

**Quantum chemistry of the Fischer-Tropsch reaction  
catalysed by a stepped Ruthenium surface**

Journal:	<i>Catalysis Science &amp; Technology</i>
Manuscript ID:	CY-ART-04-2014-000483.R1
Article Type:	Paper
Date Submitted by the Author:	19-May-2014
Complete List of Authors:	Filot, Ivo; Eindhoven University of Technology, Department of Chemical Engineering and Chemistry; van Santen, Rutger; Eindhoven University of Technology, Institute for Complex Molecular Systems; Eindhoven University of Technology, Laboratory of Inorganic Materials Chemistry, Department of Chemical Engineering and Chemistry Hensen, Emiel; Eindhoven University of Technology, Department of Chemical Engineering and Chemistry

## ARTICLE

## Quantum chemistry of the Fischer-Tropsch reaction catalysed by a stepped Ruthenium surface

Cite this: DOI: 10.1039/x0xx00000x

I.A.W. Filot, R.A. van Santen and E.J.M. Hensen

Received 00th January 2012,  
Accepted 00th January 2012

DOI: 10.1039/x0xx00000x

[www.rsc.org/](http://www.rsc.org/)

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<sub>x</sub>+CH<sub>y</sub> and CH<sub>x</sub>+CO coupling, CH<sub>x</sub>CH<sub>y</sub>-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  $\sigma$ -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.<sup>1</sup> 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 H<sub>2</sub>), 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.<sup>2-4</sup> Several mechanistic proposals for this chain-growth reaction have been reviewed elsewhere.<sup>5</sup> They can be distinguished by the assumption of the inserting species in the growing chain. The inserting species is either a CH<sub>x</sub> 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<sub>x</sub> 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.<sup>5-8</sup> The competing reaction is complete hydrogenation of the CH<sub>x</sub> surface intermediate to methane, which is an undesired by-product.

Good FT catalysts should exhibit high selectivity towards long-chain hydrocarbons<sup>4, 9</sup> which requires facile CO dissociation and slow chain-growth termination.<sup>10, 11</sup> Formation of methane, the undesired by-product of the FT reaction, should be minimized. The self-organization of monomeric C<sub>1</sub> 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.<sup>12, 13</sup> Experimentally, it is usually found that the C<sub>1</sub> and C<sub>2</sub> product selectivities deviate from the ASF distribution.<sup>12, 14-20</sup>

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.<sup>21, 22</sup> Although metal atoms with low coordination numbers are in principle more reactive, it has been argued that 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.<sup>11, 23-26</sup>

The mechanism of CO dissociation on metallic surfaces has recently been elucidated. Experimental observations<sup>27, 28</sup> and theoretical calculations<sup>24, 29, 30</sup> 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.<sup>23, 24, 31, 32</sup> 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.

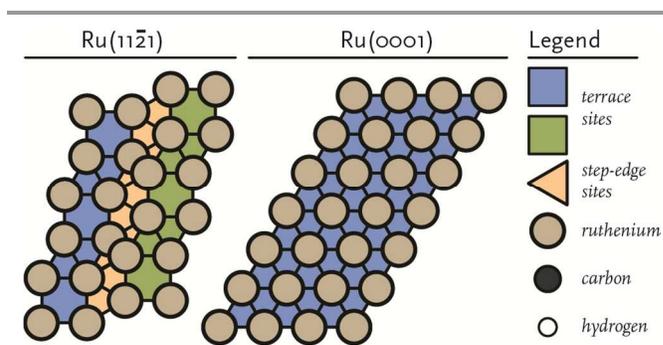
A large number of candidate reaction steps have been proposed for chain growth relevant to carbide and CO insertion mechanisms. Originally, Fischer and Tropsch proposed that surface  $CH_2$  couples with surface  $CH_3$ .<sup>2, 33</sup> Modern insights about surface reactivity and theoretical chemistry advances now learn us that this proposal is incorrect, because the predicted barrier for this coupling step is very high<sup>30</sup> and, also, because the surface coverage of  $CH_2$  and  $CH_3$  is predicted to be low.<sup>34-36</sup> Hu and coworkers<sup>37</sup> 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.<sup>38, 39</sup>  $CH_2+CO$  coupling was found to be the most favourable pathway.<sup>38</sup> The group of Maitlis has performed extensive studies on the mechanistic aspects employing a model system composed using a Co homogeneous catalyst.<sup>40-42</sup> 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.<sup>43-46</sup> 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).<sup>47, 48</sup> The Perdew-Burke-Ernzerhof (PBE) exchange-correlation functional was employed for all calculations.<sup>49</sup> To describe the interaction between nuclei and core electrons, the projector-augmented wave (PAW) method was used.<sup>50, 51</sup> 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 (Figure 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.<sup>10</sup> 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.



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

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 elastic band (NEB) method as developed by Jónsson et al. was employed.<sup>52</sup> 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  $\text{CH}_x$  intermediate and  $\text{CH}_4$ , (iii)  $\text{CH}_x$ - $\text{CH}_x$  coupling, (iv)  $\text{CH}_x$ - $\text{CH}_x$  hydrogenation, (v)  $\text{CH}_x$ -CO coupling, (vi)  $\text{CH}_x\text{CH}_y\text{O}$  hydrogenation, (vii)  $\text{CH}_x\text{CH}_y$ -O scission and (viii) O hydrogenation to  $\text{H}_2\text{O}$ . 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  $\text{CO}_2$  formation, because the  $\text{CO}_2$  selectivity for Ru catalysts in the FT reaction is low.<sup>13, 53, 54</sup>

### 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.<sup>23</sup> 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 $\text{CH}_4$

Hydrogenation of adsorbed C to  $\text{CH}_4$  has been investigated for the Ru(11 $\bar{2}$ 1) surface before.<sup>36</sup> 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  $\text{CH}_x$  ( $x = 0$ -3) to  $\text{CH}_y$  ( $y = 1$ -4) are listed in Table 1.

**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 Eact (kJ/mol)	Backward Eact (kJ/mol)
1	$\text{CO}^* + * \rightarrow \text{C}^* + \text{O}^*$	65	90
2	$\text{C}^* + \text{H}^* \rightarrow \text{CH}^* + *$	40	39
3	$\text{CH}^* + \text{H}^* \rightarrow \text{CH}_2^* + *$	75	37
4	$\text{CH}_2^* + \text{H}^* \rightarrow \text{CH}_3^* + *$	57	47
5	$\text{CH}_3^* + \text{H}^* \rightarrow \text{CH}_4 + 2^*$	94	57

The corresponding initial, transition and final states for these reactions are given in the Supporting Information (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  $\text{CH}_2$  (methylidene) is more difficult ( $E_{\text{act}} = 75$  kJ/mol) and endothermic by 38 kJ/mol. This implies that at reasonable reaction temperatures surface  $\text{CH}_2$  and its hydrogenation products are much less abundant than C and CH surface intermediates. The barrier for the endothermic  $\text{CH}_3$  (methyl) formation is 57 kJ/mol. The barrier for  $\text{CH}_4$  formation, which involves a single elementary reaction step recombining  $\text{CH}_3$  and H over a single Ru atom, has the highest barrier among the  $\text{CH}_x$  hydrogenation steps (94 kJ/mol).

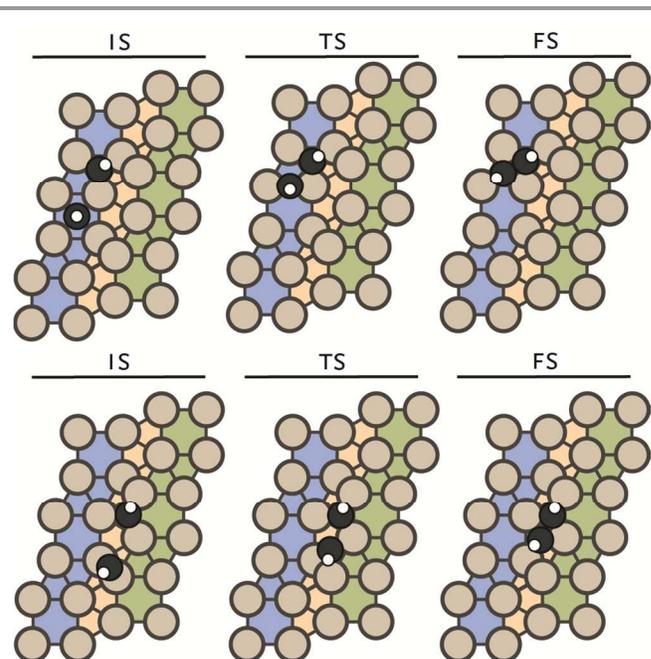
### $\text{CH}_x + \text{CH}_y$ COUPLING REACTIONS

A total of 10 reactions between  $\text{CH}_x$  species were considered for carbon-carbon bond formation to describe chain growth in the carbide mechanism of the Fischer-Tropsch reaction (Table 2).

**Table 2:** CH<sub>x</sub>+CH<sub>y</sub> coupling reactions and their forward and backward activation energies for Ru(11 $\bar{2}$ 1).

Index	Elementary reaction	Forward E <sub>act</sub> (kJ/mol)	Backward E <sub>act</sub> (kJ/mol)
6	C*+C* → CC* + *	138	144
7	C* + CH* → CCH* + *	129	75
8	C* + CH <sub>2</sub> * → CCH <sub>2</sub> * + *	<i>not found</i>	
9	C* + CH <sub>3</sub> * → CCH <sub>3</sub> * + *	92	116
10a	CH* + CH* → CHCH* + * (terrace)	149	117
10b	CH* + CH* → CHCH* + * (step)	86	55
11	CH* + CH <sub>2</sub> * → CHCH <sub>2</sub> * + *	<i>not found</i>	
12	CH* + CH <sub>3</sub> * → CHCH <sub>3</sub> * + *	<i>not found</i>	
13	CH <sub>2</sub> * + CH <sub>2</sub> * → CH <sub>2</sub> CH <sub>2</sub> * + *	54	60
14	CH <sub>2</sub> * + CH <sub>3</sub> * → CH <sub>2</sub> CH <sub>3</sub> * + *	<i>not investigated</i>	
15	CH <sub>3</sub> * + CH <sub>3</sub> * → CH <sub>3</sub> CH <sub>3</sub> * + *	<i>not investigated</i>	

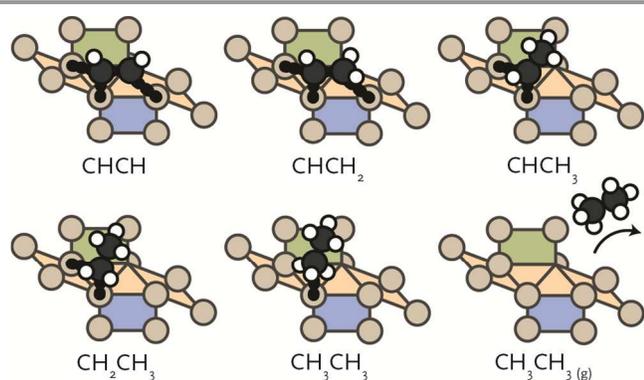
The corresponding initial, transition and final states are given in the Supporting Information. Reactions between CH<sub>2</sub> and CH<sub>3</sub> and between two CH<sub>3</sub> (reactions 14 and 15) were not investigated, because it is well known that the interaction of the spatially extended C-H bonds in CH<sub>3</sub> adsorbates results in repulsion, precluding C-C bond formation.<sup>55</sup> Despite considerable efforts, we could not identify transition states for the following reactions: C+CH<sub>2</sub>, CH+CH<sub>2</sub> and CH+CH<sub>3</sub> 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.<sup>56</sup> For the remaining five coupling steps C+C, C+CH, CH+CH, CH<sub>2</sub>+CH<sub>2</sub> and C+CH<sub>3</sub> coupling, we identified six unique transition states. For the CH+CH coupling step, two different pathways were found (see Figure 2).

**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.

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 = CH<sub>3</sub> 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 preference for coupling on the step-edge site is in accordance with several other works.<sup>37, 57, 58</sup> 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 methylidene species with a barrier of 54 kJ/mol (reaction 6). The barrier for the other coupling reactions are higher: 86 kJ/mol for CH+CH, 92 kJ/mol for C+CH<sub>3</sub>, 129 kJ/mol for C+CH and 138 kJ/mol for coupling of two carbon adatoms (C+C).

### CH<sub>x</sub>CH<sub>y</sub> HYDROGENATION REACTIONS

After formation of carbon-carbon bonds, higher hydrocarbons may leave the surface as olefins or paraffins (see Figure 3). We investigated such chain-growth termination steps for CH<sub>x</sub>-CH<sub>y</sub> species. From CCH, a large number of hydrogenation routes to ethylene and ethane needs to be considered. These possibilities and the results of a reaction energy analysis are given in Table 3.



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

**Table 3:** Hydrogenation reactions of C<sub>2</sub> surface intermediates and their forward and backward activation energies for Ru(111).

Index	Elementary reaction	Forward E <sub>act</sub> (kJ/mol)	Backward E <sub>act</sub> (kJ/mol)
16	CC* + H* → CCH* + *	104	72
17	CCH* + H* → CCH <sub>2</sub> * + *	82	129
18	CCH <sub>2</sub> * + H* → CCH <sub>3</sub> * + *	19	4
19	CHCH* + H* → CHCH <sub>2</sub> * + *	83	46
20	CHCH <sub>2</sub> * + H* → CHCH <sub>3</sub> * + *	62	34
21	CCH* + H* → CHCH* + *	140	162
22	CCH <sub>2</sub> * + H* → CHCH <sub>2</sub> * + *	82	21
23	CCH <sub>3</sub> * + H* → CHCH <sub>3</sub> * + *	82	8
24	CHCH <sub>2</sub> * + H* → CH <sub>2</sub> CH <sub>2</sub> * + *	45	42
25	CHCH <sub>3</sub> * + H* → CH <sub>2</sub> CH <sub>3</sub> * + *	19	23
26	CH <sub>2</sub> CH <sub>2</sub> * + H* → CH <sub>2</sub> CH <sub>3</sub> * + *	58	34
27	CH <sub>2</sub> CH <sub>3</sub> * + H* → CH <sub>3</sub> CH <sub>3</sub> * + *	112	71

The structures of the involved stable and transition states are collected in the Supporting Information. Hydrogenation of CCH to CCH<sub>2</sub> (reaction 17) and CHCH to CHCH<sub>2</sub> (reaction 19) have very similar forward activation barriers around 82 kJ/mol. While CCH<sub>2</sub> formation is exothermic, CHCH<sub>2</sub> formation is endothermic. Subsequent hydrogenation to CCH<sub>3</sub> (reaction 18) and CHCH<sub>3</sub> (reaction 20) proceeds with barriers of 19 and 62 kJ/mol, 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, respectively. Hydrogenation of CCH<sub>2</sub> to CHCH<sub>2</sub> (reaction 22) and of CCH<sub>3</sub> to CHCH<sub>3</sub> (reaction 23) have barriers of 82 kJ/mol. The reverse dehydrogenation reactions are very facile with barriers of 21 and 8 kJ/mol, respectively. This implies that

CHCH<sub>x</sub> intermediates are significantly more stable than the corresponding CHCH<sub>x</sub> intermediates. This difference draws similarity to the higher stability of adsorbed CH over CH<sub>2</sub>. 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 and is almost thermodynamically neutral, the hydrogenation of CHCH<sub>3</sub> to CH<sub>2</sub>CH<sub>3</sub> and, finally, CH<sub>3</sub>CH<sub>3</sub> is endothermic. Although the first hydrogenation step to CH<sub>2</sub>CH<sub>3</sub> is facile and slightly exothermic, the second hydrogenation step toward ethane has to overcome a relatively high barrier of 112 kJ/mol. It is endothermic by 41 kJ/mol. Alternatively, adsorbed ethylene can be further hydrogenated to CH<sub>2</sub>CH<sub>3</sub> (reaction 26). Although this reaction is facile with an activation barrier of 58 kJ/mol, the reverse reaction is more favourable with an activation energy of 34 kJ/mol.

### CH<sub>x</sub>+CO COUPLING

Three different reactions between CH<sub>x</sub> 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<sub>x</sub> adsorption site followed by C-C bond formation. Both carbon atoms in the final coupled CH<sub>x</sub>-CO species coordinate to surface Ru atoms. This strongly differs from mechanisms proposed based on homogeneous mononuclear coordination complexes for CO insertion or migration.<sup>59</sup>

**Table 4:** CH<sub>x</sub>+CO coupling reactions and their forward and backward activation energy.

Index	Elementary reaction	Forward E <sub>act</sub> (kJ/mol)	Backward E <sub>act</sub> (kJ/mol)
28	C* + CO* → CCO* + *	138	61
29	CH* + CO* → CHCO* + *	148	44
30	CH <sub>2</sub> * + CO* → CH <sub>2</sub> CO* + *	155	36
31	CH <sub>3</sub> * + CO* → CH <sub>3</sub> CO* + *	<i>not investigated</i>	

The corresponding initial, transition and final states are given in the Supporting Information. The reaction between CH<sub>3</sub> 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. The barrier for CH+CO coupling (reaction 29) and CH<sub>2</sub>+CO coupling (reaction 30) are somewhat higher at 148 kJ/mol and 155 kJ/mol, 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<sub>x</sub>+CH<sub>y</sub> coupling (Table 2). Moreover, all CO insertion reactions are strongly endothermic by at least 77 kJ/mol, implying that the rate constants for the reverse reactions are substantially higher..

### CH<sub>x</sub>-CO HYDROGENATION

After CO insertion, the resulting  $\text{CH}_x\text{CO}$  moiety can be hydrogenated to form a  $\text{CH}_x\text{CH}_y\text{O}$  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  $\text{CH}_x\text{CO}^*$  intermediate, the hydrogenation of either carbon atom was considered. This results in five different hydrogenation reactions (Table 5).

**Table 5:** Hydrogenation reactions of  $\text{CH}_x\text{CO}$  surface intermediates and their forward and backward activation energies for Ru(11 $\bar{1}$ 1).

Index	Elementary reaction	Forward $E_{\text{act}}$ (kJ/mol)	Backward $E_{\text{act}}$ (kJ/mol)
32	$\text{CCO}^* + \text{H}^* \rightarrow \text{CHCO}^* + *$	104	77
33	$\text{CCO}^* + \text{H}^* \rightarrow \text{CCHO}^* + *$	<i>not found</i>	
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

We could not identify a transition state for  $\text{CCO}^*$  hydrogenation (reaction 33). The orientation of the  $\text{CCO}^*$  moiety in the step-edge site (see the Supporting Information) prevents hydrogenation of the carbon atom of the CO group. The most difficult hydrogenation step is that of  $\text{CCO}^*$  to  $\text{CHCO}^*$  (reaction 32), which involves an activation energy of 104 kJ/mol. The other reactions, *i.e.* the hydrogenation of  $\text{CHCO}^*$  to  $\text{CH}_2\text{CO}^*$ , that of  $\text{CHCO}^*$  to  $\text{CHCHO}^*$  and finally the hydrogenation of  $\text{CH}_2\text{CO}^*$  to  $\text{CH}_2\text{CHO}^*$  proceed with lower activation energies of 95, 68 and 63 kJ/mol, respectively. All  $\text{CH}_x\text{CO}$  hydrogenation reactions are endothermic.

### $\text{CH}_x\text{CH}_y\text{-O}$ BOND SCISSION

For chain growth to proceed, the C-O bond in the  $\text{CH}_x\text{CH}_y\text{O}$  surface intermediate must first cleave. Five such elementary reaction steps were considered for which results are listed in Table 6.

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

Index	Elementary reaction	Forward $E_{\text{act}}$ (kJ/mol)	Backward $E_{\text{act}}$ (kJ/mol)
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

Among these reactions, it is most easy to cleave the C-O bond in  $\text{CCO}^*$  with an activation energy of 52 kJ/mol. 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. 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 to  $\text{H}_2\text{O}$  (reaction 43) or via proton migration between two OH species to form  $\text{H}_2\text{O}$  and O (reaction 44). The first hydrogenation step to produce adsorbed OH has a relatively high barrier of 97 kJ/mol. The barrier for water formation via OH hydrogenation is 89 kJ/mol. Comparatively, the reaction between two hydroxyl groups is much more facile with a barrier of 54 kJ/mol. This suggests that water formation by reaction of two OH groups to  $\text{H}_2\text{O}$  and adsorbed O is preferred.

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

Index	Elementary reaction	Forward $E_{\text{act}}$ (kJ/mol)	Backward $E_{\text{act}}$ (kJ/mol)
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

### 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  $\text{CCH}_3$  on stepped sites, because the latter reaction is relevant for formation of  $\text{C}_3$  products. Finally, we will highlight the different adsorption strengths of ethylene and propylene on stepped sites, which explains the lower than expected  $\text{C}_2$  selectivity observed during the FT reaction.

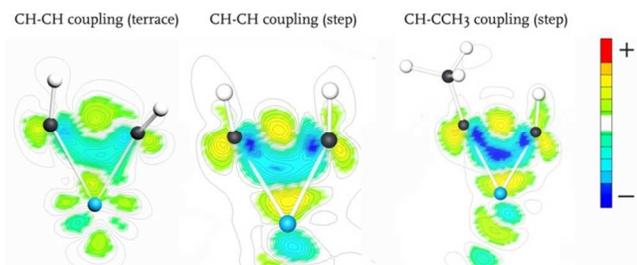
### CH+CH COUPLING

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

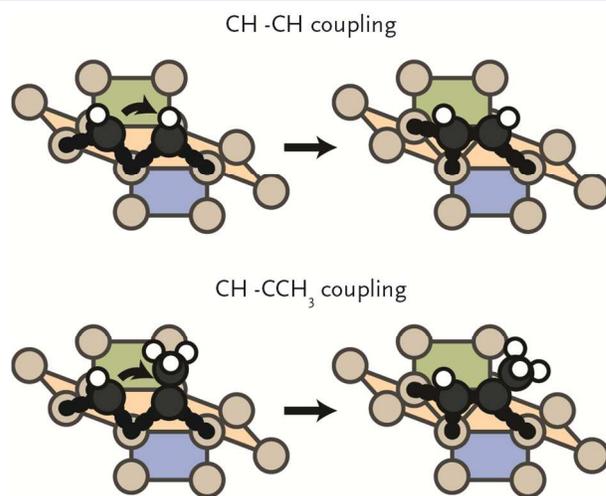
Figure 4a shows the electron density shifts that result from the formation of C-C bonds between two CH surface species in the corresponding transition state complexes. To determine these, the electron density distributions of the two separate CH fragments and the empty surface were subtracted from the 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<sub>3</sub> COUPLING

We found that methylidyne (CH) coupling to the CCH<sub>3</sub> surface intermediate (Figure 4b) proceeds with lower barrier than CH coupling to another CH species (65 vs. 86 kJ/mol). In Figure 4a, the electron density shifts for C-C bond formation are shown. It can be seen that in the transition state for CH+CCH<sub>3</sub> coupling more electron density is shifted from between the CH and CCH<sub>3</sub> species to the metal atom relative to the CH-CH case. This difference is caused by the  $\sigma$ -donating effect of the methyl group. Consequently, the Pauli repulsion between CH and CCH<sub>3</sub> is lower than between two CH fragments, explaining why the barrier for coupling of the former two fragments is lower.



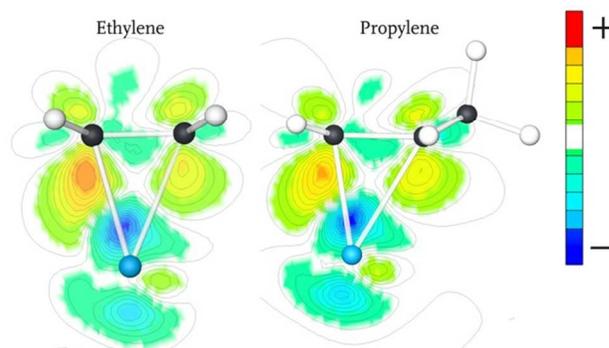
**Fig 4a** Electron density difference plots of the transition state for CH+CH coupling on the terrace (left), step-edge (middle) and of CH+CCH<sub>3</sub> coupling on the step-edge (right) of the Ru(11 $\bar{2}$ 1) surface. These plots were generated by subtracting the electron density of the individual 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.



**Fig 4b** Schematic representation of CH-CH and CH-CCH<sub>3</sub> coupling on a step-edge site of the Ru(11 $\bar{2}$ 1) surface. The initial and final state of reaction are displayed.

#### 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, respectively. The electron density shifts upon olefin adsorption are shown in Figure 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  $\pi$ -accepting nature of the CH<sub>3</sub> substituent. This results in weaker adsorption of propylene as 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.



**Fig. 5** Electron density difference plots of the adsorbed states of ethylene (left) and propylene (right) on the Ru(11 $\bar{2}$ 1) 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.



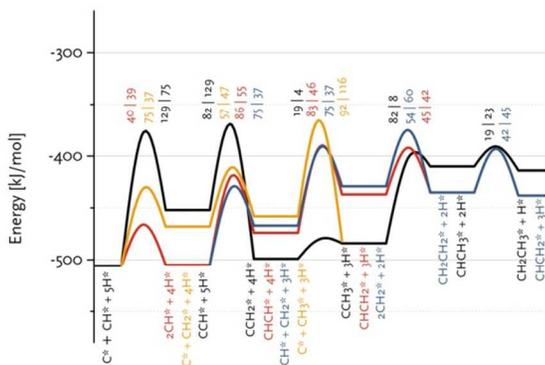


Fig. 7 Enlarged part of the reaction energy diagram showing various routes to ethane and ethylene formation. All data refer to the Ru(11 $\bar{1}$ 1) surface.

Figure 7 shows that the preferred pathway for formation of adsorbed  $\text{CH}_2\text{CH}_2$  involves  $\text{CH}+\text{CH}$  coupling followed by hydrogenation to  $\text{CH}_2\text{CH}_2$ . The  $\text{C}+\text{CH}$  coupling reaction is very unfavourable with a forward barrier of 129 kJ/mol. The overall barrier for  $\text{C}+\text{CH}_3$  coupling is also unfavourable, because  $\text{CH}_3$  formation is endothermic and the barrier for its coupling to  $\text{C}$  is also relatively high. Further reaction of  $\text{CCH}_3$  to  $\text{CH}_2\text{CH}_2$  would involve the endothermic dehydrogenation to  $\text{CCH}_2$ , followed by the endothermic hydrogenation to  $\text{CHCH}_2$  ( $E_{\text{act}} = 61$  kJ/mol) and the nearly thermoneutral hydrogenation step to adsorbed  $\text{CH}_2\text{CH}_2$  with a barrier of 45 kJ/mol. Although coupling of two  $\text{CH}_2$  fragments is facile, their formation is strongly endothermic. As  $\text{C}$  and  $\text{CH}$  are the most stable surface intermediates yet coupling reactions between  $\text{C}$  adsorbates are unfavourable,  $\text{CH}$  is predicted to be the dominant chain propagation surface intermediate. The desorption energy of adsorbed ethylene is 120 kJ/mol. The further hydrogenation of ethylene to ethane is included in Figure 6. The most difficult step towards ethane formation is the hydrogenative desorption of  $\text{CH}_2\text{CH}_3$  to  $\text{CH}_3\text{CH}_3$  with an activation energy of 112 kJ/mol.

Figure 8 summarizes the pathways relevant to the  $\text{CO}$  insertion mechanism leading towards  $\text{CHCH}^*$  formation in a reaction energy diagram. Similar to the carbide mechanism, the growing chain is initiated from adsorbed  $\text{C}^*$ , which requires  $\text{CO}$  dissociation. The most favourable path involves  $\text{C}+\text{CO}$  coupling followed by  $\text{CC}$  formation and its hydrogenation to  $\text{CHCH}$ . As  $\text{CH}$  together with  $\text{C}$  are the most stable  $\text{CH}_x$  surface intermediates, we also considered  $\text{CH}+\text{CO}$  coupling. The overall barrier to  $\text{CCH}^*$  formation via this route is higher because of the relatively low stability of the  $\text{CHCO}$  intermediate. This pathway is nevertheless relevant to the  $\text{CO}$  insertion mechanism, because only the first coupling (*i.e.*,  $\text{C}_2$  formation) can proceed via  $\text{C}+\text{CO}$ . Growth beyond  $\text{C}_2$  surface species will proceed via  $\text{CR}+\text{CO}$  with  $\text{R}$  being an alkyl group, for which the barrier of the  $\text{CH}+\text{CO}$  coupling is a reasonable estimate. Although the barrier for  $\text{CH}_2+\text{CO}$  coupling is only slightly higher than for the other considered coupling reactions,

the lower surface stability of  $\text{CH}_2$  vs.  $\text{C}$  and  $\text{CH}$  results in a substantially higher overall barrier towards the product without the  $\text{O}$  atom, namely  $\text{CH}_2\text{C}$ . The alternative pathway via  $\text{CH}_2\text{CHO}$  is even less favourable.

Comparison of these paths with that of chain growth via  $\text{CH}+\text{CH}$  coupling (Figure 8) clearly shows that the overall barrier for  $\text{CHCH}$  formation via the carbide mechanism (63 kJ/mol) is substantially lower than the preferred pathway via  $\text{CO}$  insertion (138 kJ/mol). The  $\text{CO}$  insertion route is unfavourable because of the endothermicity of the  $\text{CH}_x+\text{CO}$  coupling (Table 4) as well as of subsequent  $\text{CH}_x\text{CO}$  hydrogenation reactions (Table 5).

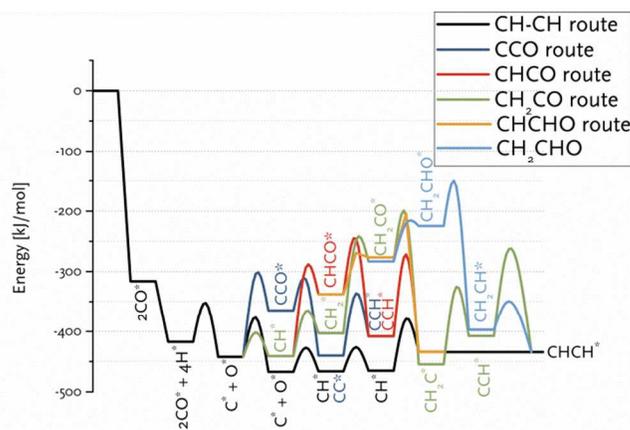


Fig. 8 Reaction energy diagram for  $\text{CHCH}^*$  formation from  $\text{CO}$  and  $\text{H}_2$ . The various routes in the  $\text{CO}$  insertion mechanism are compared to the  $\text{CH}-\text{CH}$  route in the carbide mechanism.

Figure 9 shows the reaction energy diagram for propylene formation within the carbide mechanism. This diagram involves  $\text{CH}$  as the main inserting species in view of the low barrier for  $\text{CH}$  insertion into  $\text{CCH}_3$  species as compared to the most favourable  $\text{CH}+\text{CH}$  pathway for  $\text{C}_2$  formation. Figure 9 also shows that further growth of the  $\text{CHCH}_2$  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  $\text{C}_2$  case, there will be less competition between chain growth of a  $\text{C}_3$  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  $\text{C}_2$  and higher carbon number surface intermediates explains the experimentally observed  $\text{C}_2$  selectivity below that of the ASF distribution for the higher hydrocarbon products.<sup>12</sup>

#### Carbide vs. $\text{CO}$ 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  $\alpha$ , defined as

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

where  $\theta_i$  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.<sup>25, 26, 60</sup> For the carbide mechanism, the expression is

$$\alpha_C = \frac{k_p \theta_{C_1}}{k_p \theta_{C_1} + k_t} \quad (2)$$

where  $\alpha_C$  is the chain-growth probability in the carbide mechanism,  $k_p$  the rate constant for  $\text{CH}_x + \text{CH}_y$  coupling,  $\theta_{C_1}$  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

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

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

**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]	$k_t$ [mol/s]
CHCH	Carbide	$6.79 \cdot 10^2$	$5.88 \cdot 10^{-4}$
CCH <sub>3</sub>	Carbide	$4.51 \cdot 10^{-1}$	$5.88 \cdot 10^{-4}$
CH <sub>2</sub> CH <sub>2</sub>	Carbide	$3.94 \cdot 10^{-2}$	$5.88 \cdot 10^{-4}$
CHCO	CO insertion	$1.74 \cdot 10^{-8}$	$5.88 \cdot 10^{-4}$
CHCHO	CO insertion	$1.00 \cdot 10^{-12}$	$5.88 \cdot 10^{-4}$
CH <sub>2</sub> CO	CO insertion	$3.80 \cdot 10^{-13}$	$5.88 \cdot 10^{-4}$
CH <sub>2</sub> CHO	CO insertion	$5.84 \cdot 10^{-17}$	$5.88 \cdot 10^{-4}$

To determine the reaction rate constants for the chain growth steps in Eqs. 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 Supporting Information). The rate constants are listed in Table 8. Equations 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.<sup>11, 26, 60</sup> Figure 10 shows that the chain-growth probability for the CO insertion mechanism is negligible irrespective of the CO coverage. This analysis implies that  $\text{CH}_x$  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),  $\text{CH}_2 + \text{CH}_2$  coupling (126 kJ/mol) and C+CH<sub>3</sub> coupling (136 kJ/mol) 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.

## ARTICLE

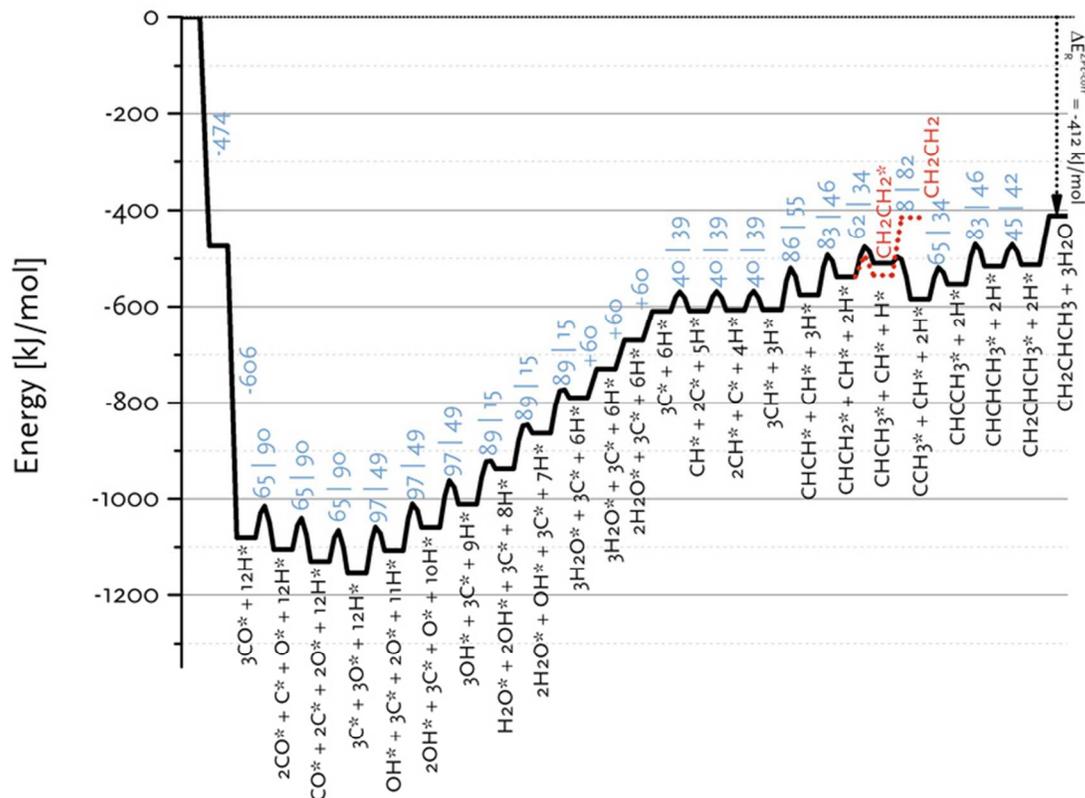


Fig 9 Reaction energy diagram for propylene from carbon monoxide and hydrogen. All data are for the Ru(11 $\bar{2}$ 1) surface.

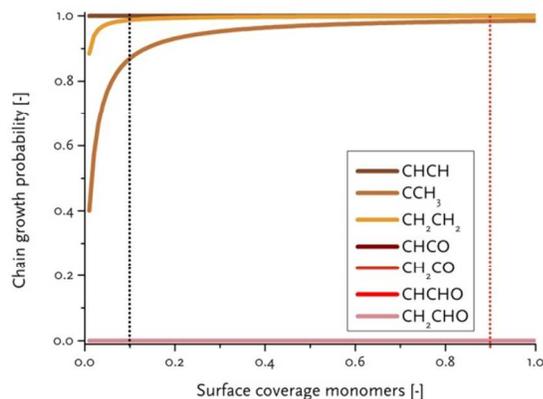


Fig 10 Computed chain-growth probability ( $\alpha$ ) as a function of the surface coverage of the chain propagating surface intermediate (CH in the carbide mechanism and CO in the CO insertion mechanism). The black and red dotted lines indicate typical respective C1 and CO coverage encountered under 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, CH<sub>x</sub>+CH<sub>y</sub> and CH<sub>x</sub>+CO coupling, hydrogenation reactions of the surface intermediates, CH<sub>x</sub>CH<sub>y</sub>-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 CCH<sub>3</sub> (or C(CH<sub>2</sub>)<sub>n</sub>CH<sub>3</sub> equivalents) intermediate, which is formed via a sequence of hydrogenation-dehydrogenation steps of adsorbed CHCH. Coupling reactions of the type CH+CCH<sub>3</sub> are more favourable than CH+CH coupling, which is attributed to the  $\sigma$ -donating effect of the alkyl substituent. CH+CH coupling is much more favourable in 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  $\pi$ -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  $\text{CH}_x\text{CO}$  intermediates are relatively unstable on the surface as compared to  $\text{CH}_x$  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.

### Acknowledgements

NWO is acknowledged for providing access to the supercomputer facilities.

### Notes and references

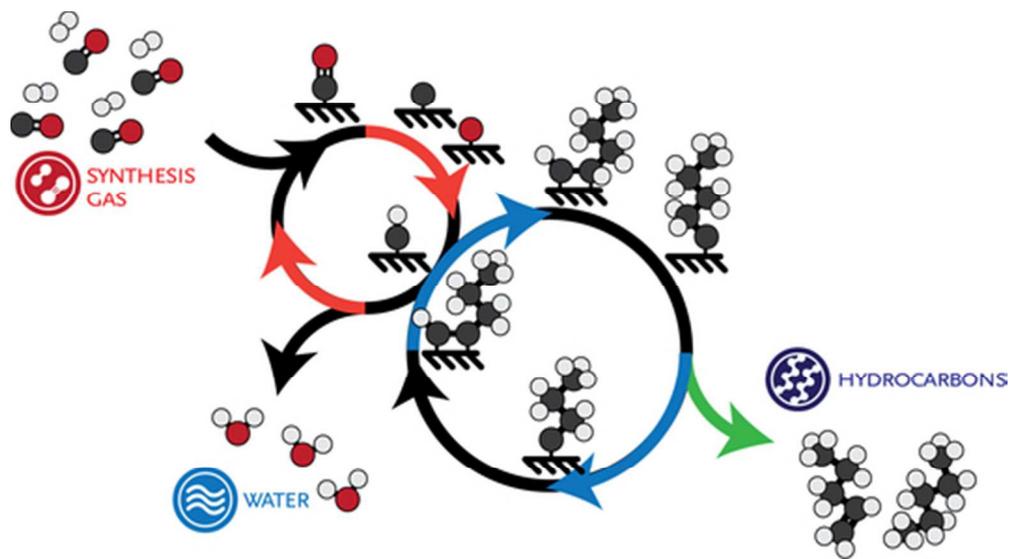
<sup>a</sup> Laboratory of Inorganic Materials Chemistry, Schuit Institute of Catalysis, Technische Universiteit Eindhoven, P.O. Box 513, 5600 MB Eindhoven, The Netherlands

Electronic Supplementary Information (ESI) available: Geometries and energies of the initial, transition and final states used in the electronic structure calculations. See DOI: 10.1039/b000000x/

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.*, eds. C. G. Bruce and K. Helmut, Academic Press, 2011, vol. Volume 54, pp. 127-187.
6. P. Biloen and W. M. H. Sachtler, in *Adv. Catal.*, eds. H. P. D.D. Eley and B. W. Paul, Academic Press, 1981, vol. Volume 30, pp. 165-216.
7. A. T. Bell, *Catal. Rev.*, 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.*, eds. S. André and D. Mark, Elsevier, 2004, vol. Volume 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.*, 1999, **41**, 255-318.
13. M. Claeys and E. van Steen, in *Stud. Surf. Sci. Catal.*, eds. S. André and D. Mark, Elsevier, 2004, vol. Volume 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.*, eds. H. P. D.D. Eley and B. W. Paul, Academic Press, 1993, vol. Volume 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.*, 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 C. Gates, Academic Press, 2000, vol. Volume 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, *J. Comput. Mater. Sci.*, 1996, **6**, 15-50.
48. G. Kresse and J. Hafner, *Phys. Rev. B*, 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*, 1994, **50**, 17953.
51. G. Kresse and J. Joubert, *Phys. Rev. B*, 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.

## Journal Name

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.



45x24mm (300 x 300 DPI)