facilitating hydrogen donation by affecting the decomposition of typical groups. Specifically, Mo/N_g-CNT is capable of preferentially activating O-H and C-H bonds, and Mo/N_p-CNT would enable the ready cleavage of the C-C bond. To gain new insights into the interaction between the catalysts and reactants, an electron density difference (EDD) analysis was carried out on the typical intermediate groups of acetyl, methyl and hydroxyl over both catalysts, as shown in Figs. 10-12. Atomic charge was also calculated and assigned to each atom as an indicator for interatomic covalency (Table S4).

The analysis reveals that N_p and N_g give rise to different electronegativity effects towards the lower fragment groups, and this is consistent with the catalytic property of both doping types. For the acetyl over Mo/N_p-CNT, electron loss and enrichment were both found around C_β, and mainly electron enrichment around methyl and oxygen is observed. As a result, C_β and methyl tend to be electronically neutral prior to the cleavage, evidenced by the net group charges of 0.01 and 0.00, respectively (Table S4). In the case of Mo/N_g-CNT, Mo had less electron loss, but there was more electron enrichment to make the C_β absolute electric neutral (charge = 0.00), owing to less electron enrichment on N_g and the delocalised electron compensation from N_g. However, fewer electrons were found to migrate to the methyl group in this circumstance, leading to an overall positive charge (0.08) for the methyl in acetyl over Mo/N_g-CNT. The electronic analyses elucidate the modelling results that the cleavage of the C-C bond is much more facilitated by Mo/N_p-CNT; the catalyst enables more electrons loss on Mo, so that it reinforces the Mo-C_β bond. Simultaneously, it gives rise to electron enrichment on C_β and the methyl group, which eventually reduces the strength of the C-C bond and makes it ready to cleave. This is confirmed by the difference in the C-C bond length (1.526 Å and 1.490 Å) over the two catalysts, as shown in Fig. 10. Such a mechanism is also supposed to contribute to the stabilisation of the Mo-O_β bond and the consequent dissociative adsorption of AcOH over Mo/N_p-CNT, where C-

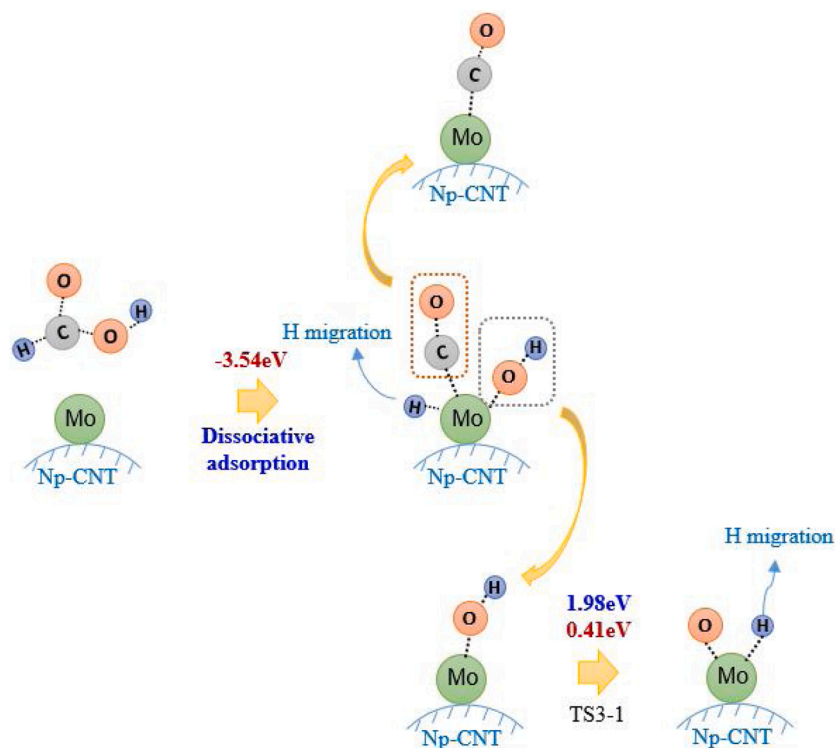


Fig. 8. Reaction flow of the decomposition of FA over Mo/Np-CNT (Blue font denotes energy barrier, and violet denotes reaction energy).

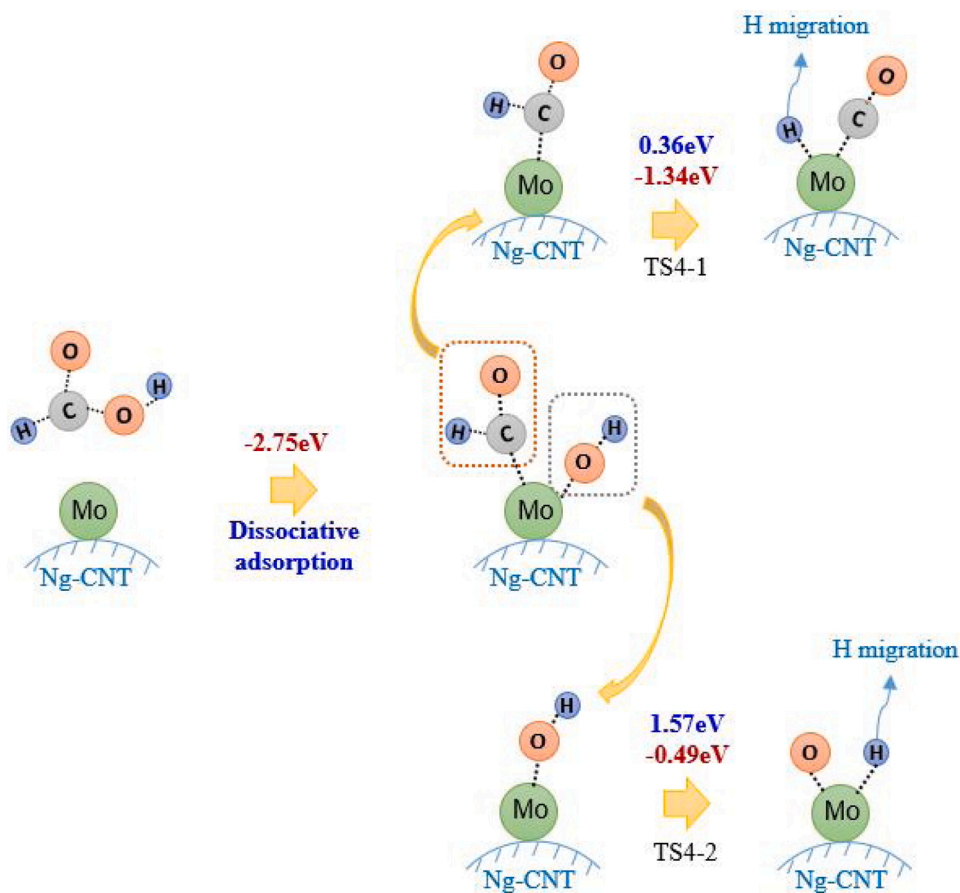


Fig. 9. Flow chart for the decomposition reaction of FA over Mo/Ng-CNT (Blue font denotes energy barrier and violet denotes reaction energy).

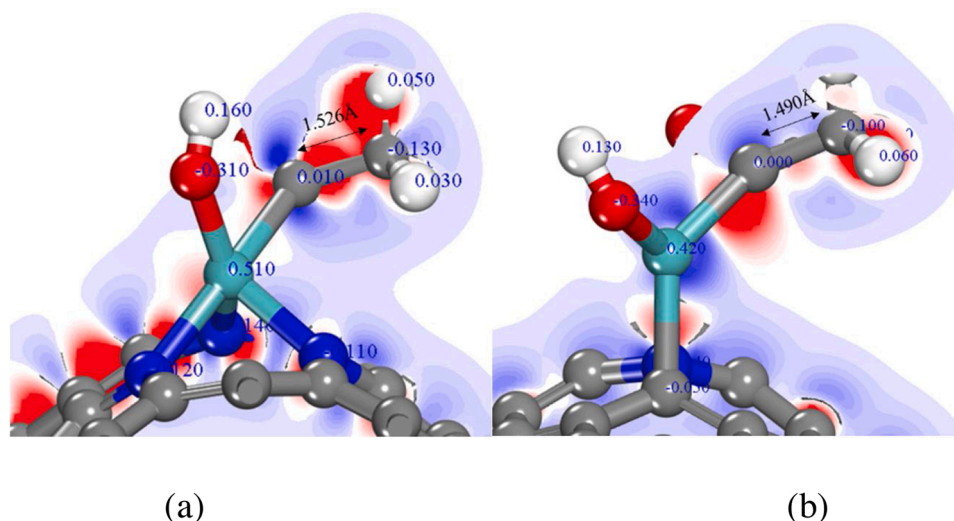


Fig. 10. EDD analyses (threshold value: ± 0.2 electrons/ \AA^3) of the C—C bond structures with atomic charges on (a) Mo/N_p-CNT (b) Mo/N_g-CNT. Red denotes electron enrichment and blue denotes electron loss.

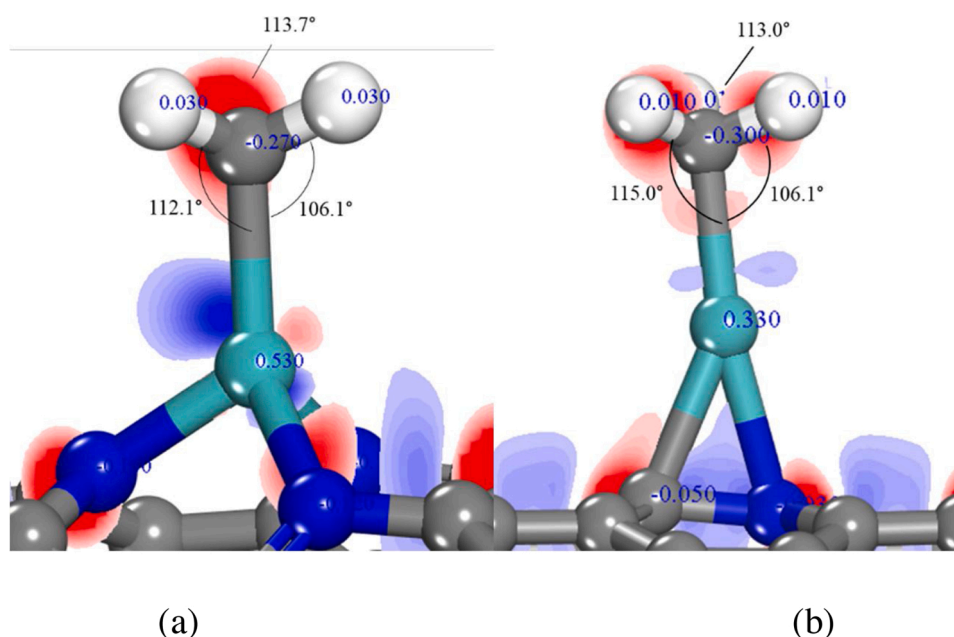


Fig. 11. EDD analyses (threshold value: ± 0.2 electrons/ \AA^3) of the methyl groups with atomic charges on (a) Mo/N_p-CNT (b) Mo/N_g-CNT. Red denotes electron enrichment and blue denotes electron loss.

O_β cleaves. Besides, the results also imply that methyl cleavage from acetyl in these circumstances tends to be homolytic, evidenced by the atomic charge analyses of the corresponding transition states (Fig. S6).

Regarding the adsorbed methyl, electron enrichment around carbon was observed when it was adsorbed on Mo/N_p-CNT, mainly resulting from the electron loss of Mo. In contrast, the methyl group in bulk gained more electrons over Mo/N_g-CNT (Table S4), revealing the effect of the delocalised electron from graphitic doping of N. It is noteworthy that not only the C atom, but also the H atoms are electron enriched in such circumstance, resulting in near charge neutrality of the H atoms and consequently facilitating the hydron donation, analogues to those in the H₂ molecule. More obvious activation of the C—H bond over Mo/N_g-CNT can be observed in Fig. 11 based on the increased Mo—C—H bond angle (to 115.0°), compared to that over Mo/N_p-CNT and 109.5° in common CH₃* tetrahedron structure. The results are in line with the reaction modelling results, and also suggest that methyl is more likely to

release hydrogen through homolytic cleavage over Mo/N-CNTs (this is also evidenced by the atomic charge analyses of the corresponding transition states in Fig. S6), benefiting from the electron enrichment between C-H.

For the adsorbed hydroxyl, less electron loss (more electron back donation) to the H atom was found to take place over Mo/N_g-CNT, as shown in Fig. 12. The adsorbed structures also exhibit stronger bent (to be 129.1°) of \angle MoOH over Mo/N_g-CNT, compared to that (137.7°) over Mo/N_p-CNT, indicating O-H is more activated in this case, with strengthened covalent character of the Mo—OH bond [63]. The release of hydroxyl H is thus facilitated compared to that over Mo/N_p-CNT. Nevertheless, even though Mo has similar electron loss to that binding to the methyl group, the hydroxyl H is not charge neutrality, but carries positive charges over both catalysts (Table S4). This may be because of the large electronegativity of oxygen, giving rise to higher energy barriers for the homolytic cleavage (Fig. S6) of H-O* than *CH₂-H. The

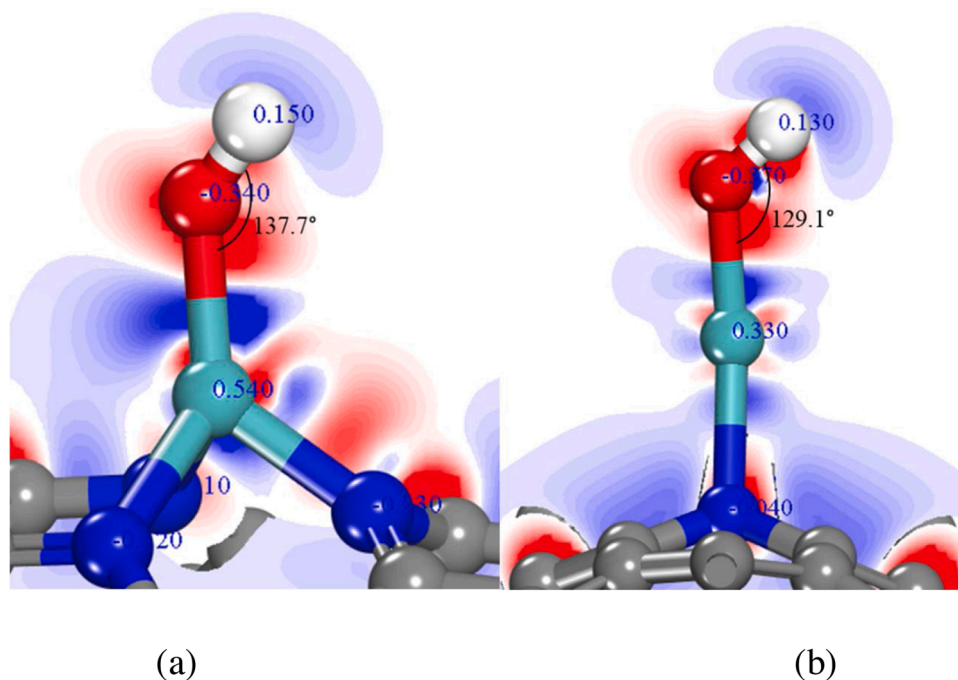


Fig. 12. EDD analyses (threshold value: ± 0.2 electrons/ \AA^3) of the hydroxyl groups with atomic charges on (a) Mo/N_p-CNT (b) Mo/N_g-CNT. Red denotes electron enrichment and blue denotes electron loss.

homolytic cleavages of hydroxyl during acid decomposition would require more electron enrichment. This agrees with the reaction modelling results shown in Figs. 5 and 6.

Generally, the results of EDD analyses in this study suggest that Mo/N_p-CNT would facilitate the C-C cleavage, and Mo/N_g-CNT favours the cleavage of H contained bonds, eventually benefiting the in-situ H donation.

4. Conclusions

This study provided predictions of the hydrogen donation behaviour of bio-acids over the metal loaded N-doped CNTs by DFT calculation. Ten different transition metals and six CNT-based catalysts were modelled. Adsorption modelling indicated that the Mo/N_p-CNT catalyst gave rise to the strongest binding with the adsorbed acid molecule, followed by Mo/CNT and Mo/N_g-CNT. The reaction modelling of the decomposition of AcOH and FA was carried out based on the catalysts of Mo/N_p-CNT and Mo/N_g-CNT, with the emphasis on the H related bond cleavages. The results revealed that Mo/N-CNTs significantly lowered the decomposition energy barriers for both AcOH and FA, while different types of N doping gave rise to various reaction pathways. The most favourable reaction pathways for the decomposition of AcOH over Mo/N_p-CNT and Mo/N_g-CNT were $\text{CH}_3\text{COOH} \rightarrow \text{CH}_3\text{CO} \rightarrow \text{CH}_3 \rightarrow \text{CH}_2 \rightarrow \text{CH} \rightarrow \text{C}$, and $\text{CH}_3\text{COOH} \rightarrow \text{CH}_3\text{CO} \rightarrow \text{CH}_3\text{C} \rightarrow \text{CH}_3 \rightarrow \text{CH}_2 \rightarrow \text{CH}_1 \rightarrow \text{C}$, respectively. In comparison, FA experienced barrier-free decomposition and released hydrogen through $\text{HCOOH} \rightarrow \text{H} + \text{CO} + \text{OH}$, during its adsorption over Mo/N_p-CNT. Over Mo/N_g-CNT, FA underwent similar dissociative adsorption through $\text{HCOOH} \rightarrow \text{HCO} + \text{OH}$, where aldehyde H could be released with a small energy barrier of 0.36 eV. The hydroxyl H release exhibited larger energy barriers over both catalysts. Further electronic analyses revealed that Mo/N_p-CNT would selectively facilitate the homolytic cleavage of the C—C bond, whereas Mo/N_g-CNT would promote the homolytic cleavage of C—H and O—H, through electron enrichment. The modelling work also predicted that FA has a superior hydrogen donating capability to AcOH over the Mo/N-CNT catalysts due to lower energy barriers, suggesting the in-situ hydrogen donation of FA would potentially occur in milder conditions than that of AcOH.

Notes

All authors agree with the content and submission of this original research, and we confirm that the manuscript is not being considered for publication elsewhere. Jiajun Zhang carried out the research and prepared the original draft of the manuscript; Xiaolei Zhang (corresponding author) did proof reading and finalised the manuscript; Amin Osa-tiashtiani did review and revision to the manuscript, Kai Hong Luo, Dekui Shen, Jun Li and Anthony Bridgwater contributed to technical discussions and proof reading of the manuscript. The authors declare no competing financial interest.

Credit author statement

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Declaration of Competing Interest

The authors report no declarations of interest.

Acknowledgement

The authors would like to acknowledge financial support from the Leverhulme Trust Research Grant (RPG-2017-254) and EPSRC First Grant (EP/R010986/1). The authors are also grateful for computational support from the UK Materials and Molecular Modelling Hub, which is partially funded by the EPSRC (EP/P020194/1), for which access was obtained via the UKCP consortium and funded by EPSRC Grant (EP/P022561/1). Additional computational resources from the EPSRC under the project UK Consortium on Mesoscale Engineering Sciences

(UKCOMES) (Grant No. EP/R029598/1) are also gratefully acknowledged.

Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.mcat.2020.111289>.

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