

Quantum chemistry of the Fischer–Tropsch reaction
catalysed by a stepped ruthenium surface†

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A comprehensive density functional theory study of the Fischer–Tropsch mechanism on the corrugated Ru(11 $\bar{2}$ 1) surface has been carried out. Elementary reaction steps relevant to the carbide mechanism and the CO insertion mechanism are considered. Activation barriers and reaction energies were determined for CO dissociation, C hydrogenation, CH_x + CH_y and CH_x + CO coupling, CH_xCH_y–O bond scission and hydrogenation reactions, which lead to formation of methane and higher hydrocarbons. Water formation that removes O from the surface was studied as well. The overall barrier for chain growth in the carbide mechanism (preferred path CH + CH coupling) is lower than that for chain growth in the CO insertion mechanism (preferred path C + CO coupling). Kinetic analysis predicts that the chain-growth probability for the carbide mechanism is close to unity, whereas within the CO insertion mechanism methane will be the main hydrocarbon product. The main chain propagating surface intermediate is CH via CH + CH and CH + CR coupling (R = alkyl). A more detailed electronic analysis shows that CH + CH coupling is more difficult than coupling reactions of the type CH + CR because of the σ -donating effect of the alkyl substituent. These chain growth reaction steps are more facile on step-edge sites than on terrace sites. The carbide mechanism explains the formation of long hydrocarbon chains for stepped Ru surfaces in the Fischer–Tropsch reaction.

Introduction

The limited supply of readily available petroleum stimulates the search for alternative energy sources.¹ Due to its abundance, natural gas is increasingly considered as a valuable alternative feedstock for the synthesis of fuels and chemicals. Synthesis gas (a mixture of CO and H₂), which can be obtained from natural gas by autothermal or steam reforming, can be used to produce long-chain hydrocarbons in the Fischer–Tropsch (FT) reaction.^{2–4} Several mechanistic proposals for this chain-growth reaction have been reviewed elsewhere.⁵ They can be distinguished by the assumption of the inserting species in the growing chain. The inserting species is either a CH_x monomer derived from surface CO dissociation in the carbide mechanism or an intermediate with the C–O bond intact in the CO insertion mechanism. The carbide mechanism involves dissociation of adsorbed CO, hydrogenation of the carbon adatom to a CH_x building block and its insertion into the growing hydrocarbon chains on the surface. The chain reaction is

terminated by desorption of the hydrocarbon chain from the surface as an alkene or alkane. An alternative termination pathway involves CO insertion, that results in aldehyde and alcohol products. In the CO insertion mechanism, chain propagation proceeds via insertion of a CO moiety and the C–O bond scission therefore occurs after C–C coupling. This is in contrast to the carbide mechanism, where C–O bond scission occurs prior to C–C coupling. Oxygen is predominantly removed from the surface as water.^{5–8} The competing reaction is complete hydrogenation of the CH_x surface intermediate to methane, which is an undesired by-product.

Good FT catalysts should exhibit high selectivity towards long-chain hydrocarbons^{4,9} which requires facile CO dissociation and slow chain-growth termination.^{10,11} Formation of methane, the undesired by-product of the FT reaction, should be minimized. The self-organization of monomeric C₁ species into growing chains can be seen as a simple polymerization process and the molecular weights of the hydrocarbon products tend to follow the Anderson–Schulz–Flory (ASF) distribution.^{12,13} Experimentally, it is usually found that the C₁ and C₂ product selectivities deviate from the ASF distribution.^{12,14–20}

Two important schools of thought exist about the nature of the catalytically active surface. On the one hand, it is assumed that the close-packed surfaces (terraces) are the active sites.^{21,22} Although metal atoms with low coordination numbers are in principle more reactive, it has been argued that

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these more reactive low-coordinated sites become blocked due to their strong interaction with the adsorbed species and, in particular, with carbon adatoms. On the other hand, the high reactivity of low-coordinated sites, especially for CO bond dissociation over stepped (B_5) sites, is considered to be crucial to provide sufficient monomer building blocks to maintain a high rate of chain growth.^{11,23–26}

The mechanism of CO dissociation on metallic surfaces has recently been elucidated. Experimental observations^{27,28} and theoretical calculations^{24,29,30} agree on the importance of ensembles of surface metal atoms arranged in such a way that a stepped site is obtained for facile CO dissociation. These sites are usually called B_5 sites. On such stepped sites of Co and Ru surfaces, the direct dissociation of CO is favoured over the hydrogen-assisted alternative involving the formyl (CHO) intermediate.^{23,24,31,32} Although the discovery of the Fischer–Tropsch reaction is almost a century ago, many open questions remain about the reactions that lead to chain growth and chain-growth termination.

A large number of candidate reaction steps have been proposed for chain growth relevant to carbide and CO insertion mechanisms. Originally, Fischer and Tropsch proposed that surface CH_2 couples with surface CH_3 .^{2,33} Modern insights about surface reactivity and theoretical chemistry advances now learn us that this proposal is incorrect, because the predicted barrier for this coupling step is very high³⁰ and, also, because the surface coverage of CH_2 and CH_3 is predicted to be low.^{34–36} Hu and coworkers³⁷ have explored various coupling reactions for a range of transition metal surfaces using Density Functional Theory (DFT). Their analysis stresses the importance of the surface abundance of particular CH_x species and, also, indicates that stepped sites are preferred for C–C bond formation reactions for Ru surfaces, with C + CH and CH + CH coupling being the most favourable pathways. On Co surfaces, however, CH_2 + CH_2 coupling and C + CH_3 coupling are more likely candidates for chain growth. Saeys and co-workers have studied the CO insertion pathway on flat Co(0001) surfaces.^{38,39} CH_2 + CO coupling was found to be the most favourable pathway.³⁸ The group of Maitlis has performed extensive studies on the mechanistic aspects employing a model system using a Co homogeneous catalyst.^{40–42} They found that the CH_2 insertion into a surface alkyl is the dominant mechanism. Water formation, which is required to remove O originating from CO dissociation, was also studied recently by DFT.^{43–46} It is generally proposed that, following OH formation, formation of adsorbed water occurs more favourably through proton migration between two hydroxyl adsorbates than *via* direct hydrogenation of the hydroxyl intermediate.

In the present theoretical study, we employed DFT to investigate all elementary reaction steps from syngas following the carbide as well as the CO insertion mechanism that lead to formation of ethylene and ethane on a stepped Ru(11 $\bar{2}$ 1) surface. These reactions include the already well-studied CO dissociation and CH_4 formation, because our aim is to build a database of kinetic parameters for all reaction steps relevant to the FT reaction at the same computational

accuracy. We also study the hydrogenation of the surface intermediates towards olefinic and paraffin products and include formation of water, which removes O atoms from the surface. We rationalize the experimentally observed lower C_2 selectivity in the ASF distribution by considering how the reactivity of C_3 surface intermediates will differ from that of C_2 intermediates. We explain the different kinetics of coupling reactions occurring on terraces and step-edge sites. Finally, we elaborate on the most likely FT pathway by comparing the different FT reaction routes comprising both the carbide and the CO insertion mechanism.

Method

Density Functional Theory (DFT) calculations were performed using the Vienna *Ab initio* simulation package (VASP).^{47,48} The Perdew–Burke–Ernzerhof (PBE) exchange–correlation functional was employed for all calculations.⁴⁹ To describe the interaction between nuclei and core electrons, the projector-augmented wave (PAW) method was used.^{50,51} For the valence electrons a plane-wave basis set with an energy cut-off of 400 eV and a Brillouin zone sampling of $5 \times 5 \times 1$ k-points were used. All elementary reactions were investigated on a stepped Ru(11 $\bar{2}$ 1) surface model (Fig. 1). The model for the stepped site consists of a slab with a thickness of at least 5 atomic layers and using a $p(2 \times 2)$ unit cell. To confirm that the thickness of the slab was sufficient, it was verified that the energy with respect to the number of layers in the slab was converged. The Ru(11 $\bar{2}$ 1) surface contains terrace and step-edge sites and is accordingly representative for a dual reaction centre mechanism as explored earlier.¹⁰ In order to avoid spurious interactions between the images of the system, a vacuum layer of at least 10 Å was added along the z-axis. To confirm that the vacuum layer was large enough, it was verified that the electron density approached zero at the border of the unit cell. To avoid dipole–dipole interactions between the super cells, adsorbates were placed on both sides of the surface retaining an inversion centre.

Ionic relaxation was carried out by the conjugate gradient method. During geometry optimization, all the degrees of freedom of the atoms in the slab as well as the adsorbed species were relaxed. To determine transition states, the nudged



Fig. 1 Schematic representation of (left) the corrugated Ru(11 $\bar{2}$ 1) surface and (right) the close-packed Ru(0001) terrace surface.



elastic band (NEB) method as developed by Jónsson *et al.* was employed.⁵² The initial images between the optimized initial and final states were obtained *via* linear interpolation. The transition state was confirmed to be a saddle point on the electronic energy surface by means of frequency analysis. Within VASP, this frequency analysis is performed by constructing a Hessian matrix using the finite difference approach. We only considered the frequencies of the adsorbed species within this analysis as the contribution of the metal atoms can be neglected. Prior to investigating the location and properties of the transition states, the structure and energetics of a large number of adsorption models of reactants, intermediates and products were determined. Based on these results, the initial guesses for the transition states were constructed.

Results and discussion

DFT calculations

DFT calculations will be presented for the elementary reaction steps relevant to the carbide as well as the CO insertion mechanism of the Fischer-Tropsch reaction. The main groups of reactions studied are (i) CO dissociation, (ii) C hydrogenation to CH_x intermediate and CH₄, (iii) CH_x-CH_x coupling, (iv) CH_x-CH_x hydrogenation, (v) CH_x-CO coupling, (vi) CH_xCH_yO hydrogenation, (vii) CH_xCH_y-O scission and (viii) O hydrogenation to H₂O. For the carbide mechanism, reactions of groups (i), (ii), (iii), (iv) and (viii) are relevant. The CO insertion mechanism is covered by reaction of groups (i), (v), (vi), (vii) and (viii). We did not explore oxygen removal *via* CO₂ formation, because the CO₂ selectivity for Ru catalysts in the FT reaction is low.^{13,53,54}

CO dissociation

The energy barriers for CO dissociation are given in Table 1 and correspond well with values reported before for the stepped Ru(11 $\bar{2}$ 1) surface.²³ Direct dissociation of CO is the preferred route, because the hydrogen-assisted route involves the thermodynamically unfavourable formyl intermediate, resulting in a much higher overall barrier for CH formation.

C hydrogenation to CH₄

Hydrogenation of adsorbed C to CH₄ has been investigated for the Ru(11 $\bar{2}$ 1) surface before.³⁶ Here, we carried out these calculations at the same computational accuracy as for the

other elementary reaction steps. The forward and backward activation energies for the consecutive hydrogenation steps of CH_x ($x = 0-3$) to CH_y ($y = 1-4$) are listed in Table 1.

The corresponding initial, transition and final states for these reactions are given in the ESI† (Table S1). The activation energies are with respect to the most stable adsorption site of the reacting surface adsorbates. In analogy with results for other transition metals, C hydrogenation to CH is relatively facile. For Ru(11 $\bar{2}$ 1), it is slightly endothermic. Further hydrogenation to CH₂ (methylidene) is more difficult ($E_{\text{act}} = 75 \text{ kJ mol}^{-1}$) and endothermic by 38 kJ mol^{-1} . This implies that at reasonable reaction temperatures surface CH₂ and its hydrogenation products are much less abundant than C and CH surface intermediates. The barrier for the endothermic CH₃ (methyl) formation is 57 kJ mol^{-1} . The barrier for CH₄ formation, which involves a single elementary reaction step recombining CH₃ and H over a single Ru atom, has the highest barrier among the CH_x hydrogenation steps (94 kJ mol^{-1}).

CH_x + CH_y coupling reactions

A total of 10 reactions between CH_x species were considered for carbon-carbon bond formation to describe chain growth in the carbide mechanism of the Fischer-Tropsch reaction (Table 2).

The corresponding initial, transition and final states are given in the ESI†. Reactions between CH₂ and CH₃ and between two CH₃ (reactions 14 and 15) were not investigated, because it is well known that the interaction of the spatially extended C-H bonds in CH₃ adsorbates results in repulsion, precluding C-C bond formation.⁵⁵ Despite considerable efforts, we could not identify transition states for the following reactions: C + CH₂, CH + CH₂ and CH + CH₃ coupling. We assume that these reaction will not occur on the Ru(11 $\bar{2}$ 1) surface. It should be noted that these reactions have been reported to be feasible on terrace surfaces.⁵⁶ For the remaining five coupling steps C + C, C + CH, CH + CH, CH₂ + CH₂ and C + CH₃ coupling, we identified six unique transition states. For the CH + CH coupling step, two different pathways were found (see Fig. 2).

The analogous coupling reaction involving longer chains adsorbed on the surface are of the CH + CR type with R being an alkyl group. Therefore, we investigated this latter reaction step for R = CH₃ in more detail in section 3.2. Table 2 shows that the barrier for CH + CH coupling is substantially lower on the step-edge site than on the terrace surface. This

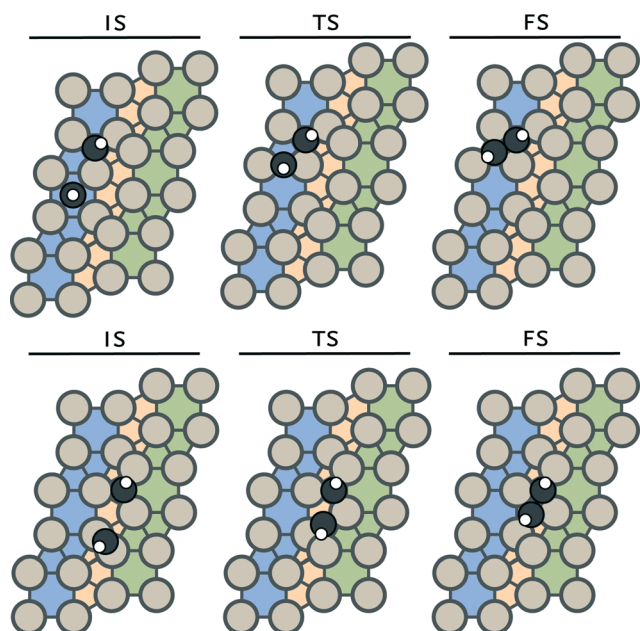
Table 1 Methanation pathway of synthesis gas on Ru(11 $\bar{2}$ 1). The reported forward and reverse energies are in relation to the most stable states found for the reactants and products

| Index | Elementary reaction | Forward E_{act} (kJ mol ⁻¹) | Backward E_{act} (kJ mol ⁻¹) |
|-------|--|--|---|
| 1 | CO* + * → C* + O* | 65 | 90 |
| 2 | C* + H* → CH* + * | 40 | 39 |
| 3 | CH* + H* → CH ₂ * + * | 75 | 37 |
| 4 | CH ₂ * + H* → CH ₃ * + * | 57 | 47 |
| 5 | CH ₃ * + H* → CH ₄ + 2* | 94 | 57 |



Table 2 $\text{CH}_x + \text{CH}_y$ coupling reactions and their forward and backward activation energies for Ru(11 $\bar{2}$ 1)

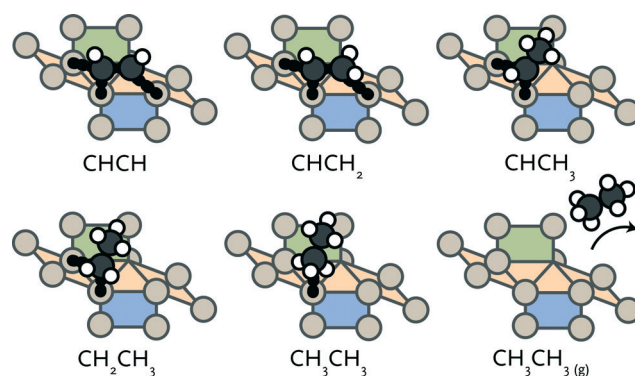
| Index | Elementary reaction | Forward E_{act} (kJ mol $^{-1}$) | Backward E_{act} (kJ mol $^{-1}$) |
|-------|--|--|---|
| 6 | $\text{C}^* + \text{C}^* \rightarrow \text{CC}^* + *$ | 138 | 144 |
| 7 | $\text{C}^* + \text{CH}^* \rightarrow \text{CCH}^* + *$ | 129 | 75 |
| 8 | $\text{C}^* + \text{CH}_2^* \rightarrow \text{CCH}_2^* + *$ | Not found | |
| 9 | $\text{C}^* + \text{CH}_3^* \rightarrow \text{CCH}_3^* + *$ | 92 | 116 |
| 10a | $\text{CH}^* + \text{CH}^* \rightarrow \text{CHCH}^* + *$ (terrace) | 149 | 117 |
| 10b | $\text{CH}^* + \text{CH}^* \rightarrow \text{CHCH}^* + *$ (step) | 86 | 55 |
| 11 | $\text{CH}^* + \text{CH}_2^* \rightarrow \text{CHCH}_2^* + *$ | Not found | |
| 12 | $\text{CH}^* + \text{CH}_3^* \rightarrow \text{CHCH}_3^* + *$ | Not found | |
| 13 | $\text{CH}_2^* + \text{CH}_2^* \rightarrow \text{CH}_2\text{CH}_2^* + *$ | 54 | 60 |
| 14 | $\text{CH}_2^* + \text{CH}_3^* \rightarrow \text{CH}_2\text{CH}_3^* + *$ | Not investigated | |
| 15 | $\text{CH}_3^* + \text{CH}_3^* \rightarrow \text{CH}_3\text{CH}_3^* + *$ | Not investigated | |

**Fig. 2** Schematic representation of the two CH + CH coupling pathways on the Ru(11 $\bar{2}$ 1) surface. The coupling of two CH moieties on (top) a terrace site and (bottom) a step-edge site. The black and white spheres represent carbon and hydrogen atoms, respectively.

preference for coupling on the step-edge site is in accordance with several other works.^{37,57,58} Based on this insight, we investigated the other coupling reactions (reactions 6, 7, 9 and 13) on the stepped surface as well. The most facile coupling reaction occurs between two methyldene species with a barrier of 54 kJ mol $^{-1}$ (reaction 6). The barrier for the other coupling reactions are higher: 86 kJ mol $^{-1}$ for CH + CH, 92 kJ mol $^{-1}$ for C + CH₃, 129 kJ mol $^{-1}$ for C + CH and 138 kJ mol $^{-1}$ for coupling of two carbon adatoms (C + C).

CH_xCH_y hydrogenation reactions

After formation of carbon–carbon bonds, higher hydrocarbons may leave the surface as olefins or paraffins (see Fig. 3). We investigated such chain-growth termination steps for $\text{CH}_x\text{--CH}_y$ species. From CCH, a large number of hydrogenation routes to ethylene and ethane needs to be considered. These possibilities

**Fig. 3** Schematic representation of hydrogenation of adsorbed acetylene to ethane.

and the results of a reaction energy analysis are given in Table 3.

The structures of the involved stable and transition states are collected in the ESI.† Hydrogenation of CCH to CCH₂ (reaction 17) and CHCH to CHCH₂ (reaction 19) have very similar forward activation barriers around 82 kJ mol $^{-1}$. While CCH₂ formation is exothermic, CHCH₂ formation is endothermic. Subsequent hydrogenation to CCH₃ (reaction 18) and CHCH₃ (reaction 20) proceeds with barriers of 19 and 62 kJ mol $^{-1}$, respectively.

Comparatively, hydrogenation of CCH to CHCH and the reverse dehydrogenation of CHCH to CCH (reaction 21) are kinetically hindered with barriers of 140 and 162 kJ mol $^{-1}$, respectively. Hydrogenation of CCH₂ to CHCH₂ (reaction 22) and of CCH₃ to CHCH₃ (reaction 23) have barriers of 82 kJ mol $^{-1}$. The reverse dehydrogenation reactions are very facile with barriers of 21 and 8 kJ mol $^{-1}$, respectively. This implies that CCH_x intermediates are significantly more stable than the corresponding CHCH_x intermediates. This difference draws similarity to the higher stability of adsorbed CH over CH₂. Finally, we consider reactions that lead to formation of ethylene (reaction 24) and ethane (reactions 25, 26 and 27). Whereas ethylene formation proceeds with an activation energy of 45 kJ mol $^{-1}$ and is almost thermodynamically neutral, the hydrogenation of CHCH₃ to CH₂CH₃ and, finally, CH₃CH₃ is endothermic. Although the first hydrogenation step to CH₂CH₃ is facile and slightly exothermic, the second



Table 3 Hydrogenation reactions of C₂ surface intermediates and their forward and backward activation energies for Ru(11 $\bar{2}$ 1)

| Index | Elementary reaction | Forward E_{act} (kJ mol ⁻¹) | Backward E_{act} (kJ mol ⁻¹) |
|-------|--|--|---|
| 16 | $\text{CC}^* + \text{H}^* \rightarrow \text{CCH}^* + *$ | 104 | 72 |
| 17 | $\text{CCH}^* + \text{H}^* \rightarrow \text{CCH}_2^* + *$ | 82 | 129 |
| 18 | $\text{CCH}_2^* + \text{H}^* \rightarrow \text{CCH}_3^* + *$ | 19 | 4 |
| 19 | $\text{CHCH}^* + \text{H}^* \rightarrow \text{CHCH}_2^* + *$ | 83 | 46 |
| 20 | $\text{CHCH}_2^* + \text{H}^* \rightarrow \text{CHCH}_3^* + *$ | 62 | 34 |
| 21 | $\text{CCH}^* + \text{H}^* \rightarrow \text{CHCH}^* + *$ | 140 | 162 |
| 22 | $\text{CCH}_2^* + \text{H}^* \rightarrow \text{CHCH}_2^* + *$ | 82 | 21 |
| 23 | $\text{CCH}_3^* + \text{H}^* \rightarrow \text{CHCH}_3^* + *$ | 82 | 8 |
| 24 | $\text{CHCH}_2^* + \text{H}^* \rightarrow \text{CH}_2\text{CH}_2^* + *$ | 45 | 42 |
| 25 | $\text{CHCH}_3^* + \text{H}^* \rightarrow \text{CH}_2\text{CH}_3^* + *$ | 19 | 23 |
| 26 | $\text{CH}_2\text{CH}_2^* + \text{H}^* \rightarrow \text{CH}_2\text{CH}_3^* + *$ | 58 | 34 |
| 27 | $\text{CH}_2\text{CH}_3^* + \text{H}^* \rightarrow \text{CH}_3\text{CH}_3^* + *$ | 112 | 71 |

hydrogenation step toward ethane has to overcome a relatively high barrier of 112 kJ mol⁻¹. It is endothermic by 41 kJ mol⁻¹. Alternatively, adsorbed ethylene can be further hydrogenated to CH₂CH₃ (reaction 26). Although this reaction is facile with an activation barrier of 58 kJ mol⁻¹, the reverse reaction is more favourable with an activation energy of 34 kJ mol⁻¹.

CH_x + CO coupling

Three different reactions between CH_x species and CO were considered for chain propagation in the CO insertion mechanism (Table 4). These reactions all involve the migration of adsorbed CO to a site adjacent to the CH_x adsorption site followed by C–C bond formation. Both carbon atoms in the final coupled CH_x–CO species coordinate to surface Ru atoms. This strongly differs from mechanisms proposed based on homogeneous mononuclear coordination complexes for CO insertion or migration.⁵⁹

The corresponding initial, transition and final states are given in the ESI.† The reaction between CH₃ and CO (reaction 30) was not considered due to the expected steric repulsion. The most facile CO insertion reaction is the C + CO coupling reaction (reaction 28) with a barrier of 138 kJ mol⁻¹. The barrier for CH + CO coupling (reaction 29) and CH₂ + CO coupling (reaction 30) are somewhat higher at 148 kJ mol⁻¹ and 155 kJ mol⁻¹, respectively. From comparison with the data in Table 2, it is immediately clear that the barriers for CO insertion are significantly higher than those for CH_x + CH_y coupling (Table 2). Moreover, all CO insertion reactions are strongly endothermic by at least 77 kJ mol⁻¹, implying that the rate constants for the reverse reactions are substantially higher.

CH_x–CO hydrogenation

After CO insertion, the resulting CH_xCO moiety can be hydrogenated to form a CH_xCH_yO species. We did not consider

formation of oxygenated products that can leave the surface, because their yield is usually very low under practical FT conditions. For each CH_xCO* intermediate, the hydrogenation of either carbon atom was considered. This results in five different hydrogenation reactions (Table 5).

We could not identify a transition state for CCO* hydrogenation (reaction 33). The orientation of the CCO* moiety in the step-edge site (see the ESI†) prevents hydrogenation of the carbon atom of the CO group. The most difficult hydrogenation step is that of CCO* to CHCO* (reaction 32), which involves an activation energy of 104 kJ mol⁻¹. The other reactions, *i.e.* the hydrogenation of CHCO* to CH₂CO, that of CHCO* to CHCHO* and finally the hydrogenation of CH₂CO* to CH₂CHO* proceed with lower activation energies of 95, 68 and 63 kJ mol⁻¹, respectively. All CH_xCO hydrogenation reactions are endothermic.

CH_xCH_y–O bond scission

For chain growth to proceed, the C–O bond in the CH_xCH_yO surface intermediate must first cleave. Five such elementary reaction steps were considered for which results are listed in Table 6.

Among these reactions, it is most easy to cleave the C–O bond in CCO* with an activation energy of 52 kJ mol⁻¹. The C–O bond cleavage barriers for the other reactions are higher. All reactions are strongly exothermic, especially those for the partially hydrogenated fragments.

Water formation

In the FT reaction, the surface oxygen atoms deriving from CO dissociation are predominantly removed as water (Table 7). Water formation can proceed *via* two mechanisms, both involving intermediate OH formation from O and H (reaction 42) followed by either direct hydrogenation of OH

Table 4 CH_x + CO coupling reactions and their forward and backward activation energy

| Index | Elementary reaction | Forward E_{act} (kJ mol ⁻¹) | Backward E_{act} (kJ mol ⁻¹) |
|-------|--|--|---|
| 28 | $\text{C}^* + \text{CO}^* \rightarrow \text{CCO}^* + *$ | 138 | 61 |
| 29 | $\text{CH}^* + \text{CO}^* \rightarrow \text{CHCO}^* + *$ | 148 | 44 |
| 30 | $\text{CH}_2^* + \text{CO}^* \rightarrow \text{CH}_2\text{CO}^* + *$ | 155 | 36 |
| 31 | $\text{CH}_3^* + \text{CO}^* \rightarrow \text{CH}_3\text{CO}^* + *$ | <i>Not investigated</i> | |



Table 5 Hydrogenation reactions of CH_xCO surface intermediates and their forward and backward activation energies for Ru(11 $\bar{2}$ 1)

| Index | Elementary reaction | Forward E_{act} (kJ mol $^{-1}$) | Backward E_{act} (kJ mol $^{-1}$) |
|-------|---|--|---|
| 32 | $\text{CCO}^* + \text{H}^* \rightarrow \text{CHCO}^* + *$ | 104 | 77 |
| 33 | $\text{CCO}^* + \text{H}^* \rightarrow \text{CCHO}^* + *$ | Not found | |
| 34 | $\text{CHCO}^* + \text{H}^* \rightarrow \text{CH}_2\text{CO}^* + *$ | 95 | 41 |
| 35 | $\text{CHCO}^* + \text{H}^* \rightarrow \text{CHCHO}^* + *$ | 68 | 7 |
| 36 | $\text{CH}_2\text{CO}^* + \text{H}^* \rightarrow \text{CH}_2\text{CHO}^* + *$ | 63 | 4 |

Table 6 $\text{CH}_x\text{CH}_y\text{-O}$ bond scission and their forward and backward activation energies

| Index | Elementary reaction | Forward E_{act} (kJ mol $^{-1}$) | Backward E_{act} (kJ mol $^{-1}$) |
|-------|--|--|---|
| 37 | $\text{CCO}^* + * \rightarrow \text{CC}^* + \text{O}^*$ | 52 | 127 |
| 38 | $\text{CHCO}^* + * \rightarrow \text{CCH}^* + \text{O}^*$ | 92 | 163 |
| 39 | $\text{CH}_2\text{CO}^* + * \rightarrow \text{CCH}_2^* + \text{O}^*$ | 77 | 248 |
| 40 | $\text{CHCHO}^* + * \rightarrow \text{CHCH}^* + \text{O}^*$ | 72 | 225 |
| 41 | $\text{CH}_2\text{CHO}^* + * \rightarrow \text{CHCH}_2^* + \text{O}^*$ | 65 | 234 |

Table 7 Elementary reaction steps leading to removal of water including their forward and backward activation energies

| Index | Elementary reaction | Forward E_{act} (kJ mol $^{-1}$) | Backward E_{act} (kJ mol $^{-1}$) |
|-------|---|--|---|
| 42 | $\text{O}^* + \text{H}^* \rightarrow \text{OH}^* + *$ | 97 | 49 |
| 43 | $\text{OH}^* + \text{H}^* \rightarrow \text{H}_2\text{O}^* + *$ | 89 | 15 |
| 44 | $\text{OH}^* + \text{OH}^* \rightarrow \text{H}_2\text{O}^* + \text{O}^* + *$ | 54 | 11 |

to H_2O (reaction 43) or *via* proton migration between two OH species to form H_2O and O (reaction 44). The first hydrogenation step to produce adsorbed OH has a relatively high barrier of 97 kJ mol $^{-1}$. The barrier for water formation *via* OH hydrogenation is 89 kJ mol $^{-1}$. Comparatively, the reaction between two hydroxyl groups is much more facile with a barrier of 54 kJ mol $^{-1}$. This suggests that water formation by reaction of two OH groups to H_2O and adsorbed O is preferred.

Electron density difference analysis

To understand the differences in activation energies of the various elementary reaction steps, we investigated electron density differences of selected reactions. Since the use of atomic charges is not preferred as it is difficult to assign electrons to a specific atom and chemical bonding typically involves contributions of several molecular orbitals, it is more instructive to map electron density differences in space. We focus here on three important issues relevant for the FT reaction. Firstly, we address the different reactivities of terrace and stepped sites. In particular, we focus on the coupling step with the lowest activation barrier, *i.e.* CH + CH coupling. Secondly, we will compare CH coupling with CH and CCH_3 on stepped sites, because the latter reaction is relevant for formation of C_3 products. Finally, we will highlight the different adsorption strengths of ethylene and propylene on stepped sites, which explains the lower than expected C_2 selectivity observed during the FT reaction.

CH + CH coupling

The barriers for CH + CH coupling on stepped and terrace sites of the Ru(11 $\bar{2}$ 1) (Fig. 2) are 86 and 149 kJ mol $^{-1}$,

respectively. Fig. 4a shows the electron density shifts that result from the formation of C–C bonds between two CH surface species in the corresponding transition state complexes. To determine these, the electron density distributions of the two separate CH fragments and the empty surface were subtracted from the electron density distribution of the transition state complex. It is seen that in the transition state for the stepped site the electron density between the two C atoms is lowered and shifts to the metal atom. The corresponding shift in electron density for the transition state formation on the terrace site is smaller. There are two reasons for this difference. Firstly, it relates to the specific geometry of the transition state complex on the stepped site, which results in increased overlap between the bonding C–C orbitals and the metal d-band. Secondly, the coordinative unsaturation of the surface metal atoms is higher for the stepped than for the terrace surface. The decrease in electron density between the two CH fragments by electron donation to the metal center results in decreased Pauli repulsion and, accordingly, in lower activation energy for coupling.

CH + CH vs. CH + CCH_3 coupling

We found that methylidyne (CH) coupling to the CCH_3 surface intermediate (Fig. 4b) proceeds with lower barrier than CH coupling to another CH species (65 vs. 86 kJ mol $^{-1}$). In Fig. 4a, the electron density shifts for C–C bond formation are shown. It can be seen that in the transition state for CH + CCH_3 coupling more electron density is shifted from between the CH and CCH_3 species to the metal atom relative to the CH–CH case. This difference is caused by the σ -donating effect of the methyl



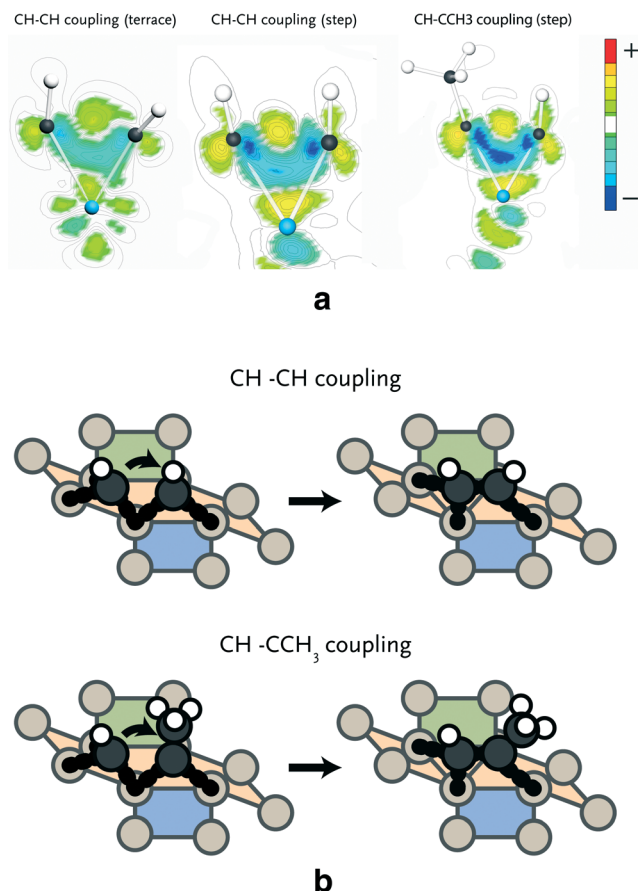


Fig. 4 a. Electron density difference plots of the transition state for CH + CH coupling on the terrace (left), step-edge (middle) and of CH + CCH₃ coupling on the step-edge (right) of the Ru(1121) surface. These plots were generated by subtracting the electron density of the individual CH moieties on the surface from the electron density of the complex. Note that the electron density difference for the step-edge site is larger than for the terrace site. 4b. Schematic representation of CH-CH and CH-CCH₃ coupling on a step-edge site of the Ru(1121) surface. The initial and final state of reaction are displayed.

group. Consequently, the Pauli repulsion between CH and CCH₃ is lower than between two CH fragments, explaining why the barrier for coupling of the former two fragments is lower.

Ethylene vs. propylene desorption

Ethylene and propylene adsorb 'side-on' on a single Ru atom through interaction with their C=C double bonds. The adsorption energies of ethylene and propylene are 120 and 100 kJ mol⁻¹, respectively. The electron density shifts upon olefin adsorption are shown in Fig. 5. Here, the electron density difference is determined by subtracting the electron density of the adsorbate in the geometry it has in the adsorbed state and the empty surface from the electron density of the surface-adsorbate complex. The difference in electron density for ethylene is larger than for propylene. The reason for the decreased electron density shift of the double bond in adsorbed propylene is the π -accepting nature of the CH₃ substituent. This results in weaker adsorption of propylene as

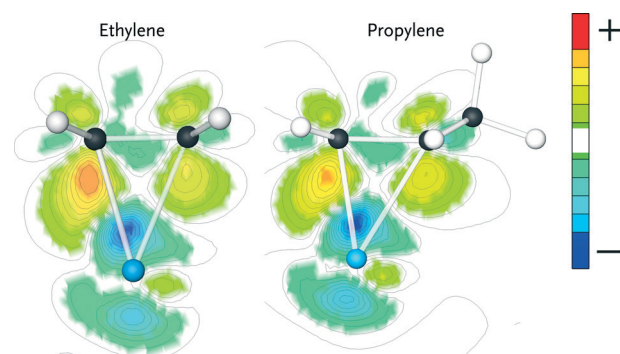


Fig. 5 Electron density difference plots of the adsorbed states of ethylene (left) and propylene (right) on the Ru(1121) surface. These plots were generated by subtracting the electron density of the individual complexes in vacuum and empty surface from the electron density of the complex. Note that the electron density difference for ethylene is larger than for propylene.

compared to ethylene. We expect that the stronger interaction of ethylene will result in increased surface residence time of this fragment, which increases the probability of its chain growth relative to desorption and hydrogenation as compared with that of its higher carbon number analogues.

Reaction energy diagrams

We constructed reaction energy diagrams based on the elementary reaction steps explored in the DFT study (Tables 2–6). Fig. 6 shows the reaction energy diagram for formation of ethylene and water from CO and H₂ in the carbide mechanism. For comparison, this diagram also contains the reaction pathway that leads to methane and water from synthesis gas. The ZPE (zero-point energy)-corrected overall reaction energies for ethylene and methane formation are -212 and -217 kJ mol⁻¹, respectively, which are close to thermodynamic data.

Fig. 7 shows that the preferred pathway for formation of adsorbed CH₂CH₂ involves CH + CH coupling followed by hydrogenation to CH₂CH₂. The C + CH coupling reaction is very unfavourable with a forward barrier of 129 kJ mol⁻¹. The overall barrier for C + CH₃ coupling is also unfavourable, because CH₃ formation is endothermic and the barrier for its coupling to C is also relatively high. Further reaction of CCH₃ to CH₂CH₂ would involve the endothermic dehydrogenation to CCH₂, followed by the endothermic hydrogenation to CHCH₂ ($E_{\text{act}} = 61$ kJ mol⁻¹) and the nearly thermoneutral hydrogenation step to adsorbed CH₂CH₂ with a barrier of 45 kJ mol⁻¹. Although coupling of two CH₂ fragments is facile, their formation is strongly endothermic. As C and CH are the most stable surface intermediates yet coupling reactions between C adsorbates are unfavourable, CH is predicted to be the dominant chain propagation surface intermediate. The desorption energy of adsorbed ethylene is 120 kJ mol⁻¹. The further hydrogenation of ethylene to ethane is included in Fig. 6. The most difficult step towards ethane formation is the hydrogenative desorption of CH₂CH₃ to CH₃CH₃ with an activation energy of 112 kJ mol⁻¹.



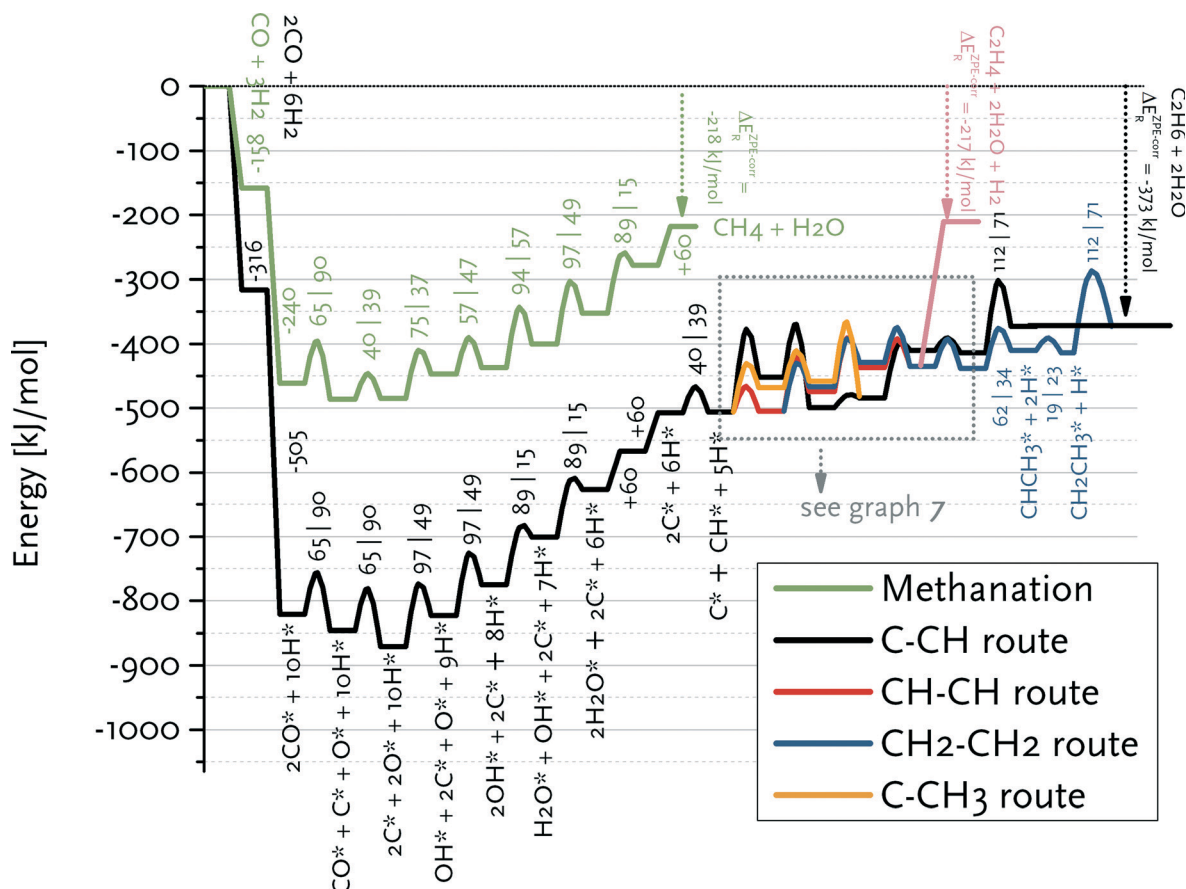


Fig. 6 Reaction energy diagram for ethane, ethylene and methane formation from carbon monoxide and hydrogen. A straight line indicates an adsorption/desorption reaction, whereas the parabolic lines denote a surface elementary reaction step. The adsorption/desorption energies as well as the forward and backward activation energies are given. The detailed pathways for the reactions from adsorbed C^* and CH^* to $CH_2CH_3^*$ and $CHCH_2^*$ are given in Fig. 7.

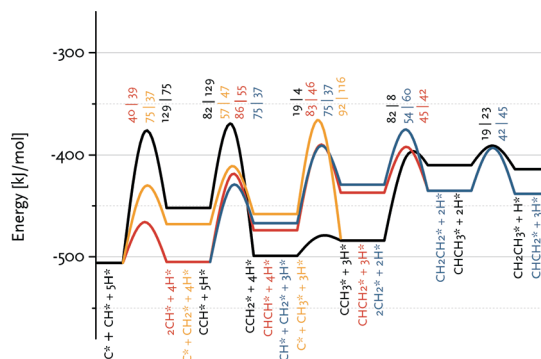


Fig. 7 Enlarged part of the reaction energy diagram showing various routes to ethane and ethylene formation. All data refer to the Ru(1121) surface.

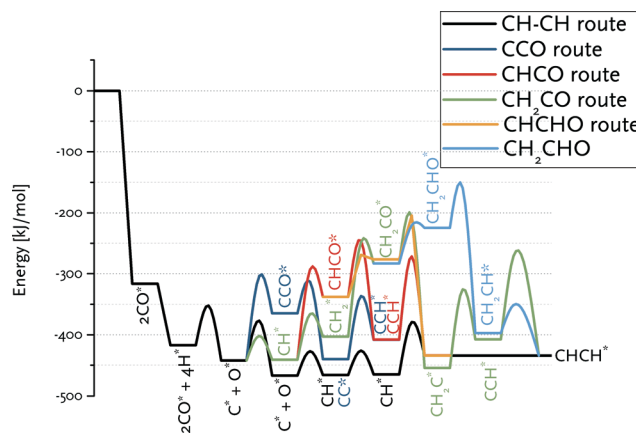


Fig. 8 Reaction energy diagram for $CHCH^*$ formation from CO and H_2 . The various routes in the CO insertion mechanism are compared to the CH-CH route in the carbide mechanism.

Fig. 8 summarizes the pathways relevant to the CO insertion mechanism leading towards $CHCH^*$ formation in a reaction energy diagram. Similar to the carbide mechanism, the growing chain is initiated from adsorbed C^* , which requires CO dissociation. The most favourable path involves $C + CO$ coupling followed by CC formation and its hydrogenation to $CHCH$. As CH together with C are the most stable CH_x surface

intermediates, we also considered $CH + CO$ coupling. The overall barrier to CCH^* formation *via* this route is higher because of the relatively low stability of the $CHCO$ intermediate. This pathway is nevertheless relevant to the CO insertion mechanism, because only the first coupling (*i.e.*, C_2 formation) can



proceed *via* C + CO. Growth beyond C₂ surface species will proceed *via* CR + CO with R being an alkyl group, for which the barrier of the CH + CO coupling is a reasonable estimate. Although the barrier for CH₂ + CO coupling is only slightly higher than for the other considered coupling reactions, the lower surface stability of CH₂ *vs.* C and CH results in a substantially higher overall barrier towards the product without the O atom, namely CH₂C. The alternative pathway *via* CH₂CHO is even less favourable.

Comparison of these paths with that of chain growth *via* CH + CH coupling (Fig. 8) clearly shows that the overall barrier for CHCH formation *via* the carbide mechanism (63 kJ mol⁻¹) is substantially lower than the preferred pathway *via* CO insertion (138 kJ mol⁻¹). The CO insertion route is unfavourable because of the endothermicity of the CH_x + CO coupling (Table 4) as well as of subsequent CH_xCO hydrogenation reactions (Table 5).

Fig. 9 shows the reaction energy diagram for propylene formation within the carbide mechanism. This diagram involves CH as the main inserting species in view of the low barrier for CH insertion into CCH₃ species as compared to the most favourable CH + CH pathway for C₂ formation. Fig. 9 also shows that further growth of the CHCH₂ surface intermediate is favoured over formation and desorption of ethylene. This already indicates that the computed kinetics are conducive to formation of long chain hydrocarbon products on this surface. Compared to the C₂ case, there will be less competition between chain growth of a C₃ intermediate

and desorption of propylene, because the desorption energy of propylene is lower than that of ethylene. We expect that the adsorption energy of longer olefins will be very similar to that of propylene. Qualitatively, the different balances between chain growth and product desorption for C₂ and higher carbon number surface intermediates explains the experimentally observed C₂ selectivity below that of the ASF distribution for the higher hydrocarbon products.¹²

Carbide *vs.* CO insertion mechanism

An important prerequisite for obtaining long hydrocarbon chains in the Fischer–Tropsch reaction is fast chain growth rate *vs.* chain-growth termination rate. This condition will lead to high values for the chain-growth probability α , defined as

$$\alpha = \frac{\theta_i}{\theta_{i-1}} \quad (1)$$

where θ_i is the concentration of growing hydrocarbon chains with length i .

We have earlier deduced explicit expressions for the chain-growth probability within the framework of lumped-sum kinetics for both the carbide and the CO insertion mechanism.^{25,26,60} For the carbide mechanism, the expression is

$$\alpha_c = \frac{k_p \theta_{C_i}}{k_p \theta_{C_i} + k_t} \quad (2)$$

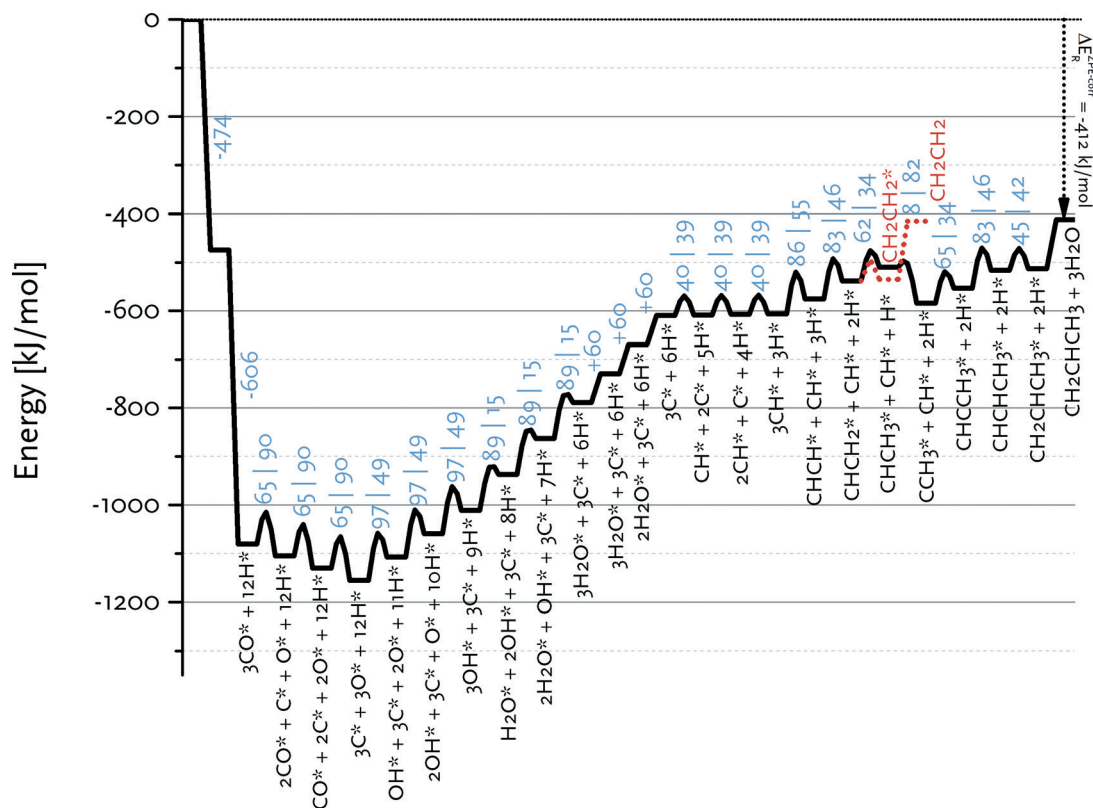


Fig. 9 Reaction energy diagram for propylene from carbon monoxide and hydrogen. All data are for the Ru(1121) surface.



where α_C is the chain-growth probability in the carbide mechanism, k_p the rate constant for $\text{CH}_x + \text{CH}_y$ coupling, θ_C the surface concentration of C_1 monomers and k_t the rate for chain-growth termination. For the CO insertion mechanism, we have deduced the expression

$$\alpha_{\text{CO}} = \frac{k_p \theta_{\text{CO}}}{k_p \theta_{\text{CO}} + k_t} \quad (3)$$

where α_{CO} is the chain-growth probability in the CO insertion mechanism and θ_{CO} the surface concentration of CO. These formulas imply that a high α -value is obtained when the rate constant for chain propagation (k_p) is significantly higher than the rate constant for chain-growth termination (k_t).

To determine the reaction rate constants for the chain growth steps in eqn (2) and (3), we determined the highest overall barrier along the reaction pathway with respect to the energy of the monomeric building block. These building blocks are surface CH and CO adsorbates, respectively, for the carbide and CO insertion mechanisms (details on other reaction steps and further assumptions are in the ESI†). The rate constants are listed in Table 8. Eqn (2) and (3) show that the chain-growth probability will also depend on the surface coverage of the chain propagating intermediate. To understand whether the proposed mechanisms can lead to desired Fischer-Tropsch products, we simply determined the chain-growth probability as a function of these surface coverages. The C_1 coverage is reported to be below 10% under FTS conditions, whereas the CO surface coverage may be assumed to be 90% or higher.^{11,26,60} Fig. 10 shows that the chain-growth probability for the CO insertion mechanism is negligible irrespective of the CO coverage. This analysis implies that CH_x hydrogenation to methane is strongly preferred over chain propagation with CO as the building block. On the contrary, all three pathways with reasonable overall barrier for chain growth in the carbide mechanism, that is CH + CH coupling (96 kJ mol⁻¹), $\text{CH}_2 + \text{CH}_2$ coupling (126 kJ mol⁻¹) and C + CH_3 coupling (136 kJ mol⁻¹) give greater than zero values for the chain-growth probability. At typical C_1 coverage below 10%, the chain-growth probability is highest for the CH + CH coupling pathway.

Table 8 Rate constants computed at $T = 220^\circ\text{C}$ for chain-propagation and chain-growth termination for the most favourable routes in carbide and CO insertion mechanisms

| Route | Mechanism | k_p [mol s ⁻¹] | k_t [mol s ⁻¹] |
|--------------------------|--------------|------------------------------|------------------------------|
| CHCH | Carbide | 6.79×10^2 | 5.88×10^{-4} |
| CCH ₃ | Carbide | 4.51×10^{-1} | 5.88×10^{-4} |
| CH_2CH_2 | Carbide | 3.94×10^{-2} | 5.88×10^{-4} |
| CHCO | CO insertion | 1.74×10^{-8} | 5.88×10^{-4} |
| CHCHO | CO insertion | 1.00×10^{-12} | 5.88×10^{-4} |
| CH_2CO | CO insertion | 3.80×10^{-13} | 5.88×10^{-4} |
| CH_2CHO | CO insertion | 5.84×10^{-17} | 5.88×10^{-4} |

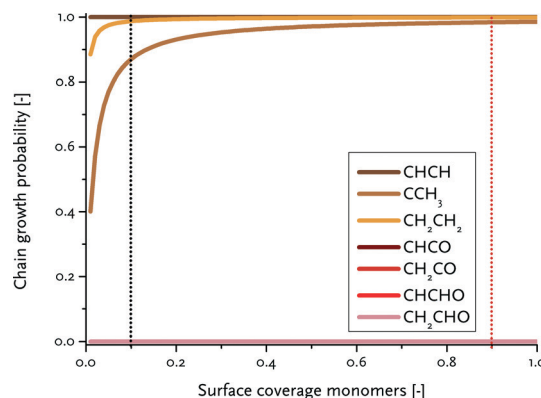


Fig. 10 Computed chain-growth probability (α) as a function of the surface coverage of the chain propagating surface intermediate (CH in the carbide mechanism and CO in the CO insertion mechanism). The black and red dotted lines indicate typical respective C_1 and CO coverage encountered under typical FT reaction conditions.

Conclusions

DFT calculations were carried out for elementary reaction steps of carbide and CO insertion mechanisms relevant to the Fischer-Tropsch reaction on the stepped Ru(11 $\bar{2}$ 1) surface. Activation barriers and reaction energies were determined for CO dissociation, C hydrogenation, $\text{CH}_x + \text{CH}_y$ and $\text{CH}_x + \text{CO}$ coupling, hydrogenation reactions of the surface intermediates, CH_xCH_y -O bond scission and O hydrogenation, which lead to formation of methane, higher hydrocarbons including olefins and alkanes and water. The preferred path for propagation in the carbide mechanism is CH + CH coupling. Higher hydrocarbons are formed by CH insertion into a CCH₃ (or C(CH₂)_nCH₃ equivalents) intermediate, which is formed *via* a sequence of hydrogenation-dehydrogenation steps of adsorbed CHCH. Coupling reactions of the type CH + CCH₃ are more favourable than CH + CH coupling, which is attributed to the σ -donating effect of the alkyl substituent. CH + CH coupling is much more favourable on the stepped site of the Ru(11 $\bar{2}$ 1) surface than on the terrace site. The difference originates from the stronger overlap of the forming π -bonds between the CH fragments with the partially empty d-orbitals of the more reactive metal surface atoms, resulting in decreased Pauli repulsion. The preferred coupling pathway in the CO insertion mechanism involves the reaction between C and CO followed by facile C-O bond cleavage in CCO. All the CH_xCO intermediates are relatively unstable on the surface as compared to CH_x and CO, so that the overall barriers for chain-growth *via* CO-insertion are much higher than those *via* CH + CH coupling. The kinetic consequence of this difference is that high chain-growth rate constants are predicted for the carbide mechanism, whereas these are very low for the CO insertion mechanism. This kinetic analysis predicts methane to be the main product during CO hydrogenation within the CO insertion mechanism. With kinetic parameters computed for the carbide mechanism, high chain-growth probability is predicted. The carbide mechanism



explains the formation of long hydrocarbon chains on the stepped Ru surface.

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Notes and references

- 1 C. Song, *Catal. Today*, 2006, **115**, 2.
- 2 F. Fischer and H. Tropsch, *Brennst.-Chem.*, 1923, **4**, 276.
- 3 G. A. Somorjai, *Introduction to Surface Chemistry and Catalysis*, Wiley, New York, 1994.
- 4 R. B. Anderson, *The Fischer-Tropsch Synthesis*, Academic Press, New York, 1984.
- 5 R. A. van Santen, I. M. Ciobica, E. van Steen and M. M. Ghouri, in *Adv. Catal.*, ed. C. G. Bruce and K. Helmut, Academic Press, 2011, vol. 54, pp. 127–187.
- 6 P. Biloen and W. M. H. Sachtler, in *Adv. Catal.*, ed. H. P. D. D. Eley and B. W. Paul, Academic Press, 1981, vol. 30, pp. 165–216.
- 7 A. T. Bell, *Catal. Rev.: Sci. Eng.*, 1981, **23**, 203–232.
- 8 R. A. van Santen, M. M. Ghouri, S. Shetty and E. J. M. Hensen, *Catal. Sci. Technol.*, 2011, **1**, 891–911.
- 9 A. P. Steynberg, in *Stud. Surf. Sci. Catal.*, ed. S. André and D. Mark, Elsevier, 2004, vol. 152, pp. 1–63.
- 10 A. J. Markvoort, R. A. van Santen, P. A. J. Hilbers and E. J. M. Hensen, *Angew. Chem., Int. Ed.*, 2012, **9**, 9012–9019.
- 11 R. A. van Santen, A. J. Markvoort, M. M. Ghouri, P. A. J. Hilbers and E. J. M. Hensen, *J. Phys. Chem. C*, 2013, **117**, 4488–4504.
- 12 G. P. Van Der Laan and A. A. C. M. Beenackers, *Catal. Rev.: Sci. Eng.*, 1999, **41**, 255–318.
- 13 M. Claeys and E. van Steen, in *Stud. Surf. Sci. Catal.*, ed. S. André and D. Mark, Elsevier, 2004, vol. 152, pp. 601–680.
- 14 S. Novak, R. J. Madon and H. Suhl, *J. Catal.*, 1982, **77**, 141–151.
- 15 E. Iglesia, S. C. Reyes, R. J. Madon and S. L. Soled, in *Adv. Catal.*, ed. H. P. D. D. Eley and B. W. Paul, Academic Press, 1993, vol. 39, pp. 221–302.
- 16 E. W. Kuipers, I. H. Vinkenburg and H. Oosterbeek, *J. Catal.*, 1995, **152**, 137–146.
- 17 E. W. Kuipers, C. Scheper, J. H. Wilson, I. H. Vinkenburg and H. Oosterbeek, *J. Catal.*, 1996, **158**, 288–300.
- 18 B. W. Wojciechowski, *Catal. Rev.: Sci. Eng.*, 1988, **30**, 629–702.
- 19 J. Abbot and B. W. Wojciechowski, *Can. J. Chem. Eng.*, 1988, **66**, 817–824.
- 20 B. Sarup and B. W. Wojciechowski, *Can. J. Chem. Eng.*, 1989, **67**, 620–627.
- 21 M. Ojeda, R. Nabar, A. U. Nilekar, A. Ishikawa, M. Mavrikakis and E. Iglesia, *J. Catal.*, 2010, **272**, 287–297.
- 22 B. T. Loveless, C. Buda, M. Neurock and E. Iglesia, *J. Am. Chem. Soc.*, 2013, **135**, 6107–6121.
- 23 S. Shetty and R. A. van Santen, *Catal. Today*, 2011, **171**, 168–173.
- 24 S. G. Shetty, A. P. J. Jansen and R. A. van Santen, *J. Phys. Chem. C*, 2008, **112**, 14027–14033.
- 25 R. A. Van Santen, A. J. Markvoort, I. A. W. Filot, M. M. Ghouri and E. J. M. Hensen, *Phys. Chem. Chem. Phys.*, 2013, **15**, 17038–17063.
- 26 R. A. van Santen and A. J. Markvoort, *ChemCatChem*, 2013, **5**, 3384–3397.
- 27 T. Zubkov, G. A. Morgan Jr and J. T. Yates Jr, *Chem. Phys. Lett.*, 2002, **362**, 181–184.
- 28 M. Mavrikakis, M. Bäumer, H. J. Freund and J. K. Nørskov, *Catal. Lett.*, 2002, **81**, 153–156.
- 29 B. Hammer and J. K. Nørskov, in *Adv. Catal.*, ed. H. K. Bruce and C. Gates, Academic Press, 2000, vol. 45, pp. 71–129.
- 30 Z.-P. Liu and P. Hu, *J. Am. Chem. Soc.*, 2002, **124**, 11568–11569.
- 31 I. M. Ciobica and R. A. van Santen, *J. Phys. Chem. B*, 2003, **107**, 3808–3812.
- 32 S. Shetty, A. P. J. Jansen and R. A. van Santen, *J. Am. Chem. Soc.*, 2009, **131**, 12874–12875.
- 33 F. Fischer and H. Tropsch, *Brennst.-Chem.*, 1926, **7**, 79–116.
- 34 X.-Q. Gong, R. Raval and P. Hu, *J. Chem. Phys.*, 2005, **122**, 024711.
- 35 I. M. Ciobica, F. Frechard, R. A. van Santen, A. W. Kleyn and J. Hafner, *J. Phys. Chem. B*, 2000, **104**, 3364–3369.
- 36 S. G. Shetty, A. P. J. Jansen and R. A. van Santen, *J. Phys. Chem. C*, 2010, **114**, 22630–22635.
- 37 J. Cheng, P. Hu, P. Ellis, S. French, G. Kelly and M. C. Lok, *J. Phys. Chem. C*, 2008, **112**, 6082–6086.
- 38 M. Zhuo, K. F. Tan, A. Borgna and M. Saeys, *J. Phys. Chem. C*, 2009, **113**, 8357–8365.
- 39 M. Zhuo, A. Borgna and M. Saeys, *J. Catal.*, 2013, **297**, 217–226.
- 40 P. M. Maitlis, H. C. Long, R. Quyoum, M. L. Turner and Z.-Q. Wang, *Chem. Commun.*, 1996, 1–8.
- 41 R. Quyoum, V. Berdini, M. L. Turner, H. C. Long and P. M. Maitlis, *J. Catal.*, 1998, **173**, 355–365.
- 42 M. L. Turner, N. Marsih, B. E. Mann, R. Quyoum, H. C. Long and P. M. Maitlis, *J. Am. Chem. Soc.*, 2002, **124**, 10456–10472.
- 43 X.-Q. Gong, R. Raval and P. Hu, *Surf. Sci.*, 2004, **562**, 247–256.
- 44 X.-Q. Gong, R. Raval and P. Hu, *Mol. Phys.*, 2004, **102**, 993–1000.
- 45 A. Michaelides, A. Alavi and D. A. King, *J. Am. Chem. Soc.*, 2003, **125**, 2746–2755.
- 46 T. W. Zhu, P. W. van Grootel, I. A. W. Filot, S. G. Sun, R. A. van Santen and E. J. M. Hensen, *J. Catal.*, 2013, **297**, 227–235.
- 47 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, **6**, 15–50.
- 48 G. Kresse and J. Hafner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **49**, 14251.
- 49 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- 50 P. E. Blöchl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953.



- 51 G. Kresse and J. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 1758.
- 52 G. Henkelman and H. Jónsson, *J. Chem. Phys.*, 2000, **113**, 9978–9985.
- 53 P. J. Lunde and F. L. Kester, *Ind. Eng. Chem. Process Des. Dev.*, 1974, **13**, 27–33.
- 54 V. Pendyala, W. Shafer and B. Davis, *Catal. Lett.*, 2013, **143**, 895–901.
- 55 C. Zheng, Y. Apeloig and R. Hoffmann, *J. Am. Chem. Soc.*, 1988, **110**, 749–774.
- 56 Z.-X. Chen, H. A. Aleksandrov, D. Basaran and N. Rösch, *J. Phys. Chem. C*, 2010, **114**, 17683–17692.
- 57 J. Chen and Z.-P. Liu, *J. Am. Chem. Soc.*, 2008, **130**, 7929–7937.
- 58 J. Cheng, X.-Q. Gong, P. Hu, C. M. Lok, P. Ellis and S. French, *J. Catal.*, 2008, **254**, 285–295.
- 59 P. W. N. M. van Leeuwen, in *Homogeneous Catalysis*, Springer, Netherlands, 2004, pp. 125–138.
- 60 R. A. van Santen, M. Ghouri and E. M. J. Hensen, *Phys. Chem. Chem. Phys.*, 2014, **16**, 10041–10058.

