

Cite this: *RSC Adv.*, 2017, 7, 13473

Theoretical insights into ω -alkynylfuran cycloisomerisation catalyzed by Au/CeO₂(111): the role of the CeO₂(111) support†

Yafei Luo,^a Zhongzhu Chen,^a Jin Zhang,^{*b} Ying Tang,^b Zhigang Xu^a
and Dianyong Tang^{*ab}

Understanding the role of CeO₂(111) supports is meaningful for designing high-performance metal oxide-supported gold nanoparticle catalysts. Here, density functional theory calculations were employed to study ω -alkynylfuran cycloisomerisation on CeO₂(111)-supported Au clusters. The various reactive sites, including the atop, interface and edge sites, and the role of oxygen vacancies on the defective CeO₂(111) surface were taken into account. On the basis of the computed results and energetic analysis, it was found that the atop and interface sites show high catalytic activity toward ω -alkynylfuran cycloisomerisation, and the edge site of Au₁₀/CeO₂(111)-s (stoichiometric) exhibits similar catalytic activity. Meanwhile, the surface oxygen vacancies of CeO₂(111) can cause some negative and positive effects on the adsorption energy and the catalytic activity. Furthermore, to shed light on the role of the CeO₂(111) support, an analysis of ω -alkynylfuran cycloisomerisation catalyzed by Au₃₋₄/CeO₂(111) and free Au₃₋₄ clusters was performed as well. Our results indicate that the presence of the CeO₂(111) support can not only change the orbital levels of the gold clusters, but also cause charge recombination and decrease the positive charge on the Au (top) atom. These virtues can effectively lead to a decrease in the adsorption energy, facilitating electrophilic attack of the C atom on the furan ring.

Received 23rd November 2016

Accepted 20th February 2017

DOI: 10.1039/c6ra27207j

rsc.li/rsc-advances

Introduction

Metal-oxide-supported gold nanoparticle (NP) catalysts have attracted extensive attention in both theory and experiments owing to their extraordinary catalytic activities for numerous reactions, including the water–gas shift reaction,^{1,2} CO oxidation,^{3,4} and so on. Among the various metal oxides, due to its high oxygen storage and release capacity, and the fact that it contains highly localized 4f orbitals,⁵⁻⁷ cerium oxide/ceria (CeO₂) has been recognized as an excellent supporting material. In addition to the above-mentioned properties, the formation of oxygen vacancies on the surface of CeO₂ is easy, which has been demonstrated *via* experimental studies.^{8,9} The presence of these oxygen vacancies can lead to strong binding between Au nanoparticles and the CeO₂ surface.¹⁰ In addition, oxygen vacancy formation on the CeO₂ surface can result in the reduction of adjacent Ce⁴⁺ ions to Ce³⁺ ions, so the concentration of Ce³⁺ ions is closely related with

the number of oxygen vacancies, exhibiting a proportional relationship.⁹ Meanwhile, according to previous reports,¹¹ oxygen vacancies on the CeO₂ surface can play a curial role in the catalytic process. In a nutshell, CeO₂-based catalysts have attracted wide attention due to the unique characteristics of the CeO₂ surface.

CeO₂-supported Au catalysts have been regarded as promising catalytic systems because they include two highly active components, namely, the CeO₂ surface and Au nanoparticles.^{12,13} Ciftci *et al.* prepared a set of catalysts with Au and Pt supported on CeO₂ nanorods and nanocubes, which were applied as catalysts in the decomposition of ethanol and formic acid.¹⁴ They found that Au catalysts are more active and selective toward H₂ than Pt catalysts in formic acid decomposition. Fu and coworkers reported that nanostructured gold–cerium oxide catalysts are active for the water–gas shift reaction.¹⁵ Based on the studied results, they found that the metal nanoparticles do not take part in the reaction, and that the reaction is actually catalyzed by non-metallic gold species strongly associated with surface cerium oxygen groups. Han and colleagues prepared low Au-loaded CeO₂ nanoparticle catalysts with high activity *via* several solution chemistries, and used them as catalysts for low-temperature CO oxidation.¹⁶ They discovered that the Au-loaded CeO₂ nanoparticle catalysts exhibit high catalytic activity because of their large surface area as well as the coexistence of Au ^{δ +} (Au⁺ and Au³⁺) and Au⁰ species, in which Au⁺ plays

^aInternational Academy of Targeted Therapeutics and Innovation, Chongqing University of Arts and Sciences, Chongqing 402160, P. R. China

^bChongqing Key Laboratory of Environmental Materials and Remediation Technologies, Research Institute for New Materials Technology, Chongqing University of Arts and Sciences, Chongqing 402160, P. R. China. E-mail: Zhangjin@cqwu.edu.cn; tangdy2008@163.com

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c6ra27207j

a predominant role. Therefore, in order to recognize and design a highly active ceria-based catalyst, it is extremely important to illustrate the nature of the active sites and the catalytic activity of CeO₂-supported Au nanoparticles in detail.

Lattice oxygen atoms¹⁷ and under-coordinated Au atoms in Au nanoparticles¹⁸ have both been reported as reactive species for CO oxidation catalyzed by CeO₂-supported Au nanoparticles. CO oxidation, as a prototypical reaction in heterogeneous catalysis,¹⁹ has attracted enormous attention and has been extensively studied. Kim *et al.* compared CO oxidation by Au₁₃ nanoparticles supported on stoichiometric and partially reduced CeO₂ with the help of DFT+U calculations, and confirmed the critical role of oxygen vacancies in the CeO₂ surface. They also suggested that lowering the vacancy formation energy of the CeO₂ surface is a promising method for enhancing the CO catalytic activity.¹¹ Additionally, analysis shows that CO oxidation by the Mars–van Krevelen mechanism at the interface between doped CeO₂ and supported Au nanoparticles is more effective than that at undoped CeO₂ surfaces.²⁰ In general, for nanocatalysts, it is meaningful to understand the interface effects of metal/support catalysts. The interface between the metal nanoparticles and the oxide supporting materials can cause a pronounced effect on the catalytic activity.^{21–23} Recently, on the basis of a theoretical study, Bing Liu *et al.* explored the three reaction mechanisms for CO oxidation on CeO₂-supported Pd nanorods, that is, the Pd–Ce³⁺ dual site mechanism, M–vK mechanism and Pd-only mechanism.²⁴ By analyzing the calculated results, they found that interface effects play a crucial role and can strongly affect the catalytic activity, which can be understood by considering the structural and electronic properties of the materials. In subsequent work, CO oxidation on CeO₂-supported Pd bimetallic nanorods was investigated by means of a theoretical method and it was found that CO and O₂ can bind at different sites on Pd–Ag/CeO₂ and Pd–Cu/CeO₂.²⁵ The results suggested that both the Pd–Ag and Pd–CeO₂ interfaces in Pd–Ag/CeO₂ can play a vital effect on CO oxidation. However, according to some previous reports,^{4,26} the catalytic activity of oxide-supported Au nanoparticles originates predominantly from the Au nanoparticles themselves, and the oxide surface is regarded as a stabilizer for the Au nanoparticles. Therefore, without doubt, knowledge of the reactive species, including the oxygen vacancies and under-coordinated Au atoms, and the interface between the Au nanoparticles and the supporting material, is meaningful for designing high-performance catalysts and understanding the origin and nature of the catalytic activity.^{27–30}

In order to achieve this goal, the mechanism of ω -alkynylfuran cycloisomerisation on CeO₂(111)-supported subnanometer gold clusters is investigated *via* a theoretical study in this article. Cycloisomerisation is an effective strategy used to obtain phenols and their derivatives, which is extensively used to synthesize natural products, polymers and pharmaceuticals.^{31–36} According to a previous study, Au/CeO₂ can efficiently catalyze cycloisomerisation, leading to a high conversion ratio.³⁵ However, to date, the mechanism of ω -alkynylfuran cycloisomerisation catalyzed by CeO₂-supported gold nanoparticles has not been elaborated or unveiled in detail. Herein, ω -alkynylfuran cycloisomerisation catalyzed by CeO₂-supported gold

nanoparticles (Au₃, Au₄, Au₁₀, Au₁₁) is explored with the help of density functional theory (DFT), as an expansion of our prior research.^{37–39} Gold nanoparticles (Au₃, Au₄, Au₁₀, Au₁₁) were employed for consistence with experimental results reported by Oliver-Meseguer *et al.* and others.^{40–42} In these studies, it was found that small gold clusters (only 3 to 10 atoms in size) can be formed during the catalytic processes, which can catalyze the ω -alkynylfuran cycloisomerisation reaction. In this paper, only the 5-*exo* Friedel–Crafts-type reaction pathway was investigated, and other possible reaction pathways were not considered. The 5-*exo* Friedel–Crafts-type mechanism (FCT) is the most feasible pathway of ω -alkynylfuran cycloisomerisation to produce the targeted phenol derivative product, catalyzed by subnanometer gold clusters and planar gold clusters, based on our previous reports.^{37–39} In order to illustrate the catalytic role of the supporting material, stoichiometric CeO₂ and partially reduced CeO₂ with one vacancy are also considered here. In addition, the possible catalytic sites of Au_{*n*}/CeO₂(111), including the atop, edge and interface sites, are investigated, aiming to illustrate the catalytic activity of the catalytic sites and interfacial effects. Other clusters, such as Pd, Pt or Ag clusters, supported on the surface of CeO₂ will be employed in future work to explore the effect of different clusters on the catalytic properties for ω -alkynylfuran cycloisomerisation.

Computational models and methods

In this work, all of the results were obtained with the help of spin-unrestricted density functional theory (DFT) in a generalized gradient approximation of the PBE functional.⁴³ For geometric optimization, all-electron scalar relativity was chosen, along with the double numerical plus p-polarization (DNP) basis set.⁴⁴ Dmol³ software was employed, which uses a local pseudopotential acting on all electrons, including those in the core, to get scalar relativistic corrections for the relevant valence orbitals. In order to accelerate convergence, a thermal smearing of 0.005 Hartree was applied to the orbital occupation, but all energies were extrapolated to 0 K. Additionally, the transition states were calculated using the linear and quadratic synchronous transit (LST/QST) method, and confirmed by nudged elastic band calculations and relaxed potential energy surface scans.^{45,46} The solvent effect of acetonitrile (dielectric constant: 37.50) was evaluated *via* the conductor-like screening model (COSMO) solvation procedure.^{47–49} All the calculations mentioned above were realized using the Dmol³ software package.^{50,51}

It is well-known that the CeO₂(111) surface is the most thermodynamically stable among the three low-index surfaces of ceria, that is, CeO₂(111), CeO₂(110) and CeO₂(100), implying that the CeO₂(111) surface is the most commonly exposed surface of ceria.^{8,52} Therefore, a 4 × 4 CeO₂(111) slab with 6 atomic layers^{53,54} was employed to describe the stoichiometric CeO₂ surface (CeO₂-s) and the reduced surface (with gold clusters on an oxygen vacancy, CeO₂-v), and 30 Å of vacuum (Fig. 1) was used in order to minimize the interaction between cells. During the geometry optimization, the three upper atomic layers of the CeO₂ slab were relaxed, while the other remaining atomic layers were fixed. Consideration of the enough large



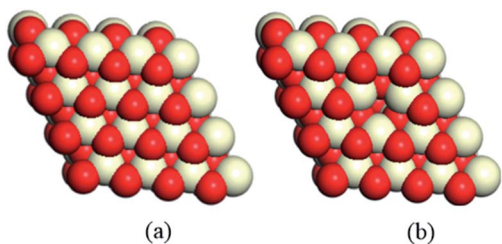


Fig. 1 A 4×4 $\text{CeO}_2(111)$ slab is used to model (a) stoichiometric CeO_2 and (b) partially reduced CeO_2 with one vacancy.

surface cells for all models, only a gamma-point mesh was chosen for Brillouin zone integration. In addition, the bulk equilibrium lattice constant (5.44 Å) was calculated by means of the PBE/DNP method with $8 \times 8 \times 8$ k -points, which is consistent with the experimental values⁵⁵ and previous theoretical studies.^{56–58}

Usually, to correct the resulting electronic structure of reduced ceria, the Hubbard-U term in the DFT+U approach is employed. However, according to a paper reported by Da Silva *et al.*, a reasonable description of the structural parameters, relative energies of different oxides and spectroscopic properties cannot be given at the same time using a unique value of U; therefore, it is not clear whether the DFT+U method provides a systematic improvement of the energetics.⁵⁹ In addition, in Bernard Delley and coworkers' reports, the stability and morphology of cerium oxide surfaces in an oxidizing environment, the adsorption and thermodynamic stability of CH_x ($x = 1-4$) on a $\text{CeO}_2(111)$ surface, and water adsorption on stoichiometric and reduced $\text{CeO}_2(111)$ surfaces were explored by employing the DFT-GGA approach using Dmol³ software and the results were reliable and comparable.^{60–62} The same choice was made to study CO adsorption on CeO_2 and investigate the interaction of water with stoichiometric and reduced CeO_2 surfaces by Yang *et al.* and Kumar *et al.*, respectively.^{63,64} Hence, in view of these considerations, the DFT-GGA approach is also adopted in this article.

Additionally, herein, the turnover frequency (TOF) was employed to evaluate the efficiency of a catalytic cycle. The energetic span model can be used to predict the TOF from theoretically obtained free energy profiles, which was proposed by Shaik and co-workers.^{65–68} The Gibbs free energy is related to the corrected enthalpy ($\Delta H_{\text{correct}}$) and entropy (ΔS). These two parameters were not considered because numerical frequency computations are difficult for large system in Dmol³. On the basis of this model, the TOF-determined transition state (TDTS) and intermediate (TDI) can be determined by evaluating the degree of TOF control ($X_{\text{TOF},i}$), as shown in eqn (1):

$$X_{\text{TOF},i} = \left| \frac{1}{\text{TOF}} \frac{\partial \text{TOF}}{\partial E_i} \right| \quad (1)$$

E_i denotes the energy of the transition state or intermediate. The larger the X_{TOF} value, the higher the influence of the corresponding state (transition state or intermediate) on the TOF. The TOF can be calculated using eqn (2):

$$\text{TOF} = \frac{\kappa_B T}{h} e^{-\delta E/RT} \quad (2)$$

where δE is the energetic span, which can be defined as shown in eqn (3):

$$\delta E = \begin{cases} E_{\text{TDTS}} - E_{\text{TDI}} & \text{if TDTS appears after TDI} \\ E_{\text{TDTS}} - E_{\text{TDI}} + \Delta E_r & \text{if TDTS appears before TDI} \end{cases} \quad (3)$$

All of the kinetic information was obtained by applying the energetic span model using the AUTOF program.⁶⁷

Results and discussion

Morphology of gold clusters on $\text{CeO}_2(111)$

Before the reaction mechanism of ω -alkynylfuran cycloisomerisation catalyzed by Au_n/CeO_2 was unveiled, the atomic structures of the gold clusters on the $\text{CeO}_2(111)$ surface were investigated. For Au_3 and Au_4 clusters supported on $\text{CeO}_2(111)$ surfaces, all possible configurations are taken into account. However, for Au_{10} and Au_{11} clusters supported on $\text{CeO}_2(111)$ surfaces, only the fcc-like and hcp-like clusters are considered, as these have been proven to be the stable structures of gold clusters on $\text{CeO}_2(111)$ surfaces.⁶⁹ Three structures for Au_3 clusters on stoichiometric $\text{CeO}_2(111)$ surfaces, including triangle perpendicular to the surface, triangle parallel to the surface and linear parallel to the surface, are considered. According to the calculated results, it is demonstrated that the triangle perpendicular to the $\text{CeO}_2(111)$ surface structure is the most stable among the three configurations. Nevertheless, only two configurations, namely, triangle perpendicular to and linear parallel to the surface Au_3 clusters on reduced $\text{CeO}_2(111)$ surfaces, were found, because during optimization the triangle parallel to the surface configuration converged to the triangle perpendicular to the surface configuration. Similarly, the triangle structure is also more stable than the linear structure. For Au_4 clusters on both stoichiometric and reduced $\text{CeO}_2(111)$ surfaces, three different structures were revealed based on the theoretical calculations. The computed results show that the tetrahedral Au_4 cluster is the most stable structure on both stoichiometric and reduced $\text{CeO}_2(111)$ surfaces. The stable structures of Au_3 and Au_4 clusters on $\text{CeO}_2(111)$ surfaces are in line with previous theoretical studies using PBE+U ($U = 5.0$ eV).⁶⁹ In addition, the tetrahedral Au_{10} cluster is more stable than the hemispherical structures, which is also in agreement with the PBE+U (U

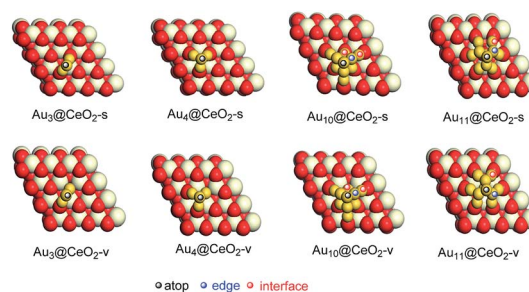


Fig. 2 The most stable structures of gold clusters on $\text{CeO}_2(111)$ and possible catalytic sites of $\text{Au}_n/\text{CeO}_2(111)$.



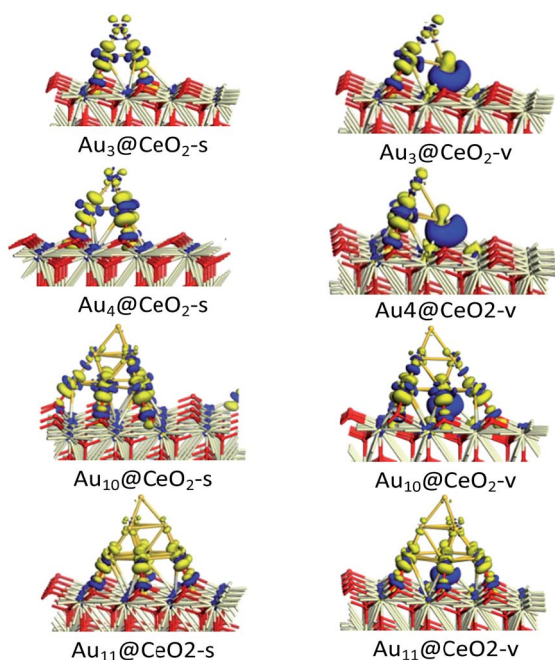
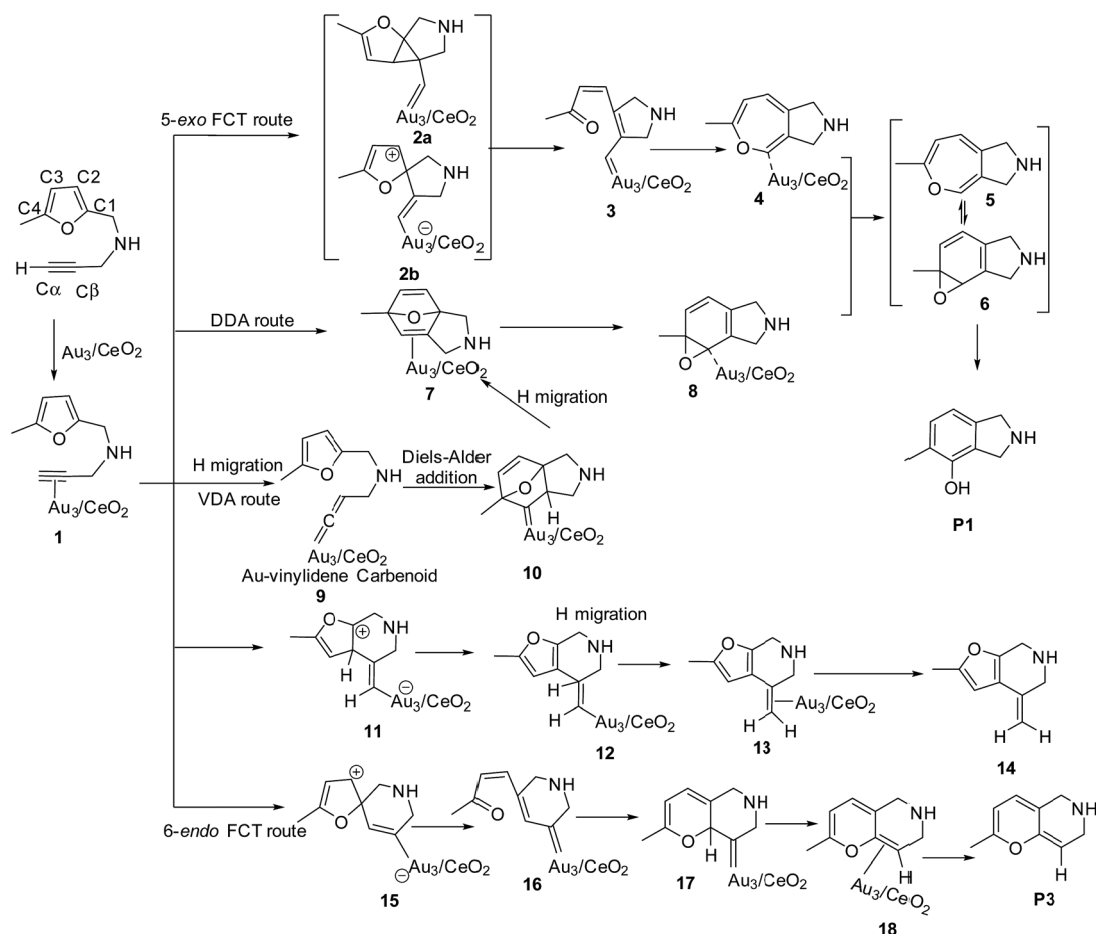


Fig. 3 Charge density difference of $Au_n/CeO_2(111)$ (blue and yellow isosurfaces represent increasing and decreasing electron densities, respectively; isovalue = 0.03).

= 5.0 eV) study.⁶⁹ It is well known that, for Au_4 – Au_{10} clusters, the most stable isomers are planar structures in the gas phase, indicating that the CeO_2 support results in structural changes from planar to three-dimensional. The corresponding optimized structures are shown in Fig. S1† and the most stable structures are also exhibited in Fig. 2, together with the possible catalytic sites.

Electronic properties of $Au_n/CeO_2(111)$

To illustrate the electronic properties of the interactions between Au clusters and CeO_2 surfaces, the charge density differences of $Au_n/CeO_2(111)$ were calculated, as shown in Fig. 3. According to Fig. 3, it can be concluded that significant charge transfer occurs between the Au cluster and the CeO_2 support. The electronic interaction is largely restricted to the bottom Au atoms of the cluster, which are in direct contact with the CeO_2 support, indicating that these atoms have been oxidized. In contrast, little decrease in the electron density is observed for the top Au layer, illustrating that this layer is much less affected electronically by the interaction with the CeO_2 surface. In addition, analysis of charge transfer between the Au cluster and the CeO_2 support indicates that the increase in electron density for $Au_n@CeO_2-v$ mainly occurs near the oxygen vacancy, and is obviously different to that seen for $Au_n@CeO_2-s$. These results indicate that charge transfer



Scheme 1 The 5-*exo* Friedel–Crafts-type (FCT) route, direct Diels–Alder-type (DDA) route and Au-vinylidene carbenoid (VDA) route to form the phenol derivative P1, the formation of β -alkenylated furan P2, and the 6-*endo* Friedel–Crafts-type (FCT) route toward the pyran derivative P3.



happens at the interface between the Au cluster and the CeO₂ support, and the presence of an oxygen vacancy can have a large effect on charge transfer.

The possible mechanisms for ω -alkynylfuran cycloisomerisation catalyzed by Au₃/CeO₂

To illustrate the most feasible pathway for ω -alkynylfuran cycloisomerisation, Au₃/CeO₂ was selected as a model catalyst to simulate the detailed mechanism of ω -alkynylfuran cycloisomerisation. The possible pathways of ω -alkynylfuran cycloisomerisation catalyzed by Au₃/CeO₂ were designed and are shown in Scheme 1. As shown in Scheme 1, after the adsorption of the substrate on Au₃/CeO₂, there are three possible pathways to produce phenol derivatives. The first one is the 5-*exo* Friedel-Crafts-type reaction route starting from the nucleophilic addition of a C β atom to the C1=C2 bond or C1 site of the furan ring (abbreviated as 5-*exo* FCT). The second pathway is [4 + 2] cycloaddition between the C \equiv C bond and the C1=C2–C3=C4 of the furan ring to produce the final product (abbreviated as DDA). The last pathway is direct hydrogen transfer from the C α atom to the C β atom to form an Au-vinylidene carbenoid intermediate, and subsequent Diels–Alder addition of the C α –C β double bond to the furan ring (abbreviated as VDA). Additionally, there is a 6-*endo* FCT pathway, which involves the nucleophilic addition of a C α atom to the furan ring in **IM1** in competition with the 5-*exo* FCT pathway, finally leading to a pyran derivative. Moreover, the C β atom can attack the C2 atom of the furan ring, finally obtaining a β -alkenylated furan *via* H migration. The energy profiles for all of the possible mechanisms are depicted in Fig. 4–7. For the VDA route, H

migration requires a large energy barrier to be overcome, and therefore it is not taken into account. According to Fig. 4–7, the rate-limiting step of the 5-*exo* FCT reaction route in the gas phase is the nucleophilic addition of the C β atom to the C1 site of the furan ring to form the spiro-intermediate **IM2**, which has an energy barrier of 13.66 kcal mol^{–1}, while the formation of **IM4** is the rate-determining step in acetonitrile. However, for the DDA pathway, the formation of the β -alkenylated furan and the 6-*endo* FCT pathway, the energy barriers of the rate-limiting steps are 46.28 (44.80), 28.79 (27.86) and 35.26 (35.87), respectively. Obviously, the 5-*exo* FCT reaction route is the most feasible pathway for ω -alkynylfuran cycloisomerisation to produce the targeted phenol derivative product.

Mechanism of ω -alkynylfuran cycloisomerisation catalyzed by Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111)

On the basis of our previous reports^{37–39} and the above discussions, the 5-*exo* Friedel–Crafts-type mechanism (FCT) is the most feasible pathway for ω -alkynylfuran cycloisomerisation. Accordingly, the 5-*exo* FCT mechanism of ω -alkynylfuran cycloisomerisation on the various reaction sites of Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111) were investigated. The corresponding reaction sites are exhibited in Fig. 2. The optimized structures and related parameters of the reaction catalyzed by Au₁₀/CeO₂(111)-s are shown in Fig. 8, and the corresponding structures and parameters for the other Au_{*n*}/CeO₂ materials are plotted in Fig. S2–S14 (ESI†). Moreover, the potential energy profiles are presented in Fig. 9–12.

Firstly, ω -alkynylfuran cycloisomerisation on the atop sites of Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111) is discussed. As shown in

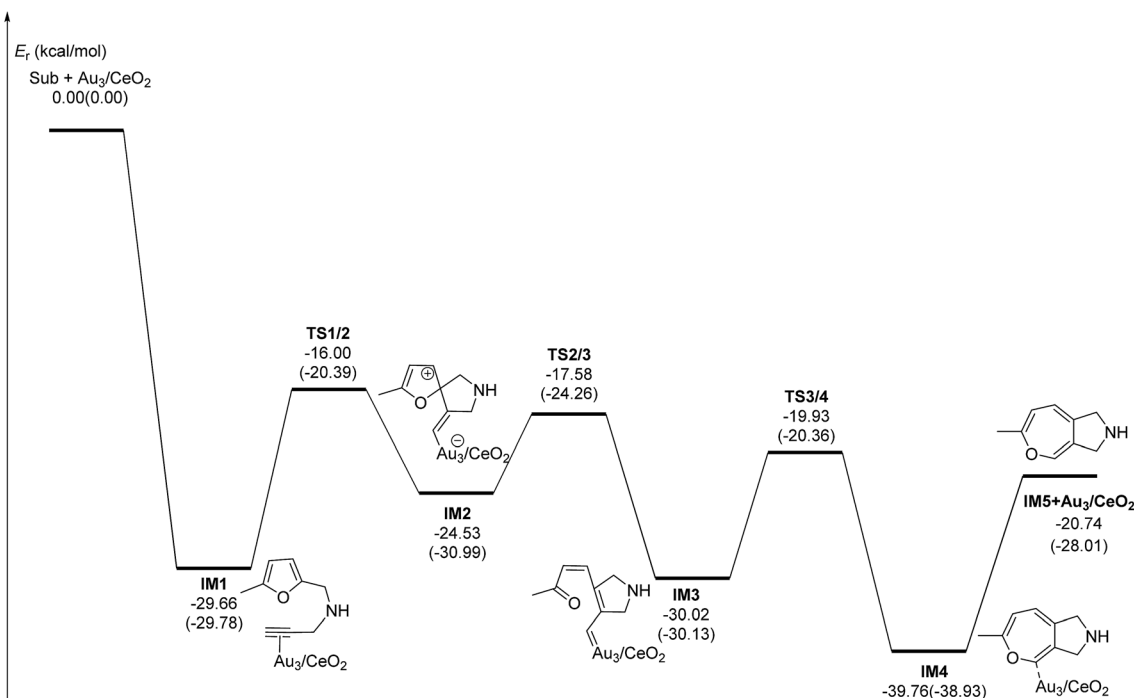


Fig. 4 Energy profile of the 5-*exo* FCT pathway that was used to form the phenol derivative **P1** in the gas phase and acetonitrile solvent (parentheses).



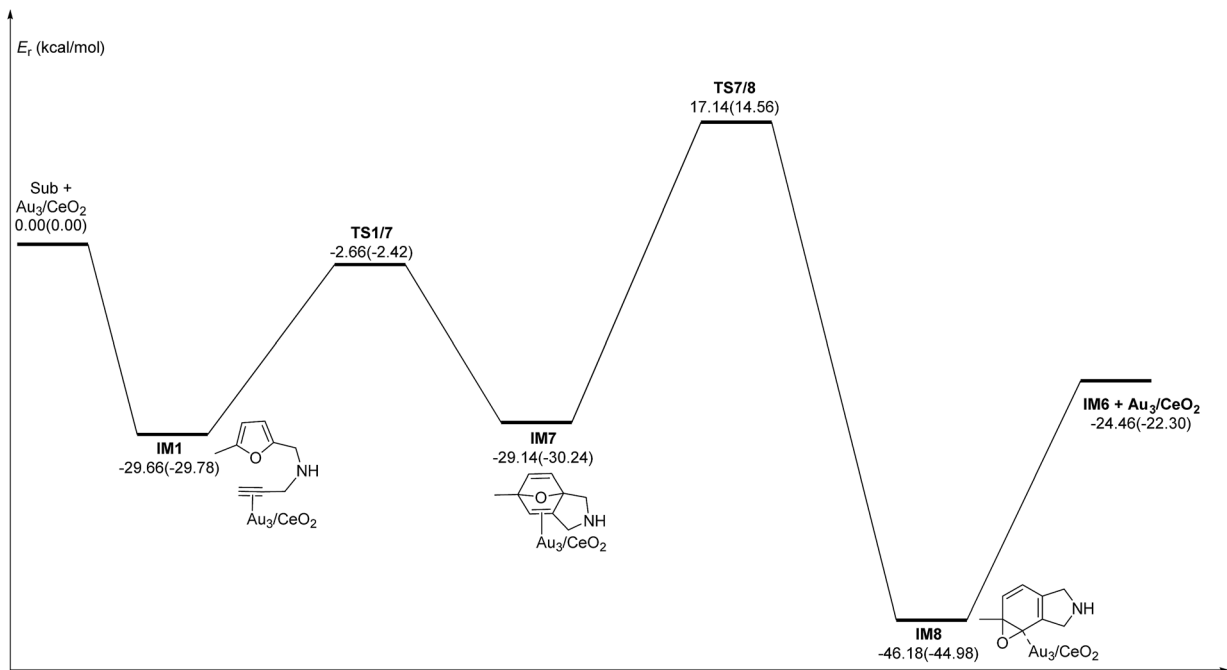


Fig. 5 Energy profile of the DDA pathway that was used to form the phenol derivative P1 in the gas phase and acetonitrile solvent (parentheses).

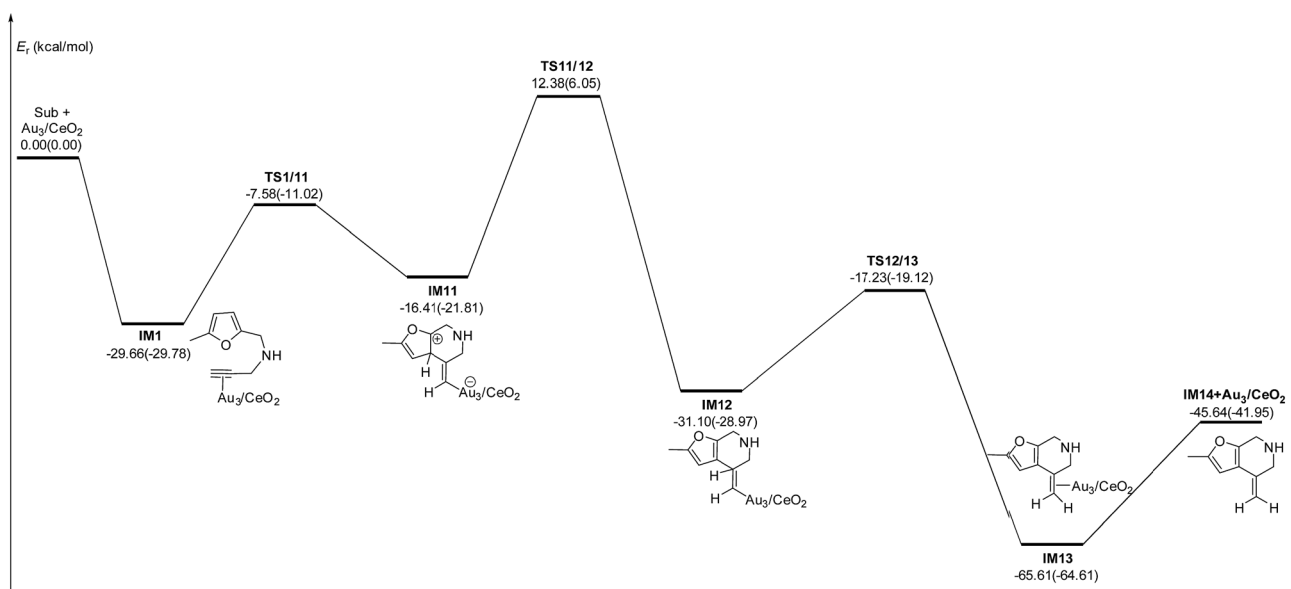


Fig. 6 Energy profile for the formation of the β -alkenylated furan P2 in the gas phase and acetonitrile solvent (parentheses).

Fig. 5–8, the adsorption of ω -alkynylfuran on the atop sites is exothermic by about $15\text{--}18\text{ kcal mol}^{-1}$ in the gas phase and acetonitrile solvent (Ca (catalyst) + Sub (substrate) \rightarrow **IM1** (intermediate)). The adsorption energies of the substrates on the atop sites of the gold clusters supported on the $\text{CeO}_2(111)$ surfaces are comparable with those on subnanometer gold clusters (Au_{38} , Au_{27} , Au_{28} , Au_{30} , Au_{35} and Au_{55}).³⁸ In addition, analysis of the computed adsorption energies exhibited in Fig. 5–8 suggests that the surface O vacancy of $\text{CeO}_2(111)$ only has a slight influence on the adsorption of the substrate on the

atop sites. Once the ω -alkynylfuran is adsorbed on the atop sites of the gold clusters supported on the $\text{CeO}_2(111)$ surfaces, the C β atom electrophilically attacks the furan ring to give rise to a spiro-intermediate, **IM2** (**IM1** \rightarrow **IM2**). This step is kinetically feasible, with an energy barrier of about 5 kcal mol^{-1} . In addition, the **IM1** \rightarrow **IM2** conversion step is exothermic by about $5\text{--}8\text{ kcal mol}^{-1}$ in acetonitrile solvent. Subsequently, the cleavage of the C–O bond of the furan ring in **IM2** produces a linear conjugate triene ketone intermediate (**IM3**) through transition state **TS2/3** with an energy barrier of $8\text{--}9\text{ kcal mol}^{-1}$. This step is



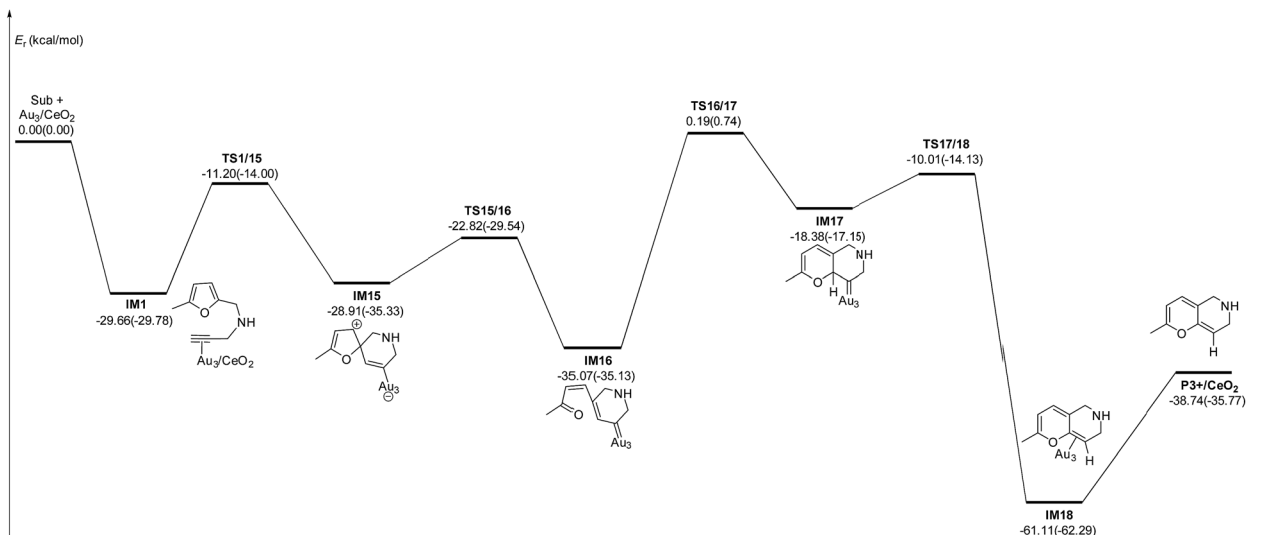


Fig. 7 Energy profile of the 6-*endo* Friedel–Crafts-type (FCT) pathway that was used to form the pyran derivative **P3** in the gas phase and acetonitrile solvent (parentheses).

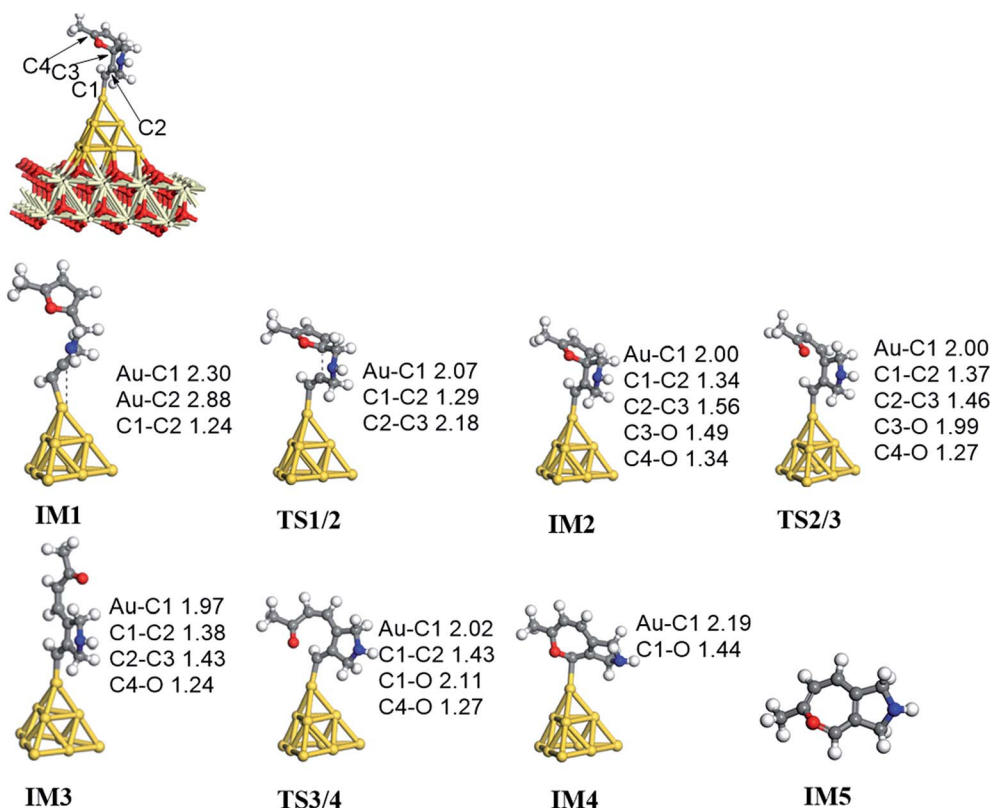


Fig. 8 The optimized structures of ω -alkynylfuran cycloisomerisation catalyzed by the atop site of $\text{Au}_{10}/\text{CeO}_2(111)\text{-s}$.

nearly thermoneutral in acetonitrile solvent. In the next step, an oxygen atom from a carbonyl group approaches the Au–C bond yielding an oxepine intermediate (**IM4**), which is exothermic in both the gas phase and acetonitrile solvent with an energy barrier of about 8–11 kcal mol^{−1}. Finally, oxepine is desorbed from the oxepine intermediate to recover the catalyst. This step

is endothermic by about 8–18 kcal mol^{−1}. The oxepine then quickly converts into the phenol product with an exothermicity of about 40 kcal mol^{−1} upon heating or exposing to acid.^{33,34,70} Additionally, on the basis of the energetic span model by Shaik and coworkers, the TOF values of ω -alkynylfuran cycloisomerisation on the atop, interface-corner, interface-edge and



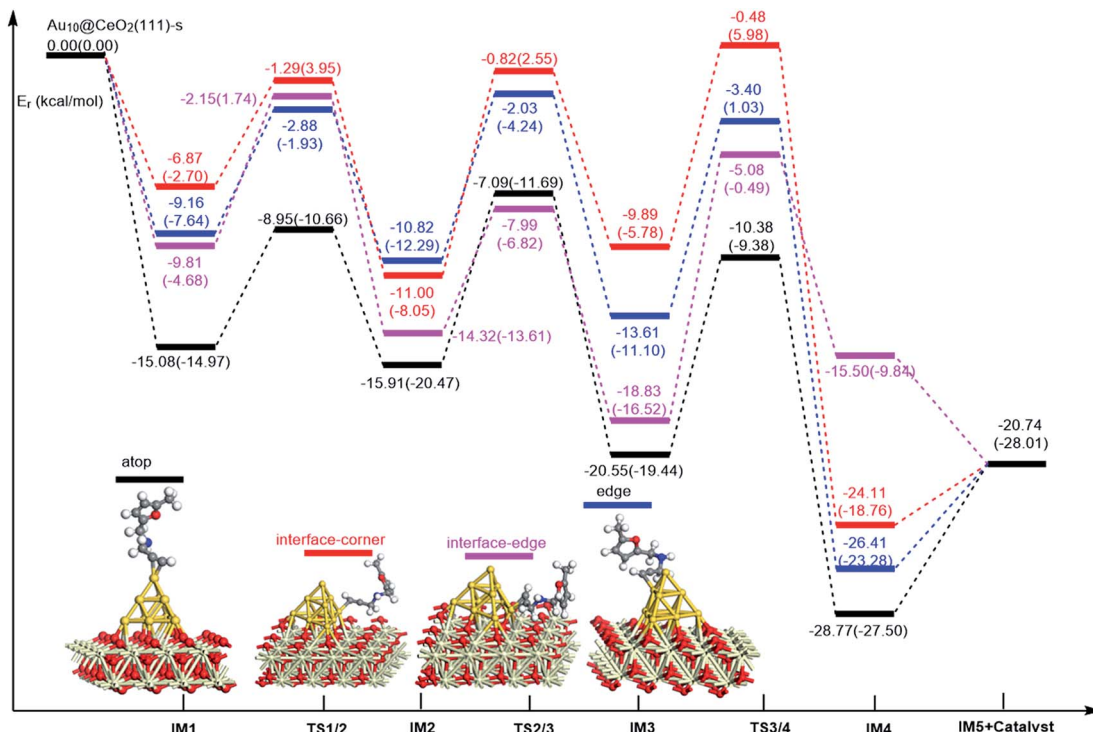


Fig. 9 Energy profile for ω -alkynylfuran cycloisomerisation on stoichiometric CeO_2 -supported Au_{10} .

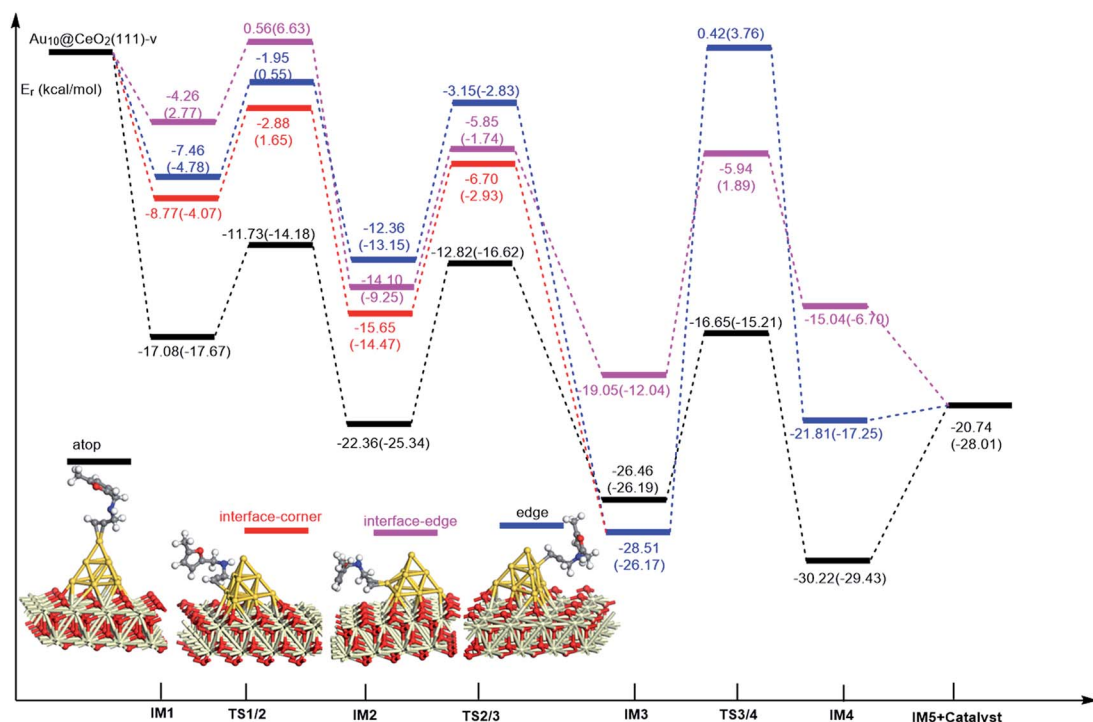


Fig. 10 Energy profile for ω -alkynylfuran cycloisomerisation on reduced CeO_2 -supported Au_{10} .

edge sites are calculated and shown in the Table 1. Compared with ω -alkynylfuran cycloisomerisation on the interface-corner, interface-edge and edge sites, the atop site has larger TOF values of about 10^5 to 10^6 s^{-1} , indicating that the atop site possesses high

catalytic activity. Clearly, the catalytic activity of the atop sites is similar to that of subnanometer gold clusters, due to the similar binding characters of the gold atoms. Acetonitrile solvent can have a slight influence on the energy barriers and the reaction rates.

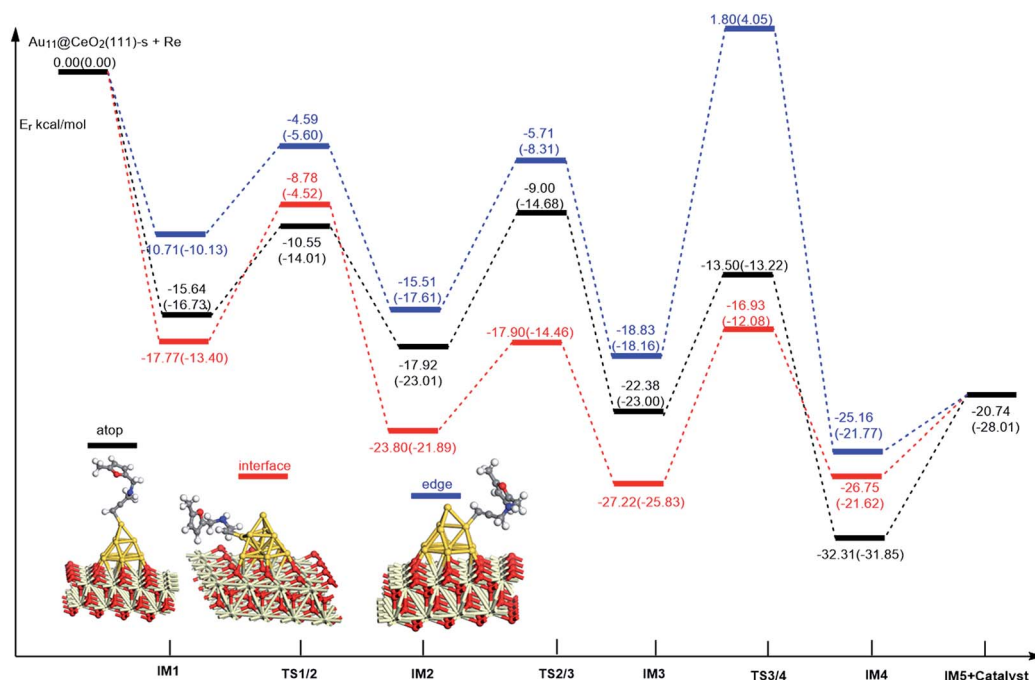


Fig. 11 Energy profile for ω -alkynylfuran cycloisomerisation on stoichiometric CeO_2 -supported Au_{11} .

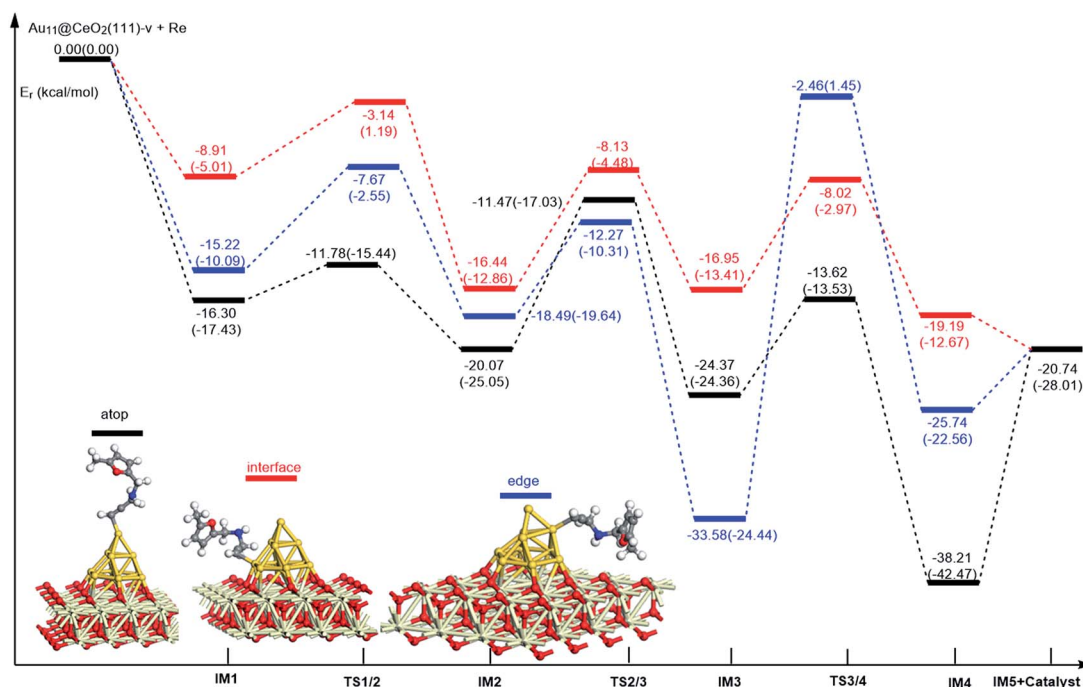


Fig. 12 Energy profile for ω -alkynylfuran cycloisomerisation on reduced CeO_2 -supported Au_{11} .

Subsequently, ω -alkynylfuran cycloisomerisation on the edge sites of $\text{Au}_{10}/\text{CeO}_2(111)$ and $\text{Au}_{11}/\text{CeO}_2(111)$ were investigated and discussed. The adsorption energies are smaller as compared to that of the atop sites because of the higher coordination numbers of the edge sites. Interestingly, a distinctive substrate adsorption behavior is observed for the edge site of $\text{Au}_{11}/\text{CeO}_2(111)$ -v. As shown in Fig. S14,[†] the adsorption of the

substrate on the edge site of $\text{Au}_{11}/\text{CeO}_2(111)$ -v induces strong structural deformation of the Au_{11} fragment due to the strong interaction between the adsorbed alkyne group and the Au atoms. The corresponding structural deformation raises the energy barrier for the electrophilic attack of the C atom on the furan ring. The electrophilic attack of the C atom on the furan ring ($\text{IM1} \rightarrow \text{IM2}$) and the cleavage of C–O in the furan ring

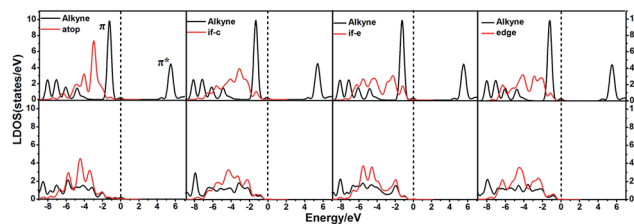


Table 1 The energetic span (δE) and TOF values for ω -alkynylfuran cycloisomerisation catalyzed by $\text{Au}_{10}/\text{CeO}_2(111)$ and $\text{Au}_{11}/\text{CeO}_2(111)$

	TDI/TDTS	δE (kcal mol $^{-1}$)	TOF (s $^{-1}$)
$\text{Au}_{10}/\text{CeO}_2(111)$-s			
Atop	IM2/TS3/4	11.09	2.9×10^5
Interface-corner	IM2/TS3/4	14.03	4.1×10^3
Interface-edge	IM3/TS3/4	16.03	4.3×10^2
Edge	IM2/TS3/4	13.32	2.0×10^4
$\text{Au}_{10}/\text{CeO}_2(111)$-v			
Atop	IM3/TS3/4	10.98	3.3×10^5
Interface-corner	IM3/TS3/4	29.93	1.6×10^{-7}
Interface-edge	IM3/TS3/4	13.93	4.9×10^3
Edge	IM3/TS3/4	29.93	1.6×10^{-7}
$\text{Au}_{11}/\text{CeO}_2(111)$-s			
Atop	IM2/TS3/4	9.79	1.2×10^6
Interface	IM3/TS3/4	13.75	6.5×10^3
Edge	IM3/TS3/4	22.21	1.3×10^{-2}
$\text{Au}_{11}/\text{CeO}_2(111)$-v			
Atop	IM4/IM5	14.46	1.1×10^3
Interface	IM3/TS3/4	10.44	6.6×10^5
Edge	IM3/TS3/4	25.89	7.1×10^{-5}

(IM2 \rightarrow IM3) steps on the edge sites have similar energy barriers. The closure of the ring to form the oxepine intermediate on the edge site of $\text{Au}_{10}/\text{CeO}_2(111)$ -s needs to overcome an energy barrier of 10.21 (12.12) kcal mol $^{-1}$ in the gas phase (acetonitrile solvent), which is comparable to the situation on the atop site. However, the closure of the ring to form the oxepine intermediate on the edge site of $\text{Au}_{10}/\text{CeO}_2(111)$ -v and $\text{Au}_{11}/\text{CeO}_2(111)$ -v entails a large energy barrier of about 29 kcal mol $^{-1}$ due to the μ^2 -coordination in IM3. Simultaneously, the closure of the ring to form the oxepine intermediate on the edge site of $\text{Au}_{11}/\text{CeO}_2(111)$ -s must overcome an energy barrier of about 20.00 kcal mol $^{-1}$ due to the large deformation of the substrate fragment. Hence, only the edge sites of $\text{Au}_{10}/\text{CeO}_2(111)$ -s show similar catalytic activity toward ω -alkynylfuran cycloisomerisation to the atop sites.

Finally, the 5-*exo* FCT mechanism can be extended to the interface sites of $\text{Au}_{10}/\text{CeO}_2(111)$ and $\text{Au}_{11}/\text{CeO}_2(111)$, so ω -alkynylfuran cycloisomerisation on the interface sites of $\text{Au}_{10}/\text{CeO}_2(111)$ and $\text{Au}_{11}/\text{CeO}_2(111)$ is also explored. As seen from Fig. 9–12, there are two types of interface sites for $\text{Au}_{10}/\text{CeO}_2(111)$ -s and $\text{Au}_{10}/\text{CeO}_2(111)$ -v. In contrast, the interface sites of $\text{Au}_{11}/\text{CeO}_2(111)$ -s and $\text{Au}_{11}/\text{CeO}_2(111)$ -v have similar bonding character, and hence there is only one type of interface site. The adsorption of the substrate on the interface sites causes a large structural deformation of the supported gold clusters due to the strong interaction between the alkyne group and gold atoms. The large structural deformation of the supported gold clusters counteracts the partial stabilizing interaction between the alkyne group and the gold atoms, which results in lower adsorption energies of the substrate on the interface sites. For the interface sites, the closure of the ring to form the oxepine intermediate is the rate-limiting step in both the gas phase and acetonitrile solvent. The interface-corner

**Fig. 13** The local density of states (LDOS) projected on triplet bonds on $\text{Au}_{10}/\text{CeO}_2(111)$ -s for the adsorption of the substrate at different sites, together with the d-projected LDOS of $\text{Au}_{10}/\text{CeO}_2(111)$ -s of the Au atom at different sites. Free C–C triplet bond and $\text{CeO}_2(111)$ (top), and adsorption (bottom). The Fermi level is set to zero (if-c = interface-corner, if-e = interface-edge).

sites of $\text{Au}_{10}/\text{CeO}_2(111)$ -v may be viewed as inactive toward ω -alkynylfuran cycloisomerisation compared to the other interface sites, due to the formation of μ^2 -coordination in IM3. The catalytic activity of the interface sites of $\text{Au}_{10}/\text{CeO}_2(111)$ is lower than that of the atop sites. Acetonitrile solvent enhances the energy barrier of the closure of the ring to form the oxepine intermediate and reduces the reaction rate.

In order to illustrate the adsorption properties of ω -alkynylfuran on the different sites of $\text{Au}_{10}/\text{CeO}_2(111)$, the corresponding local density of states (LDOS) are calculated and shown in Fig. 13 and S15.† In this article, the LDOS discussion only focuses on the d-band information because the valence orbitals are the s and d orbitals of gold. Moreover, the interaction between the clusters and the ω -alkynylfuran depends on π -back-donation and σ -coordinate bonding, that is, charge transfer between the d and π^* orbitals or π and d^* orbitals. Hence, the d-band information is mainly investigated in this article.† According to Fig. 13 and S15,† the bands of the adsorbed alkyne and $\text{Au}_{10}/\text{CeO}_2(111)$ moved to the left toward the lower energy region, compared with the free substrate and $\text{Au}_{10}/\text{CeO}_2(111)$. The main peak of the alkyne overlapped with those of the different sites of $\text{Au}_{10}/\text{CeO}_2(111)$, indicating chemical bond formation between the substrate and $\text{Au}_{10}/\text{CeO}_2(111)$. Compared with the interface-corner, interface-edge and edge sites, the band of the adsorbed alkyne is more down-shifted for the atop site and the main peaks of the alkyne are more effectively overlapped with the atop site, leading to a larger adsorption energy. In addition, the adsorption energies of ω -alkynylfuran on the interface-edge sites of $\text{Au}_{10}/\text{CeO}_2(111)$ (stoichiometric and reduced) exhibit obvious differences, which can be attributed to the intensity and overlap degree of the bands.

Similarly, the local density of states (LDOS) are calculated again to elaborate the adsorption of ω -alkynylfuran on $\text{Au}_{11}/\text{CeO}_2(111)$ (stoichiometric and reduced). Based on the computed results, the adsorption energy of ω -alkynylfuran on the interface sites is larger than that on the atop and edge site owing to the better overlap of the peaks, which can be seen from Fig. S16.† Compared with the edge sites, the band of the adsorbed alkyne is more down-shifted and there is better overlap of the peaks for the alkyne and the atop Au atoms, leading to larger adsorption energies. However, for the adsorption of ω -alkynylfuran on $\text{Au}_{11}/$



CeO₂(111) (vacancy, Fig. S17†), the adsorption energy is the smallest at the interface sites due to the weak d peak at around 8 eV and the relatively weak overlap of the bands.

In summary, according to the above-mentioned analyses, the atop and interface sites exhibit high catalytic activity toward ω -alkynylfuran cycloisomerisation *via* the FCT mechanism. The edge sites of Au₁₀/CeO₂(111)-s show similar catalytic activity toward ω -alkynylfuran cycloisomerisation to the atop sites. The adsorption energies of the substrate on the atop sites of the gold clusters supported by the CeO₂(111) surfaces are comparable with those on subnanometer gold clusters. Due to the higher coordination number of the edge and interface sites, lower adsorption energies of the substrate on the edge and interface sites were identified. In addition, based on the calculated results, it is found that the surface O vacancy of CeO₂(111) has some negative and positive influences on the adsorption energies of the substrate and the catalytic activity.

Comparison of the catalytic activity of free gold clusters and supported gold clusters

In order to clarify the role of the CeO₂ support, the 5-*exo* FCT pathway was studied on free and supported Au₃₋₄ clusters. As plotted in Fig. 14 and 15, the reaction schemes of ω -alkynylfuran cycloisomerisation on CeO₂-supported Au₃₋₄ and free Au₃₋₄ are similar to those on Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111). The adsorption of ω -alkynylfuran on the CeO₂-supported Au₃₋₄ and free Au₃₋₄ clusters generates intermediate **IM1**. An analysis of the adsorption of ω -alkynylfuran on the CeO₂-supported Au₃₋₄ and free Au₃₋₄ clusters suggests that the adsorption energies are much larger as compared to those of

ω -alkynylfuran adsorption on Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111), due to the lower coordination numbers of the Au₃₋₄ clusters. The large adsorption energies have a pronounced effect on the C β atom of the alkynyl group, which moves close to the C1 atom of the furan ring. For ω -alkynylfuran cycloisomerisation on Au₃/CeO₂(111)-s, Au₃/CeO₂(111)-v and free Au₃ clusters, overcoming the energy barriers of 13.66 (9.39), 13.76 (10.13) and 16.45 (12.17) kcal mol⁻¹ in the gas phase (acetonitrile solvent), the C β -C1 bond is formed in the spiro-intermediate **IM2**. These corresponding energy barriers are larger than those on Au₁₀/CeO₂(111) and Au₁₁/CeO₂(111). Furthermore, this elementary reaction on free Au₃ needs to overcome the highest energy barrier. Next, the C1-O bond of the furan ring is broken to form the Au-vinylidene intermediate **IM3**. The corresponding energy barriers are 6.95 (6.73), 6.48 (5.29) and 5.75 (5.22) kcal mol⁻¹ in the gas phase (acetonitrile solvent) on Au₃/CeO₂(111)-s, Au₃/CeO₂(111)-v and free Au₃ clusters, respectively. The reaction that breaks the furan ring is exothermic by about 5.49 (-0.86), 7.12 (2.45) and 9.39 (4.61) kcal mol⁻¹ in the gas phase (acetonitrile solvent) on Au₃/CeO₂(111)-s, Au₃/CeO₂(111)-v and free Au₃. Then, the O atom of the carbonyl group attacks the C α atom of the Au-vinylidene with energy barriers of 10.09 (9.77), 10.21 (11.02) and 9.54 (9.09) kcal mol⁻¹ in the gas phase (acetonitrile solvent) on Au₃/CeO₂(111)-s, Au₃/CeO₂(111)-v and free Au₃ clusters, leading to the formation of a seven-membered ring intermediate (**IM4**).

Additionally, the mechanisms of ω -alkynylfuran cycloisomerisation catalyzed by Au₄/CeO₂(111)-s, Au₄/CeO₂(111)-v and free Au₄ clusters are also unveiled. As shown in Fig. 15, the catalytic cycles of ω -alkynylfuran cycloisomerisation on Au₄/CeO₂(111)-s, Au₄/CeO₂(111)-v and free Au₄ clusters are similar to

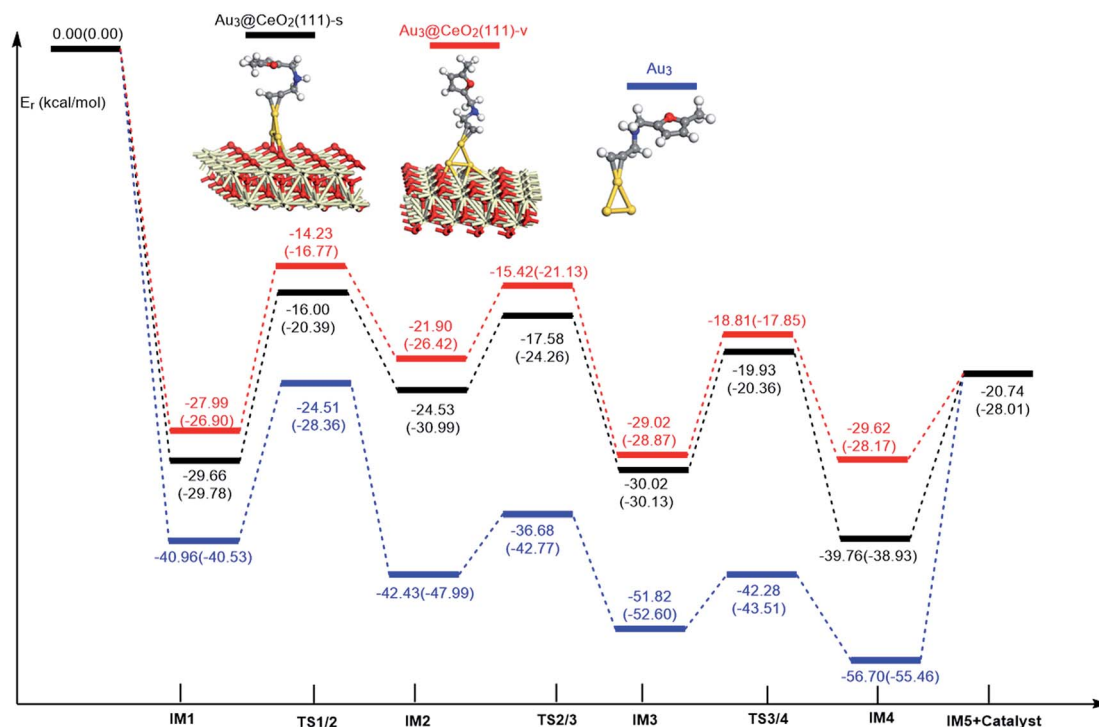


Fig. 14 Energy profile for ω -alkynylfuran cycloisomerisation on CeO₂-supported Au₃ and free Au₃.



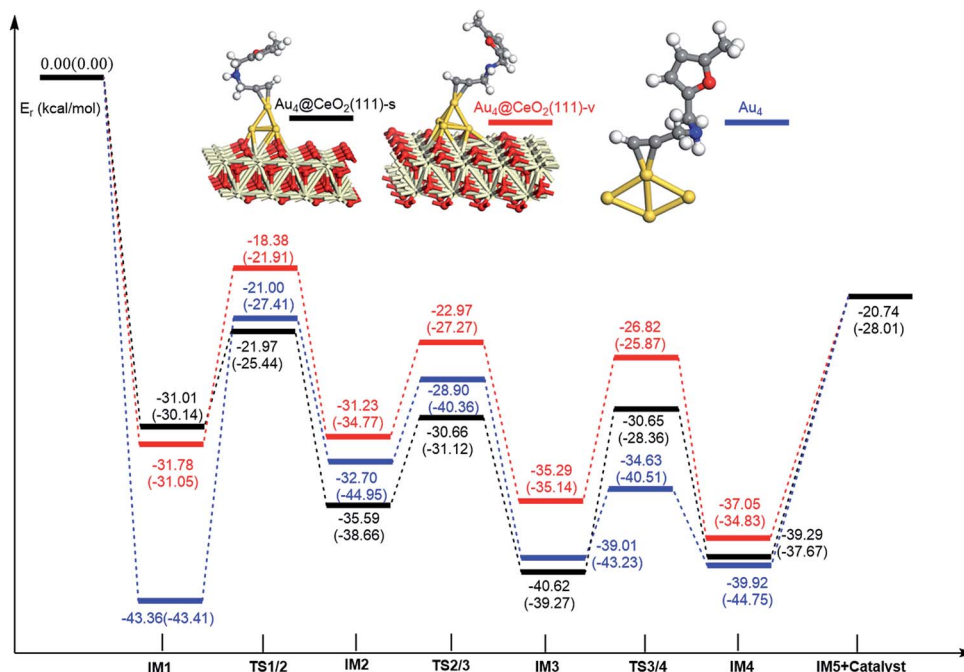


Fig. 15 Energy profile for ω -alkynylfuran cycloisomerisation on CeO_2 -supported Au_4 and free Au_4 .

those on $\text{Au}_{10}/\text{CeO}_2(111)$, $\text{Au}_{11}/\text{CeO}_2(111)$, $\text{Au}_3/\text{CeO}_2(111)$ and free Au_3 , including the adsorption of the ω -alkynylfuran, the formation of the C β –C1 bond, the breakage of the C1–O bond in the furan ring, and the formation of the seven-membered ring. On the basis of the energy profiles, the supported Au_4 clusters are clearly more active than the free Au_4 clusters toward ω -alkynylfuran cycloisomerisation.

According to the calculated results, the rate-determining steps of ω -alkynylfuran cycloisomerisation on CeO_2 -supported Au_3/Au_4 and free Au_3/Au_4 are the formation of the C β –C1 bond, that is, the **IM1** \rightarrow **IM2** conversion. Compared with free Au_3 , the CeO_2 supporting material can efficiently facilitate ω -alkynylfuran cycloisomerisation, decreasing the energy barriers for **IM1** \rightarrow **IM2** conversion to 2.69–2.79 (2.04–2.78) kcal mol^{-1} in the gas phase (acetonitrile solvent). In addition, the oxygen vacancy can have a minor effect on the energy barriers for ω -alkynylfuran cycloisomerisation on Au_3/CeO_2 in the gas phase (acetonitrile solvent). Similarly, compared with free Au_4 , CeO_2 -supported Au_4 can decrease the energy barriers of the rate-determining step to 8.96–13.28 (6.86–11.30) kcal mol^{-1} in the

gas phase (acetonitrile solvent), indicating the active role of CeO_2 in ω -alkynylfuran cycloisomerisation.

In order to better illustrate the catalytic reactivity of the free gold clusters and supported gold clusters, the corresponding energetic span (δE) and TOF values were calculated and the values are given in Table 2. As shown in Table 2, the TOF values for ω -alkynylfuran cycloisomerisation catalyzed by free Au_{3-4} are smaller than those for $\text{Au}_{3-4}/\text{CeO}_2(111)$. Therefore, the presence of $\text{CeO}_2(111)$ can effectively facilitate ω -alkynylfuran cycloisomerisation.

The role of CeO_2 surface

Based on the calculated results, the presence of the $\text{CeO}_2(111)$ surface can largely decrease the adsorption energies. In order to illustrate this interesting phenomenon, the corresponding local density of states (LDOS) were computed and plotted in Fig. 16. As shown in Fig. 16, for the adsorption of ω -alkynylfuran on Au_3 clusters, there is better peak overlap between the Au_3 cluster and the alkyne as compared to that for $\text{Au}_3/\text{CeO}_2(111)$ (stoichiometric and reduced) and Au_3 clusters, leading to a larger adsorption energy. The different orbital interactions can be attributed to the charge transfer between the $\text{CeO}_2(111)$ surface and the Au_3 cluster. Similarly, for $\text{Au}_4/\text{CeO}_2(111)$ (stoichiometric and reduced) and Au_4 , the local density of states (LDOS) shown in Fig. S24[†] clearly show that the peak overlap plays a crucial role in determining the adsorption energies. For free Au_4 clusters, the main peak of the alkyne can effectively overlap with the d-band of the Au_4 cluster, facilitating the interaction between the alkyne and the Au_4 cluster. Further, the interactions between the Au clusters and CeO_2 surface are taken into account *via* calculating the adsorption energies without the CeO_2 surface, that is, by removing the CeO_2 surface from $\text{Au}_n/$

Table 2 The energetic span (δE) and TOF values for ω -alkynylfuran cycloisomerisation catalyzed by free gold clusters and supported gold clusters

	TDI/TDTS	δE (kcal mol^{-1})	TOF (s^{-1})
Au_3	IM4/IM5	27.45	3.3×10^{-6}
$\text{Au}_3/\text{CeO}_2(111)\text{-s}$	IM4/TS3/4	10.92	2.3×10^5
$\text{Au}_3/\text{CeO}_2(111)\text{-v}$	IM3/TS3/4	11.02	3.0×10^5
Au_4	IM2/IM5	16.94	13×10^1
$\text{Au}_4/\text{CeO}_2(111)\text{-s}$	IM3/IM5	11.26	7.4×10^4
$\text{Au}_4/\text{CeO}_2(111)\text{-v}$	IM3/TS3/4	9.27	2.2×10^6



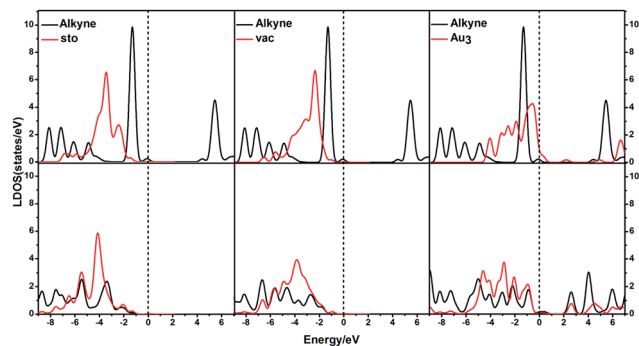


Fig. 16 The local density of states (LDOS) projected on triplet bonds on $\text{Au}_3/\text{CeO}_2(111)$ for the adsorption of the substrate at different sites, together with the d-projected LDOS of $\text{Au}_3/\text{CeO}_2(111)$ (stoichiometric and reduced) of the Au atom at different sites. Free C–C triplet bond and $\text{CeO}_2(111)$ (top), and adsorption (bottom). The Fermi level is set to zero.

CeO_2 (the resulting free Au_3 and Au_4 are represented as $\text{Au}_{3\text{-re}}$ and $\text{Au}_{4\text{-re}}$). The computed results are collected in Table S1† along with the initial values. Compared with the initial adsorption energies, the new ones possess larger values, implying that the strong interaction between the $\text{CeO}_2(111)$ surface and the substrate decreases the adsorption energies. To interpret the inherent reason for this, the d-projected LDOS were plotted in Fig. S25.† Based on Fig. S25,† near the Fermi level, the intensities of the d-bands of the $\text{Au}_{3\text{-re}}$ and $\text{Au}_{4\text{-re}}$ clusters are larger than those of $\text{Au}_3/\text{CeO}_2(111)$ and $\text{Au}_4/\text{CeO}_2(111)$, indicating that $\text{Au}_{3\text{-re}}$ and $\text{Au}_{4\text{-re}}$ clusters allow more opportunity for π -back-donation and σ -coordinate bonding with the substrate. In addition, compared with the $\text{Au}_{3\text{-re}}$ and $\text{Au}_{4\text{-re}}$ clusters, the d-bands of $\text{Au}_3/\text{CeO}_2(111)$ and $\text{Au}_4/\text{CeO}_2(111)$ are downshifted, which further illustrates that there is less opportunity to form σ -coordinate bonds with the substrate. Additionally, the Mulliken charges of the Au atoms for the free Au_3/Au_4 clusters, $\text{Au}_3/\text{Au}_4/\text{CeO}_2$ and $\text{Au}_3/\text{Au}_4/\text{CeO}_2$ were also calculated and the corresponding values are plotted in Fig. S26.† As shown in Fig. S26,† the Mulliken charges of Au (top) are 0.04, −0.02, −0.09, 0.10, 0.03 and −0.01, respectively, and the values of total charge transfer between the Au clusters and the support are 0.10, 0.43, 0.27, and 0.28. More positive charge indicates more unoccupied or partial occupied orbitals for the corresponding Au atom, which is beneficial for forming σ -coordinate bonds. Based on the computed results, the CeO_2 support can not only change the orbital levels of the gold clusters, but also cause charge recombination and decrease the positive charge on the Au (top) atom. These significant roles lead to the elevation of the adsorption energy, facilitating ω -alkynylfuran cycloisomerisation *via* the 5-*exo* FCT mechanism.

Conclusions

In summary, the catalytic properties of $\text{Au}_{10-11}/\text{CeO}_2$, $\text{Au}_{3-4}/\text{CeO}_2$, and free Au_3 and Au_4 clusters toward ω -alkynylfuran cycloisomerisation *via* the 5-*exo* FCT mechanism were systematically investigated based on density functional theory. The calculated results show that the atop and interface sites of $\text{Au}_{10-11}/\text{CeO}_2$ can

effectively facilitate ω -alkynylfuran cycloisomerisation *via* the FCT mechanism, indicating that the atop and interface sites possess excellent catalytic activity. Meanwhile, a surface oxygen vacancy on $\text{Au}_{10-11}/\text{CeO}_2$ can have a minor effect on the adsorption energies. In addition, for the sake of illustrating the role of the CeO_2 support, free Au_3 and Au_4 clusters were employed. By comparing the present results, supporting the Au_3 and Au_4 clusters on CeO_2 can decrease the energy barriers for the rate-determining step. For ω -alkynylfuran cycloisomerisation catalyzed *via* $\text{Au}_{3-4}/\text{CeO}_2$, the oxygen vacancy plays an active role. Overall, this study shows the catalytic activities of different reaction species on Au_n/CeO_2 and the role of the CeO_2 support, which is beneficial for providing some valuable information for the design of highly active metal/oxide catalysts for ω -alkynylfuran cycloisomerisation.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (Grant No. 21573030), the Chongqing Science & Technology Commission, China (Grant No. CSTC2013JCYJA50028) and the Scientific Research Foundation of Chongqing University of Arts and Sciences (R2013CJ03). The calculations were performed at the National Supercomputing Centre in Shenzhen (Shenzhen Cloud Computing Centre).

Notes and references

- 1 F. Boccuzzi, A. Chiorino, M. Manzoli, D. Andreeva and T. Tabakova, *J. Catal.*, 1999, **188**, 176–185.
- 2 W. D. Williams, M. Shekhar, W.-S. Lee, V. Kispersky, W. N. Delgass, F. H. Ribeiro, S. M. Kim, E. A. Stach, J. T. Miller and L. F. Allard, *J. Am. Chem. Soc.*, 2010, **132**, 14018–14020.
- 3 M. Haruta, T. Kobayashi, H. Sano and N. Yamada, *Chem. Lett.*, 1987, **2**, 405–408.
- 4 M. Valden, X. Lai and D. W. Goodman, *Science*, 1998, **281**, 1647–1650.
- 5 H. Y. Kim and G. Henkelman, *J. Phys. Chem. Lett.*, 2012, **4**, 216–221.
- 6 C. Sun, H. Li and L. Chen, *Energy Environ. Sci.*, 2012, **5**, 8475–8505.
- 7 C. Zhang, A. Michaelides and S. J. Jenkins, *Phys. Chem. Chem. Phys.*, 2011, **13**, 22–33.
- 8 F. Esch, S. Fabris, L. Zhou, T. Montini, C. Africh, P. Fornasiero, G. Comelli and R. Rosei, *Science*, 2005, **309**, 752–755.
- 9 N. J. Lawrence, J. R. Brewer, L. Wang, T.-S. Wu, J. Wells-Kingsbury, M. M. Ihrig, G. Wang, Y.-L. Soo, W.-N. Mei and C. L. Cheung, *Nano Lett.*, 2011, **11**, 2666–2671.
- 10 M. Baron, O. Bondarchuk, D. Stacchiola, S. Shaikhutdinov and H.-J. Freund, *J. Phys. Chem. C*, 2009, **113**, 6042–6049.
- 11 H. Y. Kim, H. M. Lee and G. Henkelman, *J. Am. Chem. Soc.*, 2012, **134**, 1560–1570.
- 12 W. Song and E. J. M. Hensen, *Catal. Sci. Technol.*, 2013, **3**, 8.
- 13 L. C. Wang, D. Widmann and R. J. Behm, *Catal. Sci. Technol.*, 2014, **5**, 925–941.



- 14 A. Ciftci, D. M. Lighthart, P. Pastorino and E. J. Hensen, *Appl. Catal., B*, 2013, **130**, 325–335.
- 15 Q. Fu, H. Saltsburg and M. Flytzani-Stephanopoulos, *Science*, 2003, **301**, 935–938.
- 16 M. Han, X. Wang, Y. Shen, C. Tang, G. Li and R. L. Smith Jr, *J. Phys. Chem. C*, 2009, **114**, 793–798.
- 17 M. F. Camellone and S. Fabris, *J. Am. Chem. Soc.*, 2009, **131**, 10473–10483.
- 18 H. Falsig, B. Hvolbæk, I. S. Kristensen, T. Jiang, T. Bligaard, C. H. Christensen and J. K. Nørskov, *Angew. Chem.*, 2008, **120**, 4913–4917.
- 19 W. Song and E. J. Hensen, *Catal. Sci. Technol.*, 2013, **3**, 3020–3029.
- 20 H. Y. Kim and G. Henkelman, *J. Phys. Chem. Lett.*, 2012, **3**, 2194–2199.
- 21 M. Cargnello, V. V. Doan-Nguyen, T. R. Gordon, R. E. Diaz, E. A. Stach, R. J. Gorte, P. Fornasiero and C. B. Murray, *Science*, 2013, **341**, 771–773.
- 22 A. Bruix, J. A. Rodriguez, P. J. Ramirez, S. D. Senanayake, J. Evans, J. B. Park, D. Stacchiola, P. Liu, J. Hrbek and F. Illas, *J. Am. Chem. Soc.*, 2012, **134**, 8968–8974.
- 23 Z. Zhou, S. Kooi, M. Flytzani-Stephanopoulos and H. Saltsburg, *Adv. Funct. Mater.*, 2008, **18**, 2801–2807.
- 24 B. Liu, J. Liu, T. Li, Z. Zhao, X.-Q. Gong, Y. Chen, A. Duan, G. Jiang and Y. Wei, *J. Phys. Chem. C*, 2015, **119**, 12923–12934.
- 25 B. Liu, Z. Zhao, G. Henkelman and W. Song, *J. Phys. Chem. C*, 2016, **120**, 5557–5564.
- 26 N. Lopez, T. Janssens, B. Clausen, Y. Xu, M. Mavrikakis, T. Bligaard and J. K. Nørskov, *J. Catal.*, 2004, **223**, 232–235.
- 27 K. Zhao, J. Qi, H. Yin, Z. Wang, S. Zhao, X. Ma, J. Wan, L. Chang, Y. Gao and R. Yu, *J. Mater. Chem. A*, 2015, **3**, 20465–20470.
- 28 J. Qi, J. Chen, G. Li, S. Li, Y. Gao and Z. Tang, *Energy Environ. Sci.*, 2012, **5**, 8937–8941.
- 29 J. Qi, K. Zhao, G. Li, Y. Gao, H. Zhao, R. Yu and Z. Tang, *Nanoscale*, 2014, **6**, 4072–4077.
- 30 H. Yin, H. Tang, D. Wang, Y. Gao and Z. Tang, *ACS Nano*, 2012, **6**, 8288–8297.
- 31 A. S. K. Hashmi, M. C. Blanco, E. Kurpejović, W. Frey and J. W. Bats, *Adv. Synth. Catal.*, 2006, **348**, 709–713.
- 32 J. H. Tyman, *Synthetic and natural phenols*, Elsevier, 1996.
- 33 A. S. K. Hashmi, T. M. Frost and J. Bats, *J. Am. Chem. Soc.*, 2000, **122**, 11553–11554.
- 34 A. S. K. Hashmi, T. M. Frost and J. W. Bats, *Org. Lett.*, 2001, **3**, 3769–3771.
- 35 S. Carrettin, M. C. Blanco, A. Corma and A. S. K. Hashmi, *Adv. Synth. Catal.*, 2006, **348**, 1283–1288.
- 36 A. S. K. Hashmi, J. Hofmann, S. Shi, A. Schütz, M. Rudolph, C. Lothschütz, M. Wietek, M. Bührle, M. Wölflle and F. Rominger, *Chem.–Eur. J.*, 2013, **19**, 382–389.
- 37 M. Yang, Z. Chen, Y. Luo, J. Zhang, D. Tang, R. He, W. Shen and M. Li, *RSC Adv.*, 2016, **6**, 22709–22721.
- 38 M. Yang, Z. Chen, Y. Luo, J. Zhang, R. He, W. Shen, D. Tang and M. Li, *ChemCatChem*, 2016, **14**, 2367–2375.
- 39 D. Tang, M. Yang, J. Zhang, R. He, W. Shen and M. Li, *ChemCatChem*, 2016, **8**, 461–470.
- 40 J. Oliver-Meseguer, A. Leyva-Pérez and A. Corma, *ChemCatChem*, 2013, **5**, 3509–3515.
- 41 J. Olivermeseguer, J. R. Cabreroantonino, I. Domínguez, A. Leyvapérez and A. Corma, *Science*, 2012, **338**, 1452–1455.
- 42 J. Oliver-Meseguer, A. Leyva-Pérez, S. I. Al-Resayes and A. Corma, *Chem. Commun.*, 2013, **49**, 7782–7784.
- 43 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- 44 B. Delley, *Int. J. Quantum Chem.*, 1998, **69**, 423–433.
- 45 S. Bell and J. S. Crighton, *J. Chem. Phys.*, 1984, **80**, 2464–2475.
- 46 S. Fischer and M. Karplus, *Chem. Phys. Lett.*, 1992, **194**, 252–261.
- 47 A. Klamt and G. Schüürmann, *J. Chem. Soc., Perkin Trans. 2*, 1993, 799–805.
- 48 J. Tomasi and M. Persico, *Chem. Rev.*, 1994, **94**, 2027–2094.
- 49 B. Delley, *Mol. Simul.*, 2006, **32**, 117–123.
- 50 B. Delley, *J. Chem. Phys.*, 1990, **92**, 508–517.
- 51 B. Delley, *J. Chem. Phys.*, 2000, **113**, 7756–7764.
- 52 A. Trovarelli, *Catalysis by ceria and related materials*, World Scientific, 2002, pp. 1–501.
- 53 H. Y. Kim and G. Henkelman, *J. Phys. Chem. Lett.*, 2012, **3**, 2194–2199.
- 54 H. Y. Kim, H. M. Lee and G. Henkelman, *J. Am. Chem. Soc.*, 2012, **134**, 1560–1570.
- 55 G. Vilé, P. Dähler, J. Vecchietti, M. Baltanás, S. Collins, M. Calatayud, A. Bonivardi and J. Pérez-Ramírez, *J. Catal.*, 2015, **324**, 69–78.
- 56 A. D. Mayernick and M. J. Janik, *J. Phys. Chem. C*, 2008, **112**, 14955–14964.
- 57 L. C. Hsu, M. K. Tsai, Y. H. Lu and H. T. Chen, *J. Phys. Chem. C*, 2013, **117**, 367–371.
- 58 Y. Zhao, C. Cui, J. Han, H. Wang, X. Zhu and Q. Ge, *J. Am. Chem. Soc.*, 2016, **138**, 10191–10198.
- 59 J. L. F. D. Silva, V. M. O. N. Ganduglia-Pirovano, J. Sauer, V. Bayer and G. Kresse, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2007, **75**, 33–39.
- 60 M. Fronzi, S. Piccinin, B. Delley, E. Traversa and C. Stampfl, *Phys. Chem. Chem. Phys.*, 2009, **11**, 9188–9199.
- 61 M. Fronzi, S. Piccinin, B. Delley, E. Traversa and C. Stampfl, *RSC Adv.*, 2013, **4**, 12245–12251.
- 62 M. Fronzi, A. Soon, B. Delley, E. Traversa and C. Stampfl, *J. Chem. Phys.*, 2009, **131**, 104701–104716.
- 63 S. Kumar and P. K. Schelling, *J. Chem. Phys.*, 2006, **125**, 261–288.
- 64 Z. Yang, T. K. Woo, M. Baudin and K. Hermansson, *J. Chem. Phys.*, 2004, **120**, 7741–7749.
- 65 S. Kozuch and S. Shaik, *Acc. Chem. Res.*, 2010, **44**, 101–110.
- 66 S. Kozuch and J. M. Martin, *ACS Catal.*, 2011, **1**, 246–253.
- 67 A. Uhe, S. Kozuch and S. Shaik, *J. Comput. Chem.*, 2011, **32**, 978–985.
- 68 S. Kozuch, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 795–815.
- 69 C. Zhang, A. Michaelides, D. A. King and S. J. Jenkins, *J. Am. Chem. Soc.*, 2010, **132**, 2175–2182.
- 70 A. S. K. Hashmi, M. Rudolph, J. P. Weyrauch, M. Wölflle, W. Frey and J. W. Bats, *Angew. Chem., Int. Ed.*, 2005, **44**, 2798–2801.
- 71 B. Hammer and J. K. Nørskov, *Adv. Catal.*, 2000, **45**, 71–129.

