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Ligand assisted hydrogenation of levulinic acid on Pt(111) from first principles calculations[†]Lars Gell,^{*a} and Karoliina Honkala^a

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In this study, we investigate the hydrogenation reaction of levulinic acid to 4-hydroxypentanoic acid on a ligand-modified Pt(111) using DFT. Modifying nanoparticle surfaces with ligands can have beneficial effects on the desired reaction such as improved selectivity or lower activation energies. The N³,N³-dimethyl-N²-(quinolin-2-yl)propane-1,2-diamine (AQ) ligand was selected to modify the surface, since it combines good surface adsorption properties with functional groups that can influence the reaction. The adsorption geometry of the AQ ligand was studied as well as the co-adsorption of a second AQ for the possibility of self-assembly. We find that dissociated hydrogen from the Pt(111) surface can protonate the AQ ligand and discuss the role this plays on the mechanism of the hydrogenation reaction of levulinic acid (LA). By comparing the ligand-modified Pt(111) surface to the bare Pt(111) surface we show that the reaction changes from a step-wise to a concerted mechanism due to the influence of the ligand molecule. This demonstrates the effect ligand-modified surfaces can have in catalyzing reactions and shows that desired reactions can be achieved by tuning the reaction environment.

1 Introduction

Enantiopure compounds, ranging from fundamental synthetic building blocks like amino acids or epoxides to commercial end products such as drugs and food additives,^{1,2} are extensively used by the chemical and pharmaceutical industry. This creates a demand for large scale enantioselective production processes. Traditionally, enantioselective syntheses are carried out using asymmetric homogeneous catalysts made of chiral transition metal complexes.³ These molecular catalysts mostly feature one central metal atom bonding with multiple, often different, ligands. The ligand molecules' influence on the overall properties of the transition metal complex is significant and therefore their role on activity and selectivity in asymmetric homogeneous catalysis as well as on the stability of the catalyst has been extensively studied.^{4–7} Metal complex-based catalysis, however, suffers from limitations such as low operation temperatures and difficulties in separating the products from the catalyst.^{4,8} Metal nano-particles can circumvent these shortcomings.^{9–14} Due to their unique prop-

erties such as high surface to volume ratio and a large number of low coordinated sites, metal nano-particles provide high stability combined with the possibility of high reactivity.^{15,16} Unfortunately, this often comes at the expense of high enantioselectivity due to the multiple reaction sites which can lead to different reaction mechanisms and, consequently, the formation of different products. Introducing ligands on metal nano-particles is one way to improve the enantioselectivity.^{17,18} Adsorbed ligands can be used to block certain reaction sites or mechanisms, tune the steric and electronic interactions with the reactant molecule, and modify the chemical properties of the surface such as hydrophilicity, acidity, and chirality.^{19,20} Ultimately, this will lead to highly specific reaction environments, similar to enzymes, that are tailored towards the desired product.

Ligand design is crucial in creating self-assembled enzyme-like reactant environments that control reaction mechanisms and achieve optimal enantioselectivity. A large variety of ligand molecules such as N-heterocyclic carbenes,^{21,22} secondary phosphine-oxides,²³ thiols,²⁴ amino acids,²⁵ and amidates²⁶ can be used to decorate metal nano-particles.^{19,27} Many ligand-modified late transition metal nano-particles have shown catalytic activity¹⁹ towards reactions such as the hydrogenation of organic acids, ketones and alcohols²⁸, the reduction of CO₂²⁹ and the oxidation of C-C-double bonds.³⁰ For hydrogenation reactions on Pt, cinchona alkaloid ligands have shown promising experimental results^{9,11,31–33} such as high enantioselectivity in multiple ketone reduction reactions.³⁴ According to DFT calculations, quinoline-

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[‡] Additional footnotes to the title and authors can be included e.g. 'Present address:' or 'These authors contributed equally to this work' as above using the symbols: ‡, §, and ¶. Please place the appropriate symbol next to the author's name and include a \footnotetext entry in the the correct place in the list.

type ligands show strong binding energies on Pt(111) surfaces, especially when co-adsorbed with hydrogen or oxygen, making them stable under moderate reaction conditions.³⁵

One key application of asymmetric catalysis is the transformation of biomass into value added products through hydrogenation reactions like the reduction of levulinic acid (LA) to γ -valerolactone (GVL).³⁶ LA is a versatile platform chemical that can be obtained on an industrial scale from biomass which is often a byproduct of agricultural production e.g. wheat straw.^{37,38} GVL is another important platform chemical for fuel and fine chemical production.^{36,39,40} Several experimental and computational studies have shown the hydrogenation of LA can occur via homogeneous as well as heterogeneous catalysis.^{41–46} The homogeneously catalyzed reduction of LA to GVL can be achieved using various transition metal complexes.^{43,45,47} Previous DFT studies have shown low activation energies (< 0.3 eV) for this transition.^{43,45} Heterogeneous reduction of LA has been addressed on bare metals,⁴¹ metal alloys⁴⁰ and metal oxides.⁴⁸ Ru-catalyzed LA reduction can follow two different mechanisms: the alkoxy mechanism and the hydroxy mechanism. In the alkoxy mechanism, hydrogen initially forms a bond to the carbonyl carbon atom followed by a second hydrogen binding to the carbonyl oxygen atom. The hydrogen addition steps are reversed in the hydroxy mechanism.⁴¹ The transformation to 4-hydroxypentanoic acid (4-HPA) is endothermic by 0.81 eV and to GVL is exothermic by 0.05 eV.⁴¹ In general, for the transformation to GVL, the alkoxy mechanism is favoured with an activation barrier of 0.5 eV whereas the hydroxy mechanism shows an activation barrier of more than 1 eV.⁴¹ Previous calculations have shown solvents affect the hydrogenation of LA. Solvents with higher polarity than methanol, such as water, create a more favourable reaction environment thereby lowering the rate determining barrier by 0.1 eV.⁴⁶ Lower reaction rates for the reduction of LA have been experimentally observed on Pt surfaces^{44,49} compared to other transition metals (e.g. Ru, Pd) suggesting higher selectivity.^{50,51}

In the present study, we address the reduction of LA on a ligand-modified Pt(111) surface. We choose a 2-amino-quinoline derivative to modify the surface because it binds sufficiently strongly to Pt(111) and allows intermolecular hydrogen bonding as well as ligand-reactant interaction. The self-assembly of these ligands and the influence of their structure on the reduction of LA are discussed. Furthermore, we demonstrate how the introduction of an amino-quinoline derivative can modify the reaction mechanism thereby opening the possibility for higher enantioselectivity.

2 Computational Details

2.1 DFT

We used density-functional theory (DFT) with a projector augmented wave method (PAW)⁵² as implemented in the real-space grid code-package GPAW.^{53–55} In the finite difference (FD) mode, the PAW setup was defined for atomic valence of Pt($5p^6 6s^1 5d^9$), O($2s^2 2p^4$) N($2s^2 2p^3$), C($2s^2 2p^2$), and H($1s^1$), with scalar-relativistic effects included for Pt. Total energy calculations and structure optimizations were done using the BEEF-vdW⁵⁶ ex-

change correlation functional together with the real space grid spacing smaller than 0.2 Å and 0.05 eV/Å convergence criteria for the residual forces acting on atoms. The Pt(111) surface was modelled as a 4-layer thick 7x7 2D slab cut from Pt bulk with a lattice constant of 4.02 Å. The two topmost layers were relaxed and the remaining bottom layers were fixed into their bulk positions. Periodic boundary conditions were used in the surface plane and more than 6 Å of empty space above and below the surface was applied. The calculations were carried out with a single Γ -point. Adsorption geometries of AQ were screened using the localized atomic orbitals (LCAO)⁵⁷ method with single zeta basis sets on all the atoms. Transition states were identified using climbing image NEB^{58,59} and confirmed through vibrational frequency analysis along the reaction coordinate. These calculations were performed on a 3-layer thick slab to reduce computational costs. The Bader method⁶⁰ was employed to evaluate individual atomic partial charges. The adsorption energies were calculated according to the equation

$$E_{\text{adsorb}} = E_{\text{ads}} - (E_{\text{surf}} + E_{\text{gas}}) \quad (1)$$

where E_{ads} is the energy of the surface and the adsorbed molecule, E_{surface} is the energy for the bare surface, and E_{gas} represents the energy of the molecule in the gas phase.

2.2 Structural Details

N³,N³-dimethyl-N²-(quinolin-2-yl)propane-1,2-diamine (AQ) (Fig. 1) was selected as a ligand to modify the Pt(111) surface. With this ligand selection, we aim to achieve three key elements: 1) Strong binding to Pt(111) in order to ensure catalyst stability under reaction conditions. Based on the binding properties of cinchona alkaloids, we choose a quinoline double ring as the anchoring unit to ensure strong ligand-surface interaction. 2) This ligand offers the possibility for attractive ligand-ligand as well as ligand-reactant interactions through intermolecular hydrogen bonds. These interactions create the possibility of self-assembled enzyme-like reaction pockets. 3) A ligand may have a beneficial influence on the enantioselectivity of the desired reaction. Potential strategies to achieve this are the limitation of adsorption configurations for the reactant (lock and key principle), the modification of the reaction mechanism or the kinetic discrimination of unwanted configurations. In the case a single AQ ligand, more than 50 different adsorbed structures were screened at the LCAO level of theory, and 20 of them were further optimized using the FD method, while in the case of two co-adsorbed AQ ligands the amount of structures was reduced to 20 and 11 respectively. On the bare Pt(111) surface, reaction pathways were calculated for both Levulinic acid with (LA1) and without (LA2) an internal hydrogen bond. Due to the higher computational costs, we focused on the reduction of LA2 in the case of the ligand modified surface.

3 Results and Discussion

3.1 Binding of AQ on Pt(111)

AQ configurations on the Pt(111) surface were divided into 3 classes according to certain structural features. Class A shows

Fig. 1 The structure of the AQ ligand molecule, N¹,N¹-dimethyl-N²-(quinolin-2-yl)propane-1,2-diamine, is shown, with the blue letters and numbers indicating the notation of individual atoms that will be referenced later.

Fig. 2 The most energetically favoured structures from each class are shown. The corresponding relative energies are a) class A, 0.00 eV, b) class B, 0.17 eV and c) class C 0.44 eV, absolute energies and interatomic distances can be found in Table 1

the N³ atom at a greater distance from the surface than its neighbouring carbon atoms and the C2'-C3' bond parallel to the surface. Class B is characterized by having the C3' and the N³ atoms being closer to the surface than their neighbouring carbon atoms. In class C, the tertiary amine moiety is close to the surface as in class B, while the C2'-C3' bond points away from the surface. The most energetically favoured configuration in each class is shown in Fig. 2. All classes bind to the Pt(111) surface through the quinoline moiety. Three energetically most favourable configurations from each class were selected and then relaxed at the FD level, as described above. The adsorption energy depends on the conformation of the AQ ligand, particularly the position of the quinoline moiety, as well as the adsorption site on Pt(111). In general, orientations in which the quinoline double ring system is not parallel to the Pt surface are very energetically unfavourable. In the most energetically favoured geometries, the quinoline rings are sitting over a rhomboid Pt₄ unit with ring-surface distances varying between 2.2 Å and 3.5 Å. These short distances, together with an exothermic adsorption energy of -1.22 eV per molecule, are a clear indication of chemisorption. The adsorption energy for AQ on Pt(111) is significantly more exothermic than cinchonidine (about -0.4 eV per molecule).^{31-33,61} Short distances between the quinoline group and the surface are energetically favoured while short distances between the N³ nitrogen and the surface are energetically unfavoured (see Table 1 as well as Fig. S2).

3.2 AQ Co-Adsorption

The co-adsorption of ligands is an important step towards a self-assembled layer that creates an enantioselective reaction environment. We therefore also explore the energetics and structural features of two co-adsorbed AQ molecules. In order to reduce the computation expense, only the three best conformers shown in

	E_{adsorb} (eV)	N ³ -Pt dist. (Å)	Ring-Pt dist. (Å)
a)	-1.22	3.78	2.71
b)	-1.06	3.84	3.10
c)	-0.79	3.10	2.40

Table 1 Adsorption energies and selected distances for the AQ structures shown in Fig. 2.

Fig. 3 The adsorption geometries of three selected co-adsorption structures are shown. The corresponding energies are a) anti-parallel, 0.00 eV, b) separated, 0.07 eV and c) parallel, 0.17 eV.

Fig. 2 were considered and combined to investigate the different orientations of two co-adsorbed AQ ligands (2AQ). The initial structures were generated by introducing the second AQ ligand into the unit cell, then translating and rotating it with respect to the initial AQ ligand while keeping similar AQ-surface distances as reported in Table 1. The three selected configurations shown in Fig. 3 highlight the effect of the alignment of class A ligands with respect to each other and the distance between them. As expected, given the higher adsorption energies for class B and class C molecules, combinations of these conformations are also less energetically favourable. In the lowest energy configuration, the AQ ligands are antiparallel to each other. This leads to the formation of two intermolecular hydrogen bonds between the secondary amine on one ligand and the quinoline nitrogen atom on the other ligand. As a result, the distance between the quinoline nitrogen and the Pt(111) surface increases slightly from 2.8 Å observed for the single AQ ligand to 3.2 Å.

The adsorption energy of a second AQ ligand in the most energetically favoured orientation is -1.34 eV. The attractive interaction of the 2AQ system lowers the energy by 0.12 eV compared to two infinitely separated AQ ligands. Ligand-ligand interactions are much weaker than the dominant ligand-surface interactions. Attractive interactions are due to the two intermolecular hydrogen bonds while repulsive interactions are attributed to unfavourable positioning on the surface as well as steric repulsion. The structure shown in Fig. 3 b) shows the 2 AQ ligands in a similar orientation to a) but separated by approximately 6 Å. Although the structural properties of each ligand are similar to those of a single adsorbed AQ ligand shown in row a) of Table 1, the separated co-adsorbed AQ ligands are slightly less energetically favoured than the co-adsorbed AQ ligands shown in Fig. 3 a). The parallel configuration shown in Fig. 3 c) eliminates

the possibility of the forming intermolecular hydrogen bonds. In fact, this configuration shows repulsive ligand-ligand interaction and, as a result, its energy is 0.17 eV higher than the antiparallel configuration. Shorter ligand-ligand distances and the orientation of ligands with respect to each other are therefore crucial in forming attractive ligand-ligand interactions, such as hydrogen bonding. Strategies to increase the attractive interaction between adsorbed ligands include strengthening hydrogen bonds by using chlorinated or fluorinated compounds, increasing the number of possible hydrogen bonds or linking more complex ligands by covalent bonds.

3.3 Ligand Protonation

The dissociative adsorption of molecular hydrogen is known to occur without any kinetic or thermodynamic barrier on Pt(111).⁶² It is therefore possible for the AQ ligand (see Fig. 3 a)) to be protonated from readily available hydrogen atoms on the Pt(111) surface when the surface is exposed to H₂ gas. The adsorption energy of hydrogen on 2AQ-covered Pt(111) is -0.18 eV, which is slightly less exothermic than experimental values (-0.3 eV) on bare Pt.⁶³ Both N² and N³ atoms of the AQ ligand were considered for protonation. Both of these nitrogens are Lewis bases and, therefore, the most likely candidates for protonation. Our calculations show that protonating the N³ atom, thus forming AQH as seen in Fig. S3 a), is thermodynamically favoured over protonating the N² atoms by 0.92 eV. The N³ atom can adopt a tetrahedral configuration more easily because the methyl groups around it are a lesser steric hindrance compared to the quinoline moiety bonded to the N² atom.

Fig. 4 The geometries of three selected structures with zero, one and two hydrogens transferred to the ligand are shown. The corresponding energies are a) 0.00 eV, b) -0.38 eV, and c) -0.80 eV. Dissociated hydrogen atoms adsorbed on the Pt(111) surface are shown in light blue and those transferred to the AQ ligand are shown in red

The hydrogen transfer to the N³ atom from the Pt(111) surface is a barrier-less exothermic process of -0.38 eV (see Fig. 4). The N³-Pt(111) distance decreases significantly from 3.8 Å to 3.2 Å as the N³ atom adopts a tetrahedral configuration upon protonation. Protonating the second AQ ligand is exothermic by an additional 0.42 eV and results in both ligands having similar structural properties (see Fig. 4).

We employ the Bader charge method⁶⁰ to determine if a protonic hydrogen has been truly transferred to the ligand. On Pt(111), H atoms are approximately neutral with a partial charge

of -0.1 e. After the transfer, the partial charge of the hydrogen atom is +0.4 e with a N-H distance of 1.07 Å, indicating the clear protonic nature while the charge of the remaining surface hydrogen is still -0.1 e due to the interaction with the conducting metal surface. The outcome of this hydrogen transfer process is an asymmetrically charged hydrogen pair resembling the products of a heterolytic hydrogen dissociation. This result is significant because asymmetrically charged hydrogen pairs can influence the mechanism and energetics of reduction reactions⁶⁴.

3.4 Levulinic acid reduction to 4-HPA on Pt(111)

In order to evaluate the effectiveness of the ligand-modified surface, we study the first step in the hydrogenation of LA, that is, the conversion of LA to 4-HPA. This reduction reaction can serve as a model for the hydrogenation of polyfunctional molecules and can help determine the effect of a ligand-modified Pt(111) surface on the reaction mechanism and energetics compared to a bare Pt(111) surface. Two different adsorption configurations for LA are considered, the first, LA1, has one internal hydrogen bond while the second, LA2, does not (see Fig. S4). Our calculations agree with a previous study⁴¹ showing that gas-phase LA1 and LA2 are almost isoenergetic. LA1 is 0.05 eV less stable than LA2 (see I₁ and I₂ in Fig. 7 and Table 2) due to two competing effects. The formation of the internal hydrogen bond stabilizes the molecule while the non-planarity of the carbon skeleton destabilizes it. In contrast, whether in gas phase or on the Pt(111) surface, LA2 lacks the internal hydrogen bond but its carbon skeleton is nearly planar, as can be seen from the minimum energy structures in Fig. S4. Nevertheless, both structures are kept to investigate LA reduction since the internal structure may influence the reaction mechanism.

The LA adsorption energies on a 2/9 ML H₂-covered Pt(111) surface are -0.59 eV for LA1 and -0.78 eV for LA2 (see I₃ and I₄ in Fig. 7 and Table 2). These adsorption energies are significantly less exothermic than the adsorption energy of -1.33 eV for LA on bare Ru(0001), which is also active for LA reduction.⁴¹ Both LA conformers favour the parallel adsorption geometry on Pt(111) lying on average about 3.70 Å above the surface. For LA1, the carboxylic oxygen are located above Pt bridge sides, whereas for LA2, they are above hollow sites, as can be seen in Figs 5 a) and S5 a). Like LA2 on Pt(111), the structure of LA adsorbed on Ru(0001) also lacks an internal hydrogen bond, but all of the oxygen atoms bind to surface bridge sites resulting in shorter distances between LA and the Ru(0001) surface.⁴¹ We also considered adsorption configurations with smaller LA-Pt distances similar to those identified on the Ru surface. Our calculations demonstrate, however, that these structures are unstable on Pt(111). During structural optimization, LA returns to a flat conformation with a larger LA-surface distance, similar that shown in Fig. 5 a).

Three possible reaction mechanisms are considered for the reduction of LA to 4-HPA: the hydroxy mechanism, the alkoxy mechanism, and the concerted mechanism. In the hydroxy mechanism, the H-O bond is initially formed, followed by C-H bond formation (light blue (LA1) and purple (LA2) path in Fig. 7). These reaction steps are reversed in the alkoxy mechanism (green

Fig. 5 Stationary points along the hydroxy mechanism for LA2 as referenced in Fig. 7 and Table 2: a) adsorbed reactants, I_4 b), intermediate I_8 , c) transition state TS_6 , and d) adsorbed 4-HPA, I_9 . The hydrogen atoms involved in the reaction are circled for better visibility.

(LA1) and red (LA2) path), and proceed simultaneously in the concerted mechanism. Although the concerted mechanism was considered, we were unable to locate a transition state as our NEB calculations always converted to either the hydroxy or the alkoxy mechanisms. Fig. 7 and Table 2 display the relative potential energies surface for all stationary points along the examined reaction paths. The O-H bond formation in the hydroxy mechanism leads to metastable intermediates I_5 at +0.33 eV and I_8 -0.09 eV for LA1 and LA2 respectively (see Fig. 7 and Table 2). Both intermediate structures resemble their respective initial adsorbed LA conformers. While no kinetic barriers are observed for the O-H bond formation, the thermodynamic barriers are +0.92 eV (LA1) and +0.69 eV (LA2) with respect to their initial states, I_3 and I_4 . The C-H bond formation producing 4-HPA (I_9 in Fig. 7) is exothermic by 1.21 eV (LA1) and 0.79 eV (LA2) but has a small kinetic barrier 0.17 eV and 0.06 eV respectively. The hydrogen atom involved in the second hydrogenation step is relatively close to the surface with H-C distances of 2.03 Å for TS_3 (LA1), and 1.94 Å for TS_6 (LA2) as shown in see Figs. S5 and 5. The overall reaction barrier, defined as the energy difference between the reactants and the highest energy structure along the reaction path, is 1.09 eV for LA1 and 0.75 eV for LA2. The reaction energy for the hydrogenation of LA to 4-HPA adsorbed on Pt(111) is exothermic by -0.22 eV for LA1 and -0.10 eV for LA2. In comparison, the reduction of LA with no internal hydrogen bond on Ru(0001) must overcome a kinetic barrier of 1.36 eV to produce a hydroxy intermediate that is 0.88 eV higher in energy than the adsorbed LA reactant. The second hydrogenation on Ru(0001) is slightly exothermic (0.05 eV) with a kinetic barrier of 0.67 eV. The overall reaction barrier on Ru(0001) is 1.64 eV and the corresponding reaction energy for the hydrogenation to 4-HPA is endothermic by 0.82 eV.⁴¹

For the alkoxy mechanism, the key reaction steps are displayed in Figs. S6 (LA1) and S7 (LA2). The formation of intermediates I_6 for LA1 and I_7 for LA2 via hydrogen transfer to carbon is endothermic by 0.77 eV and 0.75 eV relative to their corresponding adsorbed LA configurations, I_3 and I_4 , respectively (see Fig. 7). The energies for transition states involved this hydrogen transfer,

Fig. 6 Structures of stationary states referenced in Fig. 7 for the concerted hydrogenation reaction of levulinic acid on a Pt(111) surface in the presence of 2AQH a) Adsorbed LA2 in the presence of 2AQ, (I_{12}), b) transition state (TS_8), and c) adsorbed 4-HPA (I_{14}).

TS_1 (LA1) and TS_2 (LA2) are 1.21 eV and 1.34 eV respectively. The C-H distance for TS_1 is 1.45 Å and 1.44 Å for TS_2 . The second hydrogen transfer is exothermic by 1.06 eV (LA1) and 0.85 eV (LA2) and has a small kinetic barrier of 0.18 eV and 0.21 eV corresponding to TS_4 and TS_5 respectively. The O-H distance for TS_4 is 1.55 Å and 1.47 Å for TS_5 . On Ru (0001), a comparable alkoxy intermediate without an internal hydrogen bond is almost isoenergetic with its initial state, presenting activation energies of 0.67 eV and 1.29 eV for first and second hydrogenation respectively.⁴¹

We observe the hydrogenation of LA to 4-HPA follows different mechanisms on bare Pt(111) and on Ru(0001). The hydroxy mechanism is heavily preferred on Pt(111) while the alkoxy mechanism is favoured on Ru(0001). On Pt(111), the activation energy of the hydroxy mechanism is 0.59 eV lower than that of the alkoxy mechanism. In the case of Ru(0001), the activation energy for the alkoxy mechanism is 1.35 eV while the activation energy for the hydroxy mechanism is 1.64 eV relative to the adsorbed reactants. Furthermore, the overall reaction is exothermic by -0.10 eV on Pt(111) while it is endothermic by 0.82 eV on Ru(0001). Previous experimental studies on the conversion of LA have suggested 4-HPA is an intermediate, losing a water molecule to form GVL.^{44,65} We find the 4-HPA conversion to GVL on Pt(111) is exothermic by -0.15 eV. No barriers have been computed, however, since the focus in the present work is to explore the effect of a ligand's presence compared to the bare surface rather than the full reaction network to GVL. Interestingly, previous DFT calculations suggest that GVL formation on Ru(0001) occurs via an alternative mechanism that does not involve 4-HPA.⁴¹

3.5 Levulinic acid reduction on 2AQH-modified Pt(111)

Lastly, we consider the effect of adsorbed AQ on LA hydrogenation, particularly in its protonated form, AQH. The adsorption energy of LA2 on 2AQH/Pt(111) (Fig. 4 b)) is -1.04 eV, slightly more exothermic than the adsorption on the bare Pt(111) surface (-0.78 eV). This indicates that the 2AQH ligands stabilize LA2, most likely due to hydrogen bonding between the N^3H and the LA2 oxygen (see Fig. S8 a)). The average LA2-Pt(111) distance in the presence of 2AQH is 3.85 Å which is 0.15 Å larger than on the bare surface, again, most likely due to the interaction with the N^3H group.

Three possible mechanisms were also explored for the hydro-

Fig. 7 Reaction network of LA to 4-HPA on bare and 2AQ covered Pt(111) surface. The corresponding energies can be found in Table 2.

genation reaction of LA2 on ligand-modified Pt(111) (Fig. 6 a)). The major difference with the reduction of LA on the bare Pt(111) surface is that, as a result of the barrier-less exothermic AQ protonation, we consider a proton transfer from the AQ ligand to LA2. Our DFT results show that the concerted mechanism on 2AQH/Pt(111) is slightly endothermic by 0.17 eV and involves a transition state (TS₇) with an activation energy of 0.77 eV above the initial state, I₁₂, consisting of adsorbed LA2 and 2AQH (refer to Fig. 6). For the TS₇ structure, the O-H bond length is 1.40 Å, while the C-H bond length is 1.31 Å. This indicates the slightly asymmetric nature of the transition state because the O-H bond is only 69 % formed while the C-H bond is already 84 % formed. Our findings suggest that the proton at the N³ atom is crucial for the concerted mechanism as we were unable to locate the transition state for the simultaneous hydrogen transfer on the bare Pt(111) surface. In the hydroxy mechanism, the first hydrogen transfer, going from I₁₂ to I₁₃, is endothermic by 0.85 eV (see Fig. 7) with no additional kinetic barrier. The C-H bond formation in the hydroxy mechanism is exothermic by 0.69 eV. For this step, a transition state, TS₈, is observed, however, it is thermoneutral (+0.01 eV). The transition state C-H distance is 1.72 Å. This mechanism is very similar to the hydroxy mechanism observed on the bare surface except for a small barrier for the second hydrogen addition of 0.06 eV. The overall barrier for the hydroxy mechanism in the presence of 2AQ is 0.11 eV higher than on the bare surface. We were unable to locate any transition states for the alkoxy mechanism, however, a thermodynamic barrier of 0.78 eV was determined by constraining the C-H distance to 1.08 Å, which is a typical distance for the computational method used. This intermediate is highly unstable and will revert back to the initial structure during the NEB calculation once the constraints are removed.

The hydroxy mechanism and the concerted mechanism are quite similar energetically, with an activation barrier relative to the I₁₂ structure of 0.77 eV for the concerted mechanism and 0.85 eV for the hydroxy mechanism. The presence of 2AQ ligands on the Pt(111) changed the preferred reaction mechanism from the hydroxy mechanism to the concerted mechanism. By

further improving the ligand design we believe it will be possible to positively influence the reactivity as well as selectivity of this reaction.

4 Conclusions

We investigated the adsorption of AQ ligands on Pt(111), the interaction between adsorbed AQ ligands, AQ protonation, and the hydrogenation of LA to 4-HPA on bare and AQ-modified Pt(111) surfaces using DFT. The exothermic adsorption of a single AQ ligand occurs when the quinoline ring aligns parallel over a Pt₄ rhombus. Two AQ ligands aligned antiparallel to each other form the most stable co-adsorption structure due to the weak ligand-ligand attraction from two intermolecular hydrogen bonds. In the presence of ligands, the transfer of dissociated hydrogen to the N³ atom of the AQ ligand is a thermodynamically preferred non-activated process. Both LA configurations, with and without internal hydrogen bonding, bind exothermically to the Pt(111) surface. AQ ligands also enhance LA binding to the Pt(111) surface. Three reaction mechanisms were considered for LA hydrogenation to 4-HPA on the bare Pt(111) and ligand-modified Pt(111) surfaces. On the bare Pt(111) surface, the hydroxy mechanism is energetically favoured over the alkoxy mechanism. The concerted mechanism converted to either the hydroxy or alkoxy mechanisms on the bare Pt(111) surface. Our work demonstrates that adsorbed AQ can actively participate catalytic reactions. By introducing AQ ligands onto the Pt(111) surface, the hydrogenation process changes dramatically. The protonated N³ atom in AQ provides the protonic hydrogen to the ketone oxygen in LA, while the surface provides the other hydrogen atom to the carbonyl carbon thus forming 4-HPA. The concerted mechanism is favoured over the hydroxy mechanism by 0.1 eV. Overall, both bare and AQ-modified surfaces are equally active towards the conversion of LA to 4-HPA. Based on our computational findings, we believe that, with careful ligand design, it is possible to improve the activity as well as the selectivity of ligand-modified Pt catalysts for hydrogenation of polyfunctional molecules. These improved ligands could feature multiple sites that interact with the reactant to ensure selective positioning before and during the reaction as well as stronger ligand-ligand interactions. DFT calculations are a valuable tool in a systematic exploration of different ligand structures to establish trends for desirable ligand properties.

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Symbol	Intermediate structure / transition state	Energy (eV)	Figure
I ₁	LA1(g) + Pt(111) + H ₂ [*]	0.05	NA
I ₂	LA2(g) + Pt(111) + H ₂ [*]	0.00	NA
I ₃	LA1 [*] + Pt(111) + H ₂ [*]	-0.59	S4 a)
I ₄	LA2 [*] + Pt(111) + H ₂ [*]	-0.78	S4 b)
I ₅	Hydroxy LA1H [*] + Pt(111) + H ₁ [*]	0.33	S5 b)
I ₆	Alkoxy LA1H [*] + Pt(111) + H ₁ [*]	0.18	S6 b) and c)
I ₇	Alkoxy LA2H [*] + Pt(111) + H ₁ [*]	-0.03	S7 b)
I ₈	Hydroxy LA2H [*] + Pt(111) + H ₁ [*]	-0.09	5 b)
I ₉	4-HPA [*] + Pt(111)	-0.88	5 c)
I ₁₀	LA2(g) + 2AQ [*] + Pt(111) + H ₂ [*]	0.00	4 a)
I ₁₁	LA2(g) + 2AQH [*] + Pt(111) + H ₁ [*]	-0.38	4 b)
I ₁₂	LA2 [*] + 2AQH [*] + Pt(111) + H ₁ [*]	-1.42	6 a)
I ₁₃	Hydroxy LA2H [*] + 2AQ [*] + Pt(111) + H ₁ [*]	-0.57	S8 b)
I ₁₄	4-HPA [*] + 2AQ [*] + Pt(111)	-1.25	6 c)
TS ₁	Alkoxy LA1 [*] + H ₁ ^{**} + Pt(111) + H ₁ [*]	0.62	S6 a)
TS ₂	Alkoxy LA2 [*] + H ₁ ^{**} + Pt(111) + H ₁ [*]	0.56	S7 a)
TS ₃	Hydroxy LA1H [*] + Pt(111) + H ₁ ^{**}	0.50	S5 c)
TS ₄	Alkoxy LA1H [*] + Pt(111) + H ₁ ^{**}	0.36	S6 d)
TS ₅	Alkoxy LA2H [*] + Pt(111) + H ₁ ^{**}	0.18	S7 c)
TS ₆	Hydroxy LA2H [*] + Pt(111) + H ₁ ^{**}	-0.03	5 b)
TS ₇	Simultan LA2 [*] + 2AQ [*] + Pt(111) + H ₂ ^{**}	-0.65	6 b)
TS ₈	Hydroxy LA2H [*] + 2AQ [*] + Pt(111) + H ₁ ^{**}	-0.56	S8 c)

Table 2 Intermediates and transition states and their corresponding energies from the potential energy surface Fig. 7. H^{**} indicates the reacting hydrogen while ^{*} and (g) refer to adsorbed and gas-phase species. The Figure column gives the reference for the figure with the atomic structure.

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