Identification of hydrogen species on Pt/Al$_2$O$_3$ by in situ inelastic neutron scattering and their reactivity with ethylene
Identification of hydrogen species on Pt/Al₂O₃ by in situ inelastic neutron scattering and their reactivity with ethylene†

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Hydrogen (H) species adsorbed on catalyst surfaces are key intermediates in catalytic hydrogenation reactions over supported metal catalysts. However, individual identification of H species on the metal catalysts has not been established to date. Here, we elucidated the H species on Pt/Al₂O₃ by the combination of in situ inelastic neutron scattering (INS) and density functional theory (DFT). Several H species in the presence of H₂ were successfully identified at different sites on the Pt surface and Al₂O₃ support. The in situ INS and FT-IR measurements revealed that the hydride/atop Pt–H, bridged perimeter/tierce Pt–H–Pt, and threefold Pt₃–H are active intermediates in the C₂H₄ hydrogenation reaction, whereas the edge Pt–H–Pt is a rather inert species. In addition, the in situ measurements indicate that the Al₂O₃ itself acts as hydrogen storage material by mediating AlO–H and Al–H–Al species, which serve H species in the C₂H₄ hydrogenation reaction.

Introduction

Hydrogen (H) species is crucial intermediates in hydrogenation and reduction reactions in chemical synthesis and environmental chemistry. In fine chemical synthesis, the selective hydrogenation reactions of functional groups such as –NO₂, –C=O, –C≡C, –C=O and –C≡N require the use of H₂ molecules to afford corresponding amines, alkanes, alkenes, alcohols and amines, respectively, which are essential for the chemical industry. Supported metal catalysts are among the most promising candidates for selective hydrogenation reactions in practical applications because they are active, easily separated from products and reusable. The catalytic activity and selectivity of supported metal catalysts are dependent on the size, composition, geometric structure, surface coordination environment of the metal particles and also the properties of the support, which affect the electronic structure of the metals and provide the sites available for absorption and reaction in many cases. These structural factors induce a variety of structures and reactivity of the H species. One typical mechanism for the formation of the active H species proceeds by the homolytic dissociation of H₂ on metals such as Pt, Pd and Rh. The occupied d-orbitals of these metals donate their electrons to the antibonding orbital of H₂ to weaken the H–H bond. Another mechanism for the formation of the active H species proceeds by heterolytic dissociation of H₂ into H⁺ and H⁻ species, which preferentially react with polar functional groups rather than nonpolar ones, typically at the interface between metal particles and metal oxide supports. Thus, it is essential to elucidate both the adsorbed state and the dynamics of H species for the research and development of hydrogenation reactions.

The activation of H₂ and the dynamics of H species on the metal particles supported on metal oxide have been studied to understand the role of metals and supports during various hydrogenation reactions. Zaera reported that H₂ adsorption
(activation) on the Pt surface is the rate-determining step for the ethylene hydrogenation reaction. High activity and selectivity in the hydrogenation of acetylene to ethylene were achieved by promoting the activation of H₂ and anticokeing, which were induced by Pd catalysts. In the case of the de-NO reaction with H₂ on Pt/MgO and Pt/CeO₂ catalysts, the H species activated by the Pt metal was found to migrate to the MgO and CeO₂ supports and react with the NO species adsorbed on these supports. The H spillover on metal oxides well depends on the surface properties of the metal oxide supports, which may change during the hydrogenation reactions. However, these reaction mechanisms have been based on the reaction results and structural characterization, but not yet supported by direct observation of the H species involved.

Inelastic neutron scattering (INS) spectroscopy is a powerful technique to investigate the H species adsorbed on the supported metal catalysts because of the extremely large cross section of neutron from ¹H. INS spectroscopy is advantageous for analysing low-frequency vibration modes, such as vibration modes of functional groups containing H atom, and vibrational states of H atoms in bulk Pd metal and nanocrystalline Pd metal. The activated H species on metals or metal particles have been studied by INS to elucidate the dynamics of the H species and the catalysis involved. INS was utilized to discriminate the vibrational modes of H species on different adsorption sites of the Pt particles immobilized on a carbon support, which was correlated with the specific electrocatalytic activity. Recently, the active H species on a 5 wt% Pt/C catalyst were detected by INS. Evidence of the H spillover from Pt to unsaturated reactive sites in the carbon was also provided. In the case of a Pt/C catalyst, the heterolytic bond cleavage of H₂ was reported by the combination of INS and FT-IR. In a recent study, Pt–H species with n-fold coordination were detected on a Pt/Al₂O₃ catalyst by INS. These recent results have demonstrated that the INS technique is effective to detect active H species on supported metal catalysts. However, the individual assignment of each H species on catalysts is still a challenge in the field of catalytic science.

In this study, we tackled this challenging issue, specifically, individual identification of each H species on a Pt/Al₂O₃ catalyst in the ethylene hydrogenation reaction using in situ INS. We successfully identified several H species on the Pt particles and H species that migrated from the Pt particles to the Al₂O₃ support by the combination of in situ INS and density functional theory (DFT) calculations for the first time. In addition, we demonstrated that the Pt–H at atop sites, the Pt–H–Pt at bridged perimeter and terrace sites, and Pt–H at threefold sites were active intermediates in C₂H₄ hydrogenation by the in situ INS and FT-IR.

**Experimental**

**Preparation of 5 wt% Pt/Al₂O₃ catalysts**

An aqueous nitric acid solution containing cis-[Pt(NH₃)₂(NO₂)₂], in which the content of Pt was 4.64 wt%, was purchased from Furuya Metal Corporation Ltd. γ-Al₂O₃ (JRC-ALO-7) of 180 m² g⁻¹ was provided by the Catalysis Society of Japan. A Pt/Al₂O₃ catalyst was prepared by an impregnation method. The loading amount of Pt was 5.0 wt% as a metal basis. γ-Al₂O₃ (10 g) was added to an aqueous solution (100 mL) containing cis-[Pt(NH₃)₂(NO₂)₂] solution (11.3 g), followed by solvent evaporation at 80 °C. The resulting powder was calcined in air at 673 K for 5 h. Thus-obtained catalyst was reduced under a H₂ gas flow (5% H₂/N₂, 50 mL min⁻¹) at 623 K for 2 h.

**Structural characterization**

Crystal structure of the Pt/Al₂O₃ was analysed by a Ultima IV X-ray diffractometer (Rigaku, Japan). The local structure of the Pt/Al₂O₃ was investigated by X-ray absorption fine structure (XAFS). XAFS measurements were performed at the BL01B1 beamline of SPring-8 facility of the Japan Synchrotron Radiation Research Institute. A Si(111) double-crystal monochromator was used to obtain the incident X-ray beam. Pt L₂,₃-edge XAFS spectra of Pt foil and the Pt/Al₂O₃ were recorded in transmission mode using ionization chambers at room temperature in air and He, respectively. Energy was calibrated using Pt foil. Before the XAFS measurement of the Pt/Al₂O₃, it was heated at 673 K under 5% H₂/He (flow rate: 100 mL min⁻¹) conditions for 1 h. XAFS spectra were analyzed using a xTunes software. After normalization, k¹-weighted χ spectra in the k range of 3.0–16.0 Å⁻¹ were Fourier-transformed into r space to obtain FT-EXAFS spectra. The curve fitting analysis was conducted in the range of 1.9–2.9 Å using a FEFF8 program.

**Catalytic reactions**

Ethylene hydrogenation reaction was carried out using a fixed-bed flow reactor at 303 K. In this experiment, the total flow rate of gas was fixed to 100 mL min⁻¹. Before the activity test, the Pt/Al₂O₃ catalyst (10.3 mg) was reduced by 5% H₂/He at 473 K for 60 min, and then the reaction gas mixture of H₂(5%)/C₂H₄(5%)/He was introduced into the reactor. At 60 min after the start of the reaction, H₂ gas was removed from the reaction gas [i.e., C₂H₄(5%)/He] and then introduced again after 120 min. The conversion of ethylene was calculated from the following equation:

\[
\text{Conversion of ethylene} = \frac{(X_{\text{in}} - X_{\text{out}})}{X_{\text{in}}} \quad (1)
\]

where \(X_{\text{in}}\) and \(X_{\text{out}}\) are the concentrations of ethylene gas in the inlet and outlet gases, respectively.

**FT-IR measurements**

In situ diffuse reflectance infrared Fourier-transform (DRIFT) spectra were obtained using an ISDR-600 FT-IR spectrometer (JASCO, Japan) equipped with a mercury–cadmium–tellurium (MCT) at a resolution of 4 cm⁻¹ with 64 co-added scans. Diffuse reflectance cell was filled with the sample power and sealed with a KBr window from the top. In this experiment,
the total flow rate of the reaction gas was fixed to 50 mL min\(^{-1}\). The Pt/Al\(_2\)O\(_3\) sample was pretreated in a flow of 5% H\(_2\)/He for 30 min at 473 K and then cooled to room temperature under a He atmosphere. After measurement of the background spectrum at 303 K under a He atmosphere, the DRIFT spectra were obtained under the gas flow conditions of H\(_2\)(5%)/He, H\(_2\)(5%)/C\(_2\)H\(_4\)(5%)/He, and C\(_2\)H\(_4\)(5%)/He.

Inelastic neutron scattering measurements

The inelastic neutron scattering measurements were carried out with the 4SEASONS time-of-flight spectrometer, SIKI, at the MLF, J-PARC, Japan.\(^{32}\) The samples were loaded into thin double-walled cells whose shape was a traditional hollow cylindrical design, but had gas inlet and outlet cocks added to the top and bottom of the cell. The internal space of the inner cylinder had a hole to contain the same atmosphere of the cryostat exchange gas, to reduce the background contribution as much as possible.

The reduced catalyst was tableted, pulverized to a 50–100 mesh and set in the measurement cell. Before the INS experiments, the sample cells were connected to a pretreatment/reaction system, and were vacuumed and heated (up to 473 K); then, gas (5% H\(_2\)/He) was flowed to remove a surface oxide layer on the Pt metal nanoparticles at a sample preparation room on the offsite of the neutron beamline. After cooling the samples to 300 K, the reaction gas (5% H\(_2\)/He, 5% C\(_2\)H\(_4\)/He, 5% H\(_2\) + 5% C\(_2\)H\(_4\)/He) or He gas (flow rate: 50 mL min\(^{-1}\)) was introduced into the cell. After reacting for 1 h, the gas cocks were closed and then the cell was attached to the sample stick of the cryostat for the neutron spectrometer.

The spectrometer has multiple incident energy (\(E_i\)) capability.\(^{33}\) The Fermi chopper frequency was 300 Hz and setting \(E_i\) was 150 meV. Typical energy resolutions were about \(\Delta E \approx 1\) meV at \(\hbar\omega\) of 50–60 meV and \(\sim 4\) meV at 80–100 meV, respectively. Data collection was performed at 123 K, facilitated by the 4SEASONS cryostat and counted for 6–12 h. Data reduction and analysis were carried out using the software package Utsusemi.\(^{34}\) The INS spectra were obtained with integrated \(Q\) in the range of \(2 \leq |Q| \leq 10\) Å\(^{-1}\). Throughout this paper, the error bar in the spectra represents the standard deviation.

DFT calculations

DFT calculations were performed to identify the active hydrogen species and to assign the INS spectra. We employed a periodic slab model of the \(\gamma\)-alumina (110) surface with loading of a Pt rod. The supercell contains an \((\text{Al}_2\text{O}_3)_{16}\) unit as proposed by Pinto et al.\(^{35}\) with six-layer thickness. A two-layer rod-like Pt(111) structure is put on the alumina surface: the bottom layer involves a hexagonal Pt\(_8\) unit and a Pt\(_6\) unit constructs the top layer. The model contains Pt\(_{14}(\text{Al}_2\text{O}_3)_{16}\) in each supercell. We inserted a vacuum space of 20 Å above the surface and optimized all of the cell parameters and the position of ions. Spin-polarized DFT calculations were performed with the PW91 function\(^{36}\) implemented with the projector augmented wavefunction (PAW) method for representing the core electrons.\(^{37,38}\) We used a \(2 \times 2 \times 1\) \(\Gamma\)-centred \(k\)-point mesh. The plane-wave cut-off was set to 600 eV, which is the optimal value to satisfy the energy convergence. On the optimized Pt/Al\(_2\)O\(_3\) structure, hydrogen atoms were adsorbed. We fixed the ionic position of the bottom two layers during the calculations of hydrogen adsorption and vibrations. The DFT calculations were performed using the VASP package.\(^{39,40}\) The spectrum simulation was performed using the oClimax program.\(^{41}\) The atomic charges on H atoms were evaluated according to Bader partitioning.\(^{42}\)

Results

Hydrogenation of ethylene on Pt/Al\(_2\)O\(_3\)

XRD patterns of the prepared 5 wt% Pt/Al\(_2\)O\(_3\) sample and the bare Al\(_2\)O\(_3\) sample revealed that Pt metal species was highly dispersed on the Al\(_2\)O\(_3\) support because of no diffraction peaks assigned to Pt species (Fig. S1†). Pt L\(_3\)-edge XANES spectrum of the Pt/Al\(_2\)O\(_3\) (Fig. S2a†) revealed that the metallic Pt species was formed on Al\(_2\)O\(_3\) after H\(_2\) treatment because the absorption peak (electron transition from Pt 2p to Pt 5d) intensity of Pt/Al\(_2\)O\(_3\) at 11 567.5 eV was similar to that of Pt foil. The particle size of the supported Pt was estimated to be 1–2 nm from the CN (5.7 ± 0.3) of Pt–Pt, which was obtained by the curve fitting analysis of FT-EXAFS spectrum of the Pt/Al\(_2\)O\(_3\) (Fig. S2b, Table S1†). This value was in good accordance with the Pt particle size (1.8 nm) of the Pt/Al\(_2\)O\(_3\) determined by the CO pulse method using BELCAT-B (MicrotracBEL Corp., Japan).

The ethylene hydrogenation reaction was carried out over a 5 wt% Pt/Al\(_2\)O\(_3\) catalyst at 303 K. Fig. 1 shows the time course of the conversion of ethylene with H\(_2\) [(a) 0–60 min, (c) 120–150 min] and without it [(b) 60–120 min] over the Pt/Al\(_2\)O\(_3\) catalyst at 303 K under H\(_2\)(5%)/C\(_2\)H\(_4\)(5%)/He, and (c) under H\(_2\)(5%)/C\(_2\)H\(_4\)(5%)/He.
Al_{2}O_{3} catalyst. Ethylene conversion of >99% was achieved in the presence of H\textsubscript{2} and ethane was formed (Fig. 1a and c). On the other hand, the ethylene conversion was suppressed in the absence of H\textsubscript{2} (Fig. 1b). In the presence of H\textsubscript{2}, the H species, which is generated by the activation of H\textsubscript{2} on the Pt particles, reacted with ethylene to form ethane at 303 K. However, dehydrogenation reaction did not proceed at 303 K because the free energy change of the ethylene dehydrogenation is large (141 kJ mol\textsuperscript{-1}).\textsuperscript{43} 

Fig. 2 shows the in situ DRIFT spectra of the Pt/Al\textsubscript{2}O\textsubscript{3} sample under several gas conditions. The addition of H\textsubscript{2} increased the intensity of the bands at 2041 and 2112 cm\textsuperscript{-1} (Fig. 2a), which are assigned to on-top hydride and atop H species on the Pt surface, respectively, based on the previous reports.\textsuperscript{29,44-46} We could not detect the n-fold H species because of the selection rule of IR spectroscopy.\textsuperscript{29} The bands in the regions of 1600–1750 and 3100–3700 cm\textsuperscript{-1}, which are attributed to Al-OH species, appeared upon the introduction of H\textsubscript{2}. This phenomenon is explained by the H spillover from the Pt surface to the Al\textsubscript{2}O\textsubscript{3} surface.\textsuperscript{45} In the presence of C\textsubscript{2}H\textsubscript{4} without H\textsubscript{2}, π-CH=CH\textsubscript{2} (1200 and 1490 cm\textsuperscript{-1}) and ethylidyne (1339 and 2883 cm\textsuperscript{-1}) on the Pt surface were detected (Fig. 2b) along with the absorption bands of gaseous C\textsubscript{2}H\textsubscript{4} at 1444, 1889, 2989, 3086 and 3131 cm\textsuperscript{-1}.\textsuperscript{47-49} In addition, the formation of Al-OH species was also observed in the absence of H\textsubscript{2} gas. Since the reactivity of π-CH=CH\textsubscript{2} is higher than that of ethylidyne in the C\textsubscript{2}H\textsubscript{4} hydrogenation reaction,\textsuperscript{50} the Al-OH species would be formed by the migration of H species, which were generated in the ethylidyne formation process on the Pt surface as shown in reaction (2).

\[
\text{Pt} + \text{CH}_{2}=\text{CH}_{2} \rightarrow \text{CH}_{3}\text{C-Pt (ethylidyne)} + \text{H}_{\text{ads}} \tag{2}
\]

The H species on the Pt surface were not detected in the region of 2000–2150 cm\textsuperscript{-1} under the C\textsubscript{2}H\textsubscript{4} atmosphere without H\textsubscript{2} (Fig. 2b) because reaction (2) is suppressed by the saturation of ethylidyne species on the proper Pt sites.\textsuperscript{51} In the presence of both C\textsubscript{2}H\textsubscript{4} and H\textsubscript{2} (Fig. 2c), gaseous product C\textsubscript{2}H\textsubscript{6} was detected at 1400–1550, 2775 and 2840–3100 cm\textsuperscript{-1} overlapping with the small amount of unreacted C\textsubscript{2}H\textsubscript{4}.\textsuperscript{47,52} The signal intensities of hydride (2046 cm\textsuperscript{-1}), H species (2118 cm\textsuperscript{-1}) and π-CH=CH\textsubscript{2} (1200 and 1490 cm\textsuperscript{-1}) on the Pt surface in Fig. 2c were weakened compared to those in Fig. 2a and b because these species were consumed in the C\textsubscript{2}H\textsubscript{4} hydrogenation reaction, which is in good accordance with previous works.\textsuperscript{29,44,48}

Observation of adsorbed hydrogen species on Pt/Al\textsubscript{2}O\textsubscript{3} by INS

Identification of the H species that are silent for FT-IR and investigation of their reactivity are crucial to understand the ethylene hydrogenation catalysis.\textsuperscript{29} To achieve this goal, INS measurements were performed at the BL01 beamline “4SEASONS” of J-PARC. To determine the motions and positions of atoms, INS is the unique technique, in particular for light atoms, e.g. H, Li, and so on. Fig. 3A shows the typical powder averaged color contour map of INS intensity of the 5 wt% Pt/Al\textsubscript{2}O\textsubscript{3} sample treated with H\textsubscript{2} as functions of momentum transfer vector \(Q\) (space correlation information) and energy transfer \(\hbar \omega\) (time correlation information) measured at 123 K with \(E_{i} = 150\) meV. In powder sample, only the information of radial direction (momentum transfer vector \(Q\)) is available and crystal orientations are sacrificed by the integration of powder averaged. Due to the nature of
incoherent scattering of hydrogen, INS spectrum can be integrated over the $Q$ range further, in this case 2–10 Å$^{-1}$ (red spectrum in Fig. 3A), to obtain the better statistics of the INS intensities.

Fig. 3B shows the INS spectra of Pt/Al$_2$O$_3$ measured at several conditions. In the spectrum for the Pt/Al$_2$O$_3$ sample without H$_2$ (Fig. 3Bd), a peak was observed in the range of 100–120 meV, which is attributed to surface OH groups on the support (discussed later). The intensity increased in the wide range of 50–120 meV due to the presence of H$_2$ as shown in Fig. 3Ba. These signals were not detected for the bare Al$_2$O$_3$ support treated with H$_2$ (Fig. S3†). Therefore, the signals in 50–120 meV would be assigned to the hydrogen species produced by the Pt metal catalyst and adsorbed on the Pt surface, the Al$_2$O$_3$ support, and the interface between them. These signals in the region of 50–120 meV were reduced by the addition of C$_2$H$_4$ (Fig. 3Bc). This drastic change is due to the reaction of C$_2$H$_4$ with the H species on the Pt, the Al$_2$O$_3$ surfaces, and/or the interface between them. The signal intensities were also increased in the C$_2$H$_4$ atmosphere without H$_2$, as shown in Fig. 3Bb, indicating that the H species are formed on the Pt/Al$_2$O$_3$ sample according to reaction (2). The signal intensities of H species at 80–120 meV were as high as those for the Pt/Al$_2$O$_3$ sample in the H$_2$ atmosphere, whereas the intensities at 60–80 meV were similar to those obtained after the reaction of C$_2$H$_4$ + H$_2$ in Fig. 3Bc.

**DFT calculation of activated H species on Pt/Al$_2$O$_3$ and simulation of INS spectrum**

DFT calculations were employed to determine feasible adsorption structures of single H atom on the Pt/Al$_2$O$_3$ catalyst using the supercell of Pt$_{14}$(Al$_2$O$_3$)$_{16}$ as a model of the Pt/Al$_2$O$_3$ surface structure. We performed the geometry optimization using probable initial structures that involves atop and bridge sites on Al$_2$O$_3$, atop, bridge, and threefold sites on Pt, and Pt/Al$_2$O$_3$ perimeter sites. The representative adsorption structures obtained by geometry optimization are shown in Fig. 4a and Table S2† and categorized as follows. Five binding structures were found on the Al$_2$O$_3$ surface: AlO–H (1–3) and bridged Al–H–Al (4,5), whereas three structures were observed on the Pt$_{14}$ moiety: bridged Pt–H–Pt at the terrace (6), edge (7) and perimeter (8) sites. We also found threefold Pt$_3$–H structures at the face-centered cubic (fcc)-like hollow site Pt$_3$–H$_{fcc}$ (9), where a Pt atom exists directly below the site, and the hexagonal close-packed (hcp)-like hollow site Pt$_3$–H$_{hcp}$ (10) without a Pt atom below the site. The H atoms of AlO–H (1–3) and Al–H–Al (4,5) were assigned to proton and hydride, respectively (Table S3†). Atomic (neutral) hydrogen was found on Pt–H–Pt (6,7), Pt$_3$–H$_{fcc}$ (9) and Pt$_3$–H$_{hcp}$ (10), whereas the perimeter Pt–H–Pt (8) involved a slightly negative charge, indicating a hydride-like character (Table S3†). The binding structures similar to terrace Pt–H–Pt (6), edge Pt–H–Pt (7) and threefold Pt$_3$–H$_{fcc}$

![Fig. 4](image-url)  
(a) Optimized structure of ten H atoms on the supercell of Pt$_{14}$(Al$_2$O$_3$)$_{16}$. (H: green, Pt: grey, Al: light blue, O: red). The adsorption sites of H atoms are shown as follows: (1) AlO–H, (2) AlO–H, (3) AlO–H, (4) bridge Al–H–Al, (5) bridge Al–H–Al, (6) terrace Pt–H–Pt, (7) edge Pt–H–Pt, (8) perimeter Pt–H–Pt, (9) threefold Pt$_3$–H$_{fcc}$ and (10) threefold Pt$_3$–H$_{hcp}$. (b) Simulated INS spectrum of ten H atoms on the supercell of Pt$_{14}$(Al$_2$O$_3$)$_{16}$. (c) Assignment of the experimental INS spectra.
species, as shown in Fig. 4c. In addition, we successfully identified that the signals appearing at 100–130 meV are mainly assigned to ALO–H (1–3) and Al–H–Al (4,5) species, although these signals were not mentioned in the previous study.5

Discussion

Activated surface species of H2 and C2H4 on Pt/Al2O3 catalyst

We summarized the activated H species on Pt/Al2O3 obtained by INS and FT-IR spectroscopies in Scheme 1a. We demonstrated the formation of surface H species [edge and terrace Pt–H–Pt (6,7), Pt–H–Hfcc (9), and Pt–H–hcp (10)], which has been predicted by DFT calculations using model Pt structures,53–57 on the Pt/Al2O3 surface in the presence of H2 by the INS and DFT (Fig. 3B and 4c) in this study. In addition, the perimeter Pt–H–Pt (6) at the interface between Pt particles and Al2O3 support was also detected in H2 conditions.

We found some ALO–H (1–3) and Al–H–Al (4,5) species on the Pt/Al2O3 sample even in the He condition, in contrast to the bare Al2O3 in the presence of H2, as shown in Fig. S3†. The introduction of H2 during the pretreatment should generate the additional ALO–H (1–3) and Al–H–Al (4,5) species (Fig. 3Ba). It is reported that H species travel a short distance from the Pt surface to the Al2O3 surface.5 This H spillover from Pt particles gives the ALO–H (1–3) and Al–H–Al (4,5) species (Scheme 1a), which is consistent with the fact that the AlO–H (1–3) and Al–H–Al (4,5) species were not detected on the bare Al2O3 support, even in the presence of H2 (Fig. S3†).

The C2H4 provided not only adsorbed π-CH2=CH2 and ethyldiene, but also ALO–H and Al–H–Al species on the Pt/Al2O3 catalyst, which is evidenced by the FT-IR and INS studies as shown in Scheme S1†. In addition, the edge and/or terrace Pt–H–Pt (6 and/or 7) species appearing at 50–55 meV in Fig. 3Bb were detected despite the absence of H2. Tan

Scheme 1 (a) Activated H species on Pt/Al2O3 in the presence of H2, (b) Reaction mechanism of active H species on Pt/Al2O3 in C2H4 hydrogenation.
et al. reported that the H adsorption on a bridged terrace site is unstable and requires high H₂ coverage on the Pt surface.⁵⁴ On the other hand, the edge site is strongest adsorption site in the ridged Pt surface.⁵⁵ Furthermore, the H adsorption on the edge site is favourable in the model of H-covered Pt₁₂₇/(100)MgO.⁵³ Therefore, the Pt–H species formed at 50–55 meV is edge Pt–H (Scheme S1). This means that the H species generated through the formation of ethylidyne are strongly adsorbed on Al₂O₃, which remained even in the He reaction conditions (Scheme 1b), whereas the H species stored on both the Pt and Al₂O₃ surfaces.

Active H species on Pt/Al₂O₃ for C₂H₄ hydrogenation reaction

The perimeter Pt–H–Pt (8), Pt₃–H₆Cc (9), and hydride and atop Pt–H species appearing in the presence of H₂ were drastically reduced by the addition of C₂H₄ with the decrease in the peak intensities of π-CH₂=CH₂ (1200 and 1490 cm⁻¹), which is the active species in the C₂H₄ hydrogenation reaction.⁵⁰ These results indicate that the perimeter Pt–H–Pt (8), Pt₃–H₆Cc (9), and hydride and atop Pt–H species can react with π-CH₂=CH₂ [Langmuir–Hinshelwood (L–H) reaction mechanism] and gaseous C₂H₄ [Eley–Rideal (E–R) reaction mechanism]⁵⁸ to form C₂H₆ (Scheme 1b). In addition, the signal intensity at 55–60 meV also decreased upon the introduction of C₂H₄ (Fig. 3Ba and c). This signal is attributed to terrace Pt–H–Pt (6), which is unstable on the Pt surface.⁵⁴,⁵⁵ Thus, the terrace Pt–H–Pt (6) is also an active species in the C₂H₄ hydrogenation reaction (Scheme 1b). On the other hand, the edge Pt–H–Pt (7) was still observed after the C₂H₄ hydrogenation reaction (Fig. 3Bc). The edge Pt–H–Pt (7) is the most stable species on the ridge Pt surface.⁵⁵ The strong interaction between H and Pt on the edge site inhibits the reaction of edge Pt–H–Pt (7) with C₂H₄.

The additionally formed AlO–H (1–3) and Al–H–Al (4,5) in the presence of H₂ or C₂H₄ were not observed after the reaction of C₂H₄ and H₂ although these AlO–H (1–3) and Al–H–Al (4,5) could not react with C₂H₄ directly (Fig. 3B). Thus, the H species on Al₂O₃ provided by H₂ and/or C₂H₄ can re-migrate to the Pt particle to react with C₂H₄ under the reaction conditions (Scheme 1b), whereas the H species strongly adsorbed on Al₂O₃, which remained even in the He condition (Fig. S3), cannot re-migrate. It was reported that the metal–organic frameworks⁵⁹ and carbons⁶⁰ acted as a hydrogen storage material via the H spillover supported Pt and Pd particles, as revealed by H₂ adsorption/desorption experiments. In addition, H species were provided from metal to carbon materials through Al₂O₃ in the case of Pd/Al₂O₃-decorated graphene sheets.⁶¹ The present study provides evidence that the Al₂O₃ support itself also acts like a hydrogen store and the formed AlO–H (1–3) and Al–H–Al (4,5) can be used as H sources in the C₂H₄ hydrogenation reaction.

Conclusions

In summary, we demonstrated that the H species such as bridged Pt–H–Pt (edge, terrace, perimeter), three-fold Pt₃–H, AlO–H, and bridged Al–H–Al species on Pt/Al₂O₃ catalyst were identified by the combination of INS spectroscopy and DFT calculations for the first time. The reactivity of the H species was successfully observed in situ INS and FT-IR. The atop and hydride Pt–H, terrace and perimeter Pt–H–Pt (6,8), Pt₃–H₆Cc (9) and Pt₃–H₆cp (10) were active intermediates in C₂H₄ hydrogenation to produce C₂H₆ via L–H and/or E–R reaction mechanisms whereas the edge Pt–H–Pt (7) was inert species in the C₂H₄ hydrogenation. In addition, we obtained the direct evidence that the Al₂O₃ support itself acted as a hydrogen storage material and the formed AlO–H (1–3) and Al–H–Al (4,5) species can be used as H sources in the C₂H₄ hydrogenation reaction. This hydrogen storage property should also contribute to the catalytic activity. The results obtained in this study provide the significant insights for the elucidation of reaction mechanism of the catalytic hydrogenation reactions.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

1. F. Zaera, Phys. Chem. Chem. Phys., 2013, 15, 11988–12003.
2. D. Wang and D. Astruc, Chem. Rev., 2015, 115, 6621–6686.
3. J. A. Delgado, O. Benkirane, C. Claver, D. Curulla-Ferré and C. Godard, Dalton Trans., 2017, 46, 12381–12403.
4. F. Meemken and A. Baiker, Chem. Rev., 2017, 117, 11522–11569.
5. F. Zaera, ACS Catal., 2017, 7, 4947–4967.
6. R. Ye, A. V. Zhukovitskiy, C. V. Deraedt, F. D. Toste and G. A. Somorjai, Acc. Chem. Res., 2017, 50, 1894–1901.
7. L. Zhang, M. Zhou, A. Wang and T. Zhang, Chem. Rev., 2020, 120, 683–733.
8. P. Kluson, Appl. Catal., A, 1995, 128, 12–31.
9. J. Harris and S. Andersson, Phys. Rev. Lett., 1985, 55, 1583–1586.
10. G. D. Frey, V. Lavallo, B. Donnadieu, W. W. Schoeller and G. Bertrand, Science, 2007, 316, 439–441.
11. S. Klacar and H. Grönbeck, Catal. Sci. Technol., 2013, 3, 183–190.
12. E. D. German, H. Abir and M. Sheintuch, J. Phys. Chem. C, 2013, 117, 7475–7486.
Catalysis Science & Technology

13 F. Ahmed, M. K. Alam, A. Suzuki, M. Koyama, H. Tsuboi, N. Hatakeyama, A. Endou, H. Takaba, C. A. Del Carpio, M. Kubo and A. Miyamoto, J. Phys. Chem. C, 2009, 113, 15676–15683.
14 R. Prins, Chem. Rev., 2012, 112, 2714–2738.
15 W. Karim, C. Spreafico, A. Kleibert, J. Gobrecht, J. VandeVondele, Y. Ekinci and J. A. van Bokhoven, Nature, 2017, 541, 68.
16 T. Mitsudome, Y. Mikami, M. Matoba, T. Mizugaki, K. Jitsukawa and K. Kaneda, Angew. Chem., Int. Ed., 2012, 51, 136–139.
17 Y. Cao, Z. Sui, Y. Zhu, X. Zhou and D. Chen, ACS Catal., 2017, 7, 7835–7846.
18 P. G. Savva and C. N. Costa, Catal. Rev.: Sci. Eng., 2011, 53, 91–151.
19 V. F. Sears, Neutron News, 1992, 3, 26–37.
20 A. Pawlukoń, I. Natkaniec, E. Grech, J. Baran, Z. Malarski and L. Sobczyk, Spectrochim. Acta, Part A, 1998, 54, 439–448.
21 P. W. Albers, G. Prescher, K. Seibold and S. F. Parker, Chem. Eng. Technol., 1999, 22, 135–137.
22 M. Kofu, N. Hashimoto, H. Akiba, H. Kobayashi, H. Kitagawa, K. Iida, M. Nakamura and O. Yamamura, Phys. Rev. B, 2017, 96, 054304.
23 P. W. Albers, E. Auer, K. Ruth and S. F. Parker, J. Catal., 2000, 196, 174–179.
24 P. W. Albers, M. Poniatowski, S. F. Parker and D. K. Ross, J. Phys.: Condens. Matter, 2000, 12, 4451–4463.
25 P. W. Albers, J. Pietsch, J. Krauter and S. F. Parker, Phys. Chem. Chem. Phys., 2003, 5, 1941–1949.
26 P. W. Albers, M. Lopez, G. Sextl, G. Jeske and S. Parker, J. Catal., 2004, 223, 44–53.
27 M. Carosso, A. Lazzarini, A. Piovano, R. Pellegrini, S. Morandi, M. Manzoli, J. G. Vitillo, M. Jimenez Ruiz, C. Lamberti and E. Grop-po, Faraday Discuss., 2018, 208, 227–242.
28 R. Juárez, S. F. Parker, P. Concepción, A. Corma and H. García, Chem. Sci., 2010, 1, 731–738.
29 M. Carosso, E. Vettero, A. Lazzarini, S. Morandi, M. Manzoli, K. A. Lomachenko, M. J. Ruiz, R. Pellegrini, C. Lamberti, A. Pio-vano and E. Groppo, ACS Catal., 2019, 9, 7124–7136.
30 H. Asakura, S. Yamazoe, T. Misumi, A. Fujita, T. Tsukuda and T. Tanaka, Radiat. Phys. Chem., 2020, 175, 108270.
31 A. L. Ankudinov, B. Ravel, J. J. Rehr and S. D. Conradson, Phys. Rev. B: Condens. Matter Mater. Phys., 1998, 58, 7565.
32 R. Kajimoto, M. Nakamura, Y. Inamura, F. Mizuno, K. Nakajima, S. O. Kawamura, T. Yokoo, T. Nakatani, R. Maruyama, K. Soyama, K. Shibata, K. Suzuki, S. Sato, K. Aizawa, M. Arai, S. Wakimoto, M. Ishikado, S. Shamoto, M. Fujita, H. Hiraka, K. Ohoyama, K. Yamada and C. H. Lee, J. Phys. Soc. Jpn., 2011, 80, SB025.
33 M. Nakamura, R. Kajimoto, Y. Inamura, F. Mizuno, M. Fujita, T. Yokoo and M. Arai, J. Phys. Soc. Jpn., 2009, 78, 093002.
34 Y. Inamura, T. Nakatani, J. Suzuki and T. Otomo, J. Phys. Soc. Jpn., 2013, 82, SA031.
35 H. P. Pinto, R. M. Nieminen and S. D. Elliott, Phys. Rev. B: Condens. Matter Mater. Phys., 2004, 70, 125402.
36 J. P. Perdew, J. A. Chevary, S. H. Vosko, K. A. Jackson, M. R. Pederson, D. J. Singh and C. Fiolhais, Phys. Rev. B: Condens. Matter Mater. Phys., 1992, 46, 6671–6687.
37 P. E. Blöchl, Phys. Rev. B: Condens. Matter Mater. Phys., 1994, 50, 17953–17979.
38 G. Kresse and D. Joubert, Phys. Rev. B: Condens. Matter Mater. Phys., 1999, 59, 1758–1775.
39 G. Kresse and J. Furthmüller, Phys. Rev. B: Condens. Matter Mater. Phys., 1996, 54, 11169–11186.
40 G. Kresse and J. Furthmüller, Comput. Mater. Sci., 1996, 6, 15–50.
41 K. Ramić, C. Wendorff, Y. Cheng, A. I. Kolesnikov, D. L. Aber-nathy, L. Daemen, G. Arbanas, L. Leal, Y. Danon and L. Liu, Ann. Nucl. Energy, 2018, 120, 778–787.
42 G. Henkelman, A. Arnaldsson and H. Jónsson, Comput. Mater. Sci., 2006, 36, 354–360.
43 F. Cataldo, J. Photochem. Photobiol., A, 1996, 99, 75–81.
44 Y. Soma, J. Catal., 1979, 59, 239–247.
45 M. Haneda, T. Watanabe and M. Ozawa, J. Jpn. Pet. Inst., 2012, 53, 191–196.
46 Y. Dong, G. Hu, X. Hu, G. Xie, J. Lu and M. Luo, J. Phys. Chem. C, 2013, 117, 12537–12543.
47 G. B. Lebrón and T. L. Tan, Int. J. Spectrosc., 2012, 2012, 474639.
48 M. K. Ko and H. Frei, J. Phys. Chem. B, 2004, 108, 1805–1808.
49 W. Wasylenko and H. Frei, J. Phys. Chem. B, 2005, 109, 16873–16878.
50 T. Miura, H. Kobayashi and K. Domen, J. Phys. Chem. B, 2000, 104, 6809–6814.
51 J. Kubota, S. Ichihara, J. N. Kondo, K. Domen and C. Hirose, Langmuir, 1996, 12, 1926–1927.
52 FT-IR data of ethane, NIST Chemistry WebBook, Standard Reference Database 69.
53 S. M. Kozlov, H. A. Aleksandrov and K. M. Neyman, J. Phys. Chem. C, 2015, 119, 5180–5186.
54 T. L. Tan, L. Wang, D. D. Johnson and K. Bai, J. Phys. Chem. C, 2013, 117, 22696–22704.
55 S. Gudmundsdóttir, E. Skúlason, K.-J. Weststrate, L. Juulink and H. Jönsson, Phys. Chem. Chem. Phys., 2013, 15, 6323–6332.
56 L. Yan, Y. Sun, Y. Yamamoto, S. Kasamatsu, I. Hamada and O. Sugino, J. Chem. Phys., 2018, 149, 164702.
57 N. B. Arboleda Jr, H. Kasai, W. A. Diño and H. Nakashima, Jpn. J. Appl. Phys., 2007, 46, 4233–4237.
58 Y. Dong, M. Ebrahimi, A. Tillekaratne and F. Zaera, J. Phys. Chem. Lett., 2016, 7, 2439–2443.
59 Y. Li and R. T. Yang, J. Am. Chem. Soc., 2006, 128, 8136–8137.
60 C.-S. Tsao, Y.-R. Tseng, C.-Y. Wang, H.-H. Tseng, T.-Y. Chung, H.-C. Wu, T. Yamamoto, K. Kaneko and S.-H. Chen, J. Phys. Chem. Lett., 2010, 1, 1060–1063.
61 Z. G. Bajestani, A. Yürüm and Y. Yürüm, Int. J. Hydrogen Energy, 2016, 41, 9810–9818.