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Selective Dimerization of Ethene to 2-Butene on Zn²⁺-Modified ZSM-5 Zeolite

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KEYWORDS. Ethene dimerization, 2-butene, Zn-modified zeolites, ZSM-5, solid-state NMR, FTIR spectroscopy, DFT.

ABSTRACT. Selective dimerization of ethene to 2-butene on Zn²⁺-containing ZSM-5 zeolite (Zn²⁺/ZSM-5) at 296–523 K has been discovered. The intermediate but-3-en-1-ylzinc species is identified with ¹³C CP/MAS NMR and FTIR spectroscopy. DFT study of two alternative dimerization pathways reveals that the intermediate is formed with the involvement of the saturated bridged dimeric Zn–(CH₂)₄–O species. It is also shown that ethene conversion to 2-butene increases with the increase in the quantity of Brønsted acid sites in Zn²⁺/ZSM-5 zeolite, however, the selectivity of the reaction decreases. The results obtained are of potential interest for

developing industrially relevant Zn-containing zeolite catalysts for the selective conversion of ethene to 2-butene.

1. INTRODUCTION

The modern chemical industry requires the synthesis of significant amounts of linear butenes to be widely used in manufacturing polymers and other valuable chemicals (plasticizers, antioxidants, oils, etc.).¹ One of the options for the synthesis of 1- and 2-butene is the dimerization of ethene,¹⁻³ which is an abundant raw material. Several catalysts based on different transition metals (Co, Ni, Pd, Rh, Zr) with or without alkylaluminium co-catalysts have been tested for ethene dimerization.^{1,4} Among the catalysts used, Ni-based systems dominate this field both in homogeneous and heterogeneous catalysis.²⁻⁴ Since nickel complexes are often stable only at low temperatures, heterogeneous catalysts such as nickel supported on silica, silica-alumina, zeolites, and mesoporous materials are preferred.⁵⁻¹¹ Despite the notable development in Ni-modified zeolites synthesis and investigation, some limitations exist concerning the selectivity toward the target product and the catalyst stability. Furthermore, the nature of the active species and the mechanism of dimerization are still under discussion.^{8, 11-12} Moreover, Ni-containing catalysts provide quite selective 1-butene formation, whereas 2-butene is also demanded olefin for the polymer industry. In this regard, it is of particular interest to search for alternative catalytic systems capable of converting ethene to 2-butene.

Zn-containing zeolites provide the strong interaction of Zn^{2+} cations with $\text{C}_3\text{--C}_4$ alkenes giving very stable π -complexes.¹³⁻¹⁵ This interaction facilitates di- and trimerization of the alkenes with the involvement of Zn^{2+} cations and prevents alkene oligomerization on Brønsted acid sites (BAS). Therefore, it is appealing to test Zn-modified zeolite for ethene dimerization.

In this work, Zn²⁺-modified ZSM-5 zeolite is investigated with respect to ethene transformation. We demonstrate for the first time the potential of Zn-loaded zeolites to perform remarkably selective ethene dimerization to 2-butene. To gain insight into the reaction mechanism, a combination of experimental and computational techniques was applied. Solid-state NMR spectroscopy (¹H, ¹³C MAS NMR), was used to follow the pathway of ethene transformation with the assistance of Zn²⁺ sites and detect the reaction intermediate. To further clarify the structure of the intermediate, FTIR spectroscopy was used. The alternative pathways of the detected intermediate formation were examined with DFT calculations.

2. EXPERIMENTAL SECTION

2.1. Materials and Reagents. Ethene-¹³C₁ (99% ¹³C isotope enrichment), benzene-¹³C₆ (99% ¹³C isotope enrichment), ethene, and metallic zinc were purchased from Aldrich Chemical Co. Inc. and were used without further purification. Industrially produced n-butene (thermodynamic mixture, trans:cis:1-butene = 100:26:4)¹⁶ was used after purification from possible traces of water vapor or other admixtures via freezing and thawing in the liquid nitrogen trap.

2.2. Zeolite Samples Preparation. The parent H-ZSM-5 zeolite, Si/Al = 12, was kindly provided by Tricat Zeolites. Before use, the zeolite powder was calcined in airflow at 773 K. The detailed characterization of this zeolite including XRD, low-temperature nitrogen adsorption, ²⁹Si, ¹H, and ²⁷Al MAS NMR data was reported in our earlier work.¹⁵ Table 1 demonstrates the main characteristics of the parent and Zn-modified ZSM-5 zeolite samples.

Table 1. The Properties of the Parent and Zn-Modified Zeolite Samples.

Zeolite sample	Si/Al	Zn / wt% (calculated)	Si–O(H)–Al / $\mu\text{mol g}^{-1}$	Zeolite unit cell
H-ZSM-5	12	–	1290	$\text{Al}_{0.5}^{\text{oct}} \text{H}_{7.4} \text{Al}_{7.4} \text{Si}_{88.6} \text{O}_{192}$
Zn^{2+} /ZSM-5	12	3.6	150	$\text{Zn}^{2+}_{3.3} \text{Al}_{0.5}^{\text{oct}} \text{H}_{0.9} \text{Al}_{7.4} \text{Si}_{88.6} \text{O}_{192}$
Zn^{2+} /H-ZSM-5	12	0.9	1010	$\text{Zn}^{2+}_{0.8} \text{Al}_{0.5}^{\text{oct}} \text{H}_{5.9} \text{Al}_{7.4} \text{Si}_{88.6} \text{O}_{192}$

To introduce isolated Zn^{2+} cations, the method of H-ZSM-5 reaction with Zn vapor was applied.¹⁷⁻²⁰ Before the reaction, the parent zeolite was activated via evacuation at 673 K for 16 h (residual pressure $< 10^{-2}$ Pa). After the activation, the reaction between H-ZSM-5 and zinc vapor was carried out in a vacuumed glassware at 773 K for 2 h followed by evacuation at 773 K for 20 h to remove molecular hydrogen and unreacted zinc. To study the particular effects of Zn sites and BAS on ethene transformation, two samples of Zn-modified ZSM-5 zeolites were prepared with different degrees of BAS exchange for Zn^{2+} cations. To synthesize ZSM-5 zeolite with minimum BAS content, the excess of metallic zinc (the ratio Zn/Al = 2.8) was used. This sample was designated as Zn^{2+} /ZSM-5. To obtain a zeolite sample with appreciable amounts of both BAS and Zn^{2+} sites, the ratio Zn/Al was chosen to be 0.13. This sample was designated as Zn^{2+} /H-ZSM-5.

The concentration of residual BAS (Si–O(H)–Al groups) in both samples was determined with ^1H MAS NMR approach²¹ using benzene- $^{13}\text{C}_6$ as an internal standard. The known amount of benzene- $^{13}\text{C}_6$ was loaded into the ampoules with Zn^{2+} -modified ZSM-5 samples (the pressure was controlled with vacuum gauge, Thermovac 101, Leybold, Germany). Afterward, the ampoules were sealed with a torch flame and heated at 373 K for 46 h to achieve benzene uniform distribution

over the zeolite pore volume. ^1H MAS NMR spectra of the zeolite samples with adsorbed benzene- $^{13}\text{C}_6$ are presented in Figure S1. It was obtained that the concentration of BAS amounted to $150\ \mu\text{mol g}^{-1}$ in $\text{Zn}^{2+}/\text{ZSM-5}$ and $1010\ \mu\text{mol g}^{-1}$ in $\text{Zn}^{2+}/\text{H-ZSM-5}$ (Table S1), giving the degree of BAS exchange for zinc cations 88 % and 22 %, respectively. It was decided to use ^{13}C -labelled benzene to be able to record high-quality ^{13}C CP/MAS NMR spectra of the samples and ensure that benzene did not react under these conditions and was an appropriate internal standard for OH concentration measurements. Indeed, the related spectra (not shown) contain only the signal from benzene at 132 ppm.

2.3. Samples Preparation for MAS NMR Experiments. To prepare the samples for MAS NMR measurements, the powder of activated zeolite sample, $\text{Zn}^{2+}/\text{ZSM-5}$ or $\text{Zn}^{2+}/\text{H-ZSM-5}$, was placed into a highly symmetrical ampoule, 3.5 mm in diameter, made of Pyrex glass. Such ampoules fit tightly into a 4 mm MAS NMR rotor and can be spun at high speed to obtain high-resolution solid-state NMR spectra. The adsorption of ethene- $^{13}\text{C}_1$ was conducted as follows. First, the required amount of the reagent was loaded to the calibrated volume (the pressure was controlled with a vacuum gauge, Thermovac 101, Leybold, Germany). Second, ethene was transferred to the zeolite sample by the ampoule freezing with liquid N_2 . Eventually, the ampoule with ethene adsorbed on Zn-containing zeolite sample was sealed with a torch flame and kept in liquid N_2 to prevent any transformation of the alkene. The total concentration of ethene amounted to $316\ \mu\text{mol}$ per gram of the Zn-modified zeolite.

2.4. MAS NMR Measurements. MAS NMR spectra were recorded at 296 K on a Bruker Avance-400 spectrometer (9.4 T), equipped with a broad-band double-resonance 4 mm MAS NMR probe. ^1H MAS NMR spectra (the resonance frequency of 400.13 MHz) were recorded with a 60 s delay between the scans and a spinning rate of 7 kHz, 16 scans were accumulated for one spectrum. To

obtain ^{13}C (CP) MAS NMR spectra at the resonance frequency of 100.63 MHz, the NMR rotors with the glass ampoules were spun at 9 kHz with dry compressed air. To suppress spin-spin coupling, high-power proton decoupling was applied at the power level corresponding to 5.0 μs length of 90° ^1H pulse and nutation frequency of 50 kHz. For each ^{13}C MAS NMR spectrum, 2000 scans were acquired (5 s repetition delay) with a 5.4 μs 90° excitation pulse. ^{13}C CP/MAS NMR spectra (20000–70000 scans with a 2 s delay) were recorded with the contact time of 2 ms at the Hartmann-Hahn matching condition of 50 kHz. Tetramethylsilane (TMS) was used as an external standard to determine ^1H and ^{13}C chemical shifts with an accuracy of ± 0.1 ppm. The parameters of ^{13}C CP/MAS NMR were optimized using adamantane as the standard.

2.5. FTIR Spectroscopy. Ethene transformation on a $\text{Zn}^{2+}/\text{ZSM-5}$ sample was additionally investigated with FTIR spectroscopy. First, the parent H-ZSM-5 zeolite was pressed in a self-supporting wafer, 12.3 mg cm^{-2} in density, and placed to the homemade quartz holder. The zeolite sample was activated via gradual heating to 673 K (1 K min^{-1}) under vacuum and further evacuation at 673 K for 16 h (residual pressure $< 10^{-2}$ Pa). To introduce isolated Zn^{2+} cations, the excess of metallic zinc ($\text{Zn}/\text{Al} = 22$) was transferred to the zeolite wafer in a sealed glass system and heated at 773 K for 1 h followed by evacuation at 773 K for 5 h. Afterward, the wafer in the holder was transferred into the homemade IR cell (CaF_2 windows) in a glovebox under Ar. To remove Ar and possible water impurities, the sample was later evacuated at 723 K for 2.5 h. The IR cell was connected to a valve for gaseous reagents dosage and a special compartment for ex situ sample heating.

FTIR spectra were recorded on Shimadzu IRTracer-100 FTIR spectrophotometer equipped with a purge control kit PCK-100. Purging of the sample compartment, detector, and interferometer with Ar flow was carried out before and during the spectra acquisition to minimize the effect of

atmospheric water and CO₂ on the resulting FTIR spectra. The spectral range was chosen to be 1100–4000 cm⁻¹, with the resolution being 4 cm⁻¹. For each spectrum, 500 scans were accumulated. The spectrum of pure Zn²⁺/ZSM-5 zeolite was first recorded. Then, incremental amounts of ethene were adsorbed from the gas phase on the zeolite wafer held in liquid N₂ to achieve a total ethene concentration of 50, 100, 300, and 500 μmol g⁻¹, with the FTIR spectrum being acquired after each dosage. To monitor ethene transformation at 373–623 K, the wafer was transferred to the special compartment, heated at the required temperature, cooled to 296 K, and transferred back to the IR cell followed by the spectrum recording. The same procedure was applied to perform the experiments with n-butene.

2.6. DFT Calculations

2.6.1. Computational Model. Molecular cluster containing 66 T-atoms (Figure S2a) was constructed from the MFI framework structure obtained from the Database of Zeolite Structures.²² Dangling O-atoms were substituted by H atoms at 1.47 Å distance from a neighboring Si-atom.²³ To compensate for the charge of the Zn²⁺ cation, two aluminum atoms were placed at the T7 and T12 positions (Figure S2b).²⁴

2.6.2. Computational Methods. DFT spin-unrestricted calculations were carried out using the ORCA 5.0.1 program package.²⁵⁻²⁶ PBE0 hybrid exchange-correlational functional²⁷ with D3BJ dispersion correction²⁸⁻²⁹ was used to describe the exchange-correlational term. 6-31G* basis set³⁰⁻³² was used for the framework atoms (Si, Al, O, H_{terminal}) while the def2-TZVP³³⁻³⁴ one was used for the atoms of the extra-framework species (Zn, C, H). The geometry of the models was considered optimized when the RMS energy gradient and maximum energy gradient were less

than 5×10^{-6} a.u. and 5×10^{-5} a.u., respectively. During the geometry optimization, the cartesian coordinates of the terminal H-atoms were fixed.

Transition states were identified using the CI-NEB procedure as implemented in ORCA.³⁵ The structure of these states was confirmed by the presence of only one imaginary frequency along the reaction coordinate in the vibrational spectra. These spectra were calculated using the partial hessian vibrational analysis (PHVA). During PHVA, only extra-framework atoms were allowed to relax.

3. RESULTS AND DISCUSSION

Figure 1 shows ^{13}C (CP) MAS NMR spectra of ethene- $^{13}\text{C}_1$ adsorbed on $\text{Zn}^{2+}/\text{ZSM-5}$ zeolite with high content of Zn^{2+} cations (Table 1, Figure S1). At 296 K, mainly the signal at 134 ppm of the adsorbed ethene is detected in the spectra (Figure 1a,b). The significant downfield shift of the ethene signal with respect to that expected in solution (123.5 ppm)³⁶ is accounted for by the C=C bond perturbation due to the formation of the π -complex with Zn^{2+} cations.³⁷ The ^{13}C CP/MAS NMR spectrum (Figure 1b), which emphasizes strongly adsorbed and immobile species, contains additional minor signals at 10, 18, 31, 105, and 180 ppm. The comparison of the ^1H MAS NMR spectrum of ethene on $\text{Zn}^{2+}/\text{ZSM-5}$ with that of pure zeolite (Figure S3) reveals that some additional amount of BAS is formed at ethene adsorption. At 423–473 K (Figure 1c–f), the signals of 2-butene are detected in the spectra, with no other products formed from ethene being observed. The signal at 142 ppm belongs to C=C atoms, while those at 18 and 13 ppm arise from the methyl groups of trans- and cis-2-butene, respectively.^{14, 38} This is unequivocal evidence for the selective ethene dimerization to 2-butene on $\text{Zn}^{2+}/\text{ZSM-5}$ at $T \leq 473$ K.

The formation of 2-butene was observed on Zn²⁺/H-ZSM-5 zeolite with the larger quantity of BAS (Table 1, Figure S1) at 296 K (Figure S4). However, no specific signals at 10, 31, 105, and 180 ppm were detected. Also, the further transformation of 2-butene occurs on Zn²⁺/H-ZSM-5 at T ≥ 423 K as seen by the signals at 26–30, 93, and 115 ppm¹⁴ (Figure S4). Therefore, the larger quantity of BAS increases ethene conversion to 2-butene but decreases the dimerization selectivity at the expense of the oligomerization process at T ≥ 423 K.

The four signals at 10, 31, 105, and 180 ppm, disappearing at T > 296 K, should belong to some intermediate preceding the formation of 2-butene (Figure 1b). Since the generation of additional BAS upon ethene adsorption was detected (vide supra), the formation of the Zn-attached C₄-intermediate can be assumed. Additionally, the observed chemical shifts (105, 180 ppm) indicate the presence of a C=C bond in the intermediate. Therefore, two alternative structures can be proposed (Scheme 1). The chemical shifts exhibited by η¹,η²-allyl species on ZnO³⁹ are in favor of but-2-en-1-ylzinc (**1**) with an internal C=C bond. On the other hand, the structure of the intermediate can be represented by but-3-en-1-ylzinc (**2**), with a terminal C=C bond, based on the chemical shifts of bis(2-methylallyl)zinc⁴⁰ with η¹,η²-like fashion of bonding to Zn. In both cases, the C=C bonds should have π-bonding to some surface site, presumably a neighboring BAS, to explain the particular chemical shifts observed.

$C_2H_4-^{13}C_1 @ Zn^{2+}/ZSM-5$

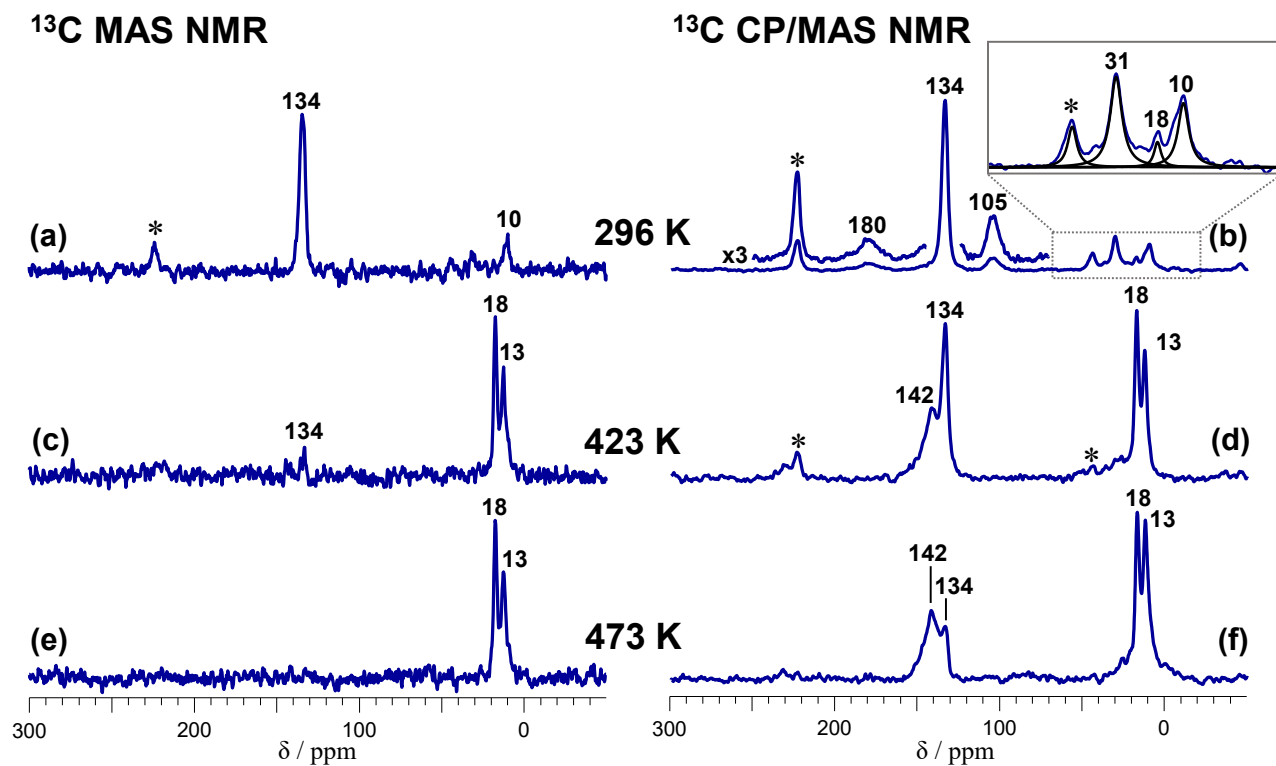
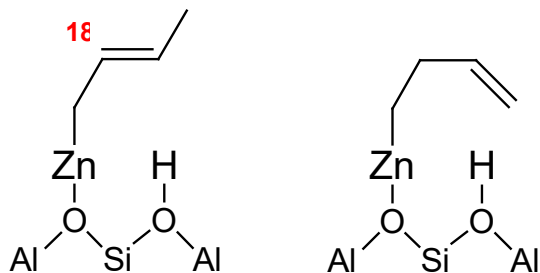


Figure 1. ^{13}C MAS NMR (a,c,e) and ^{13}C CP/MAS NMR (b,d,f) spectra of ethene- $^{13}C_1$ adsorbed on $Zn^{2+}/ZSM-5$ zeolite. The sample was successively heated ex situ for 5 min at 296 K (a,b), 423 K (c,d), and 473 K (e,f). Asterisks (*) denote spinning sidebands.



Scheme 1. The Alternative Structures of the Ethene Dimerization Intermediate Detected with ^{13}C MAS NMR. The Assigned ^{13}C Chemical Shifts are Shown in Red.

To further clarify the structure of the intermediate of ethene dimerization, FTIR spectroscopy was applied (Figures 2, S5). Upon ethene adsorption on $\text{Zn}^{2+}/\text{ZSM-5}$ at 296 K, the characteristic bands of ethene are seen in the region of C–H bending (δ_{CH} , 1332 cm^{-1}) and stretching (ν_{CH} , 3087 and 2989 cm^{-1}) vibrations.⁴¹ The frequency of ethene C=C stretching vibration is expectedly redshifted to 1586 cm^{-1} (compared to 1623 cm^{-1} for gaseous ethene⁴¹⁻⁴² and 1612 cm^{-1} for ethene adsorbed on zeolite BAS)⁴³ due to the formation of a π -complex with Zn^{2+} sites.^{37, 42} The strong interaction of ethene with Zn^{2+} sites is also indicated by the observation of the symmetry-forbidden^{37, 42} δ_{CH} and $\nu_{\text{C=C}}$ bands. The band at 3611 cm^{-1} corresponding to ν_{OH} of residual zeolite BAS (Si–O(H)–Al groups) decreases while the broad feature appears at around $3300\text{--}3500\text{ cm}^{-1}$ from the perturbed O–H stretching vibration due to the interaction of BAS with the adsorbed hydrocarbon species. The bands at 2954 (ν_{CH}), 1637 ($\nu_{\text{C=C}}$), and centered at 1458 and 1411 cm^{-1} (δ_{CH}) do not belong to adsorbed ethene and can be assigned to either the intermediate or the product of ethene conversion. Heating the sample at 373 K results in the change of the spectrum with a set of bands being almost identical to the one of 2-butene adsorbed on $\text{Zn}^{2+}/\text{ZSM-5}$ (Figures S6, S7). The band at 1637 cm^{-1} , however, disappears entirely which means it belongs, presumably, to the intermediate detected with ^{13}C CP/MAS NMR at 296 K. The position of this band is very similar to the band of 1-butene C=C stretching,⁴⁴ and the bands at 2954 , 1458 , and 1411 cm^{-1} are due to the stretching and bending modes (ν_{CH} , δ_{CH}) of $-\text{CH}_2-$ and $-\text{CH}=\text{CH}_2$ fragments. Therefore, the intermediate indeed represents the C_4 -fragment with the terminal C=C bond. Hence, the FTIR data are in favor of but-3-en-1-ylzinc (**2**) being the intermediate of ethene dimerization.

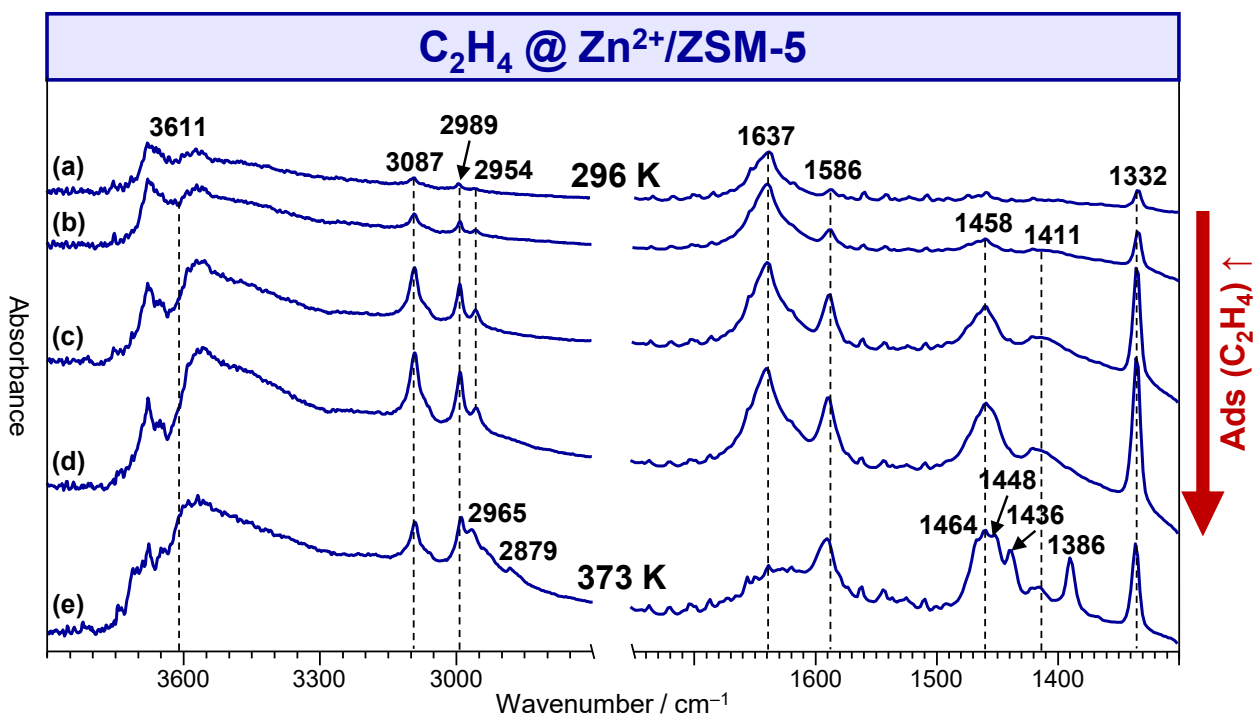
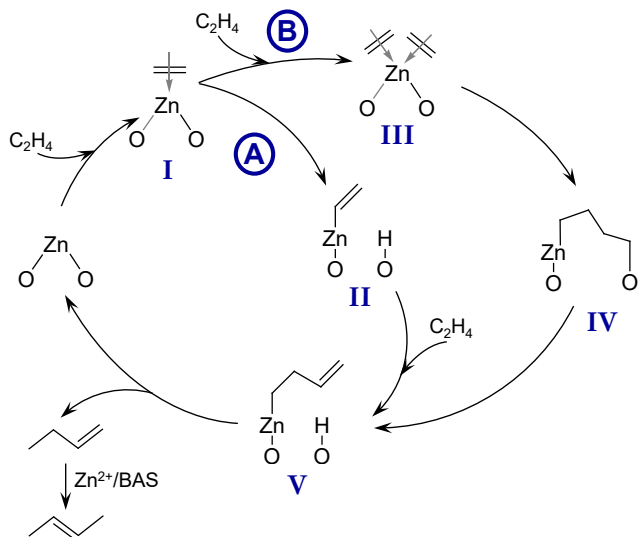


Figure 2. FTIR spectra of ethene adsorbed on Zn²⁺/ZSM-5 zeolite with the spectrum of pure Zn²⁺/ZSM-5 zeolite being subtracted (see Figure S5 for the original spectra). Incremental amounts of ethene were adsorbed on the zeolite sample (a–d) at 296 K. The sample was further heated at 373 K for 5 min (e).

Thus, both MAS NMR and FTIR spectroscopy provide evidence for selective dimerization of ethene to 2-butene on Zn²⁺/ZSM-5 zeolite at 296–473 K. Moreover, the intermediate species, including ethene π -complex with Zn²⁺ site and but-3-en-1-ylzinc, were identified with both methods.



Scheme 2. Possible Pathways of Ethene Dimerization on $\text{Zn}^{2+}/\text{ZSM-5}$ Zeolite: via Vinylzinc Species **II** (A) and via the Formation of the Bridged Intermediate **IV** (B).

The formation of ethene π -complex and but-3-en-1-ylzinc implies the occurrence of ethene dimerization on a single Zn^{2+} site. Adapting the mechanisms of ethene dimerization proposed earlier for Ni- and Ga-based systems^{1, 10, 12, 45} to Zn-containing zeolite, it is reasonable to suggest two alternative pathways, **A** and **B**, of but-3-en-1-ylzinc formation (Scheme 2). But-3-en-1-ylzinc (shown as intermediate **V** in Scheme 2) is desorbed from the Zn^{2+} site as 1-butene which is transformed to 2-butene either on the Zn sites, with the involvement of allylic intermediates, or via the olefin protonation-deprotonation on BAS.¹⁴ Obtained spectroscopic data, however, do not allow us to distinguish between pathways **A** and **B**. While **V** is experimentally detected, neither zinc-vinyl species (**II**) nor the bridged structure (**IV**) were observed with NMR and FTIR spectroscopy. This makes it virtually impossible to experimentally determine the mechanism of **V** formation. For this reason, we have carried out a theoretical study of ethene dimerization on $\text{Zn}^{2+}/\text{ZSM-5}$ zeolite with the DFT method to obtain more detailed information about the reaction mechanism.

The calculated energy profiles for pathways **A** and **B** are presented in Figure 3. The energy of ethene adsorption and stabilization in the form of a π -complex with a Zn^{2+} site (**I**) is 114 kJ mol^{-1} . The next step, for pathway **A**, is ethene dissociation on the Zn^{2+} site to give **II** and Si–O(H)–Al group. This reaction is endothermic and has an activation energy of 136 kJ mol^{-1} . The adsorption of the second ethene molecule on the Zn^{2+} site lowers the system energy to -109 kJ mol^{-1} . The following ethene insertion into the Zn–C bond of vinylic species giving **V_A** has a rather high activation barrier of 171 kJ mol^{-1} . Following pathway **B** after **I** formation, the second ethene molecule is adsorbed lowering the system energy by 53 kJ mol^{-1} (**III**). The formation of **IV** is exothermic and kinetically feasible, with the activation energy being equal to 109 kJ mol^{-1} . The next step of **IV** deprotonation to give **V_B** is characterized by the activation barrier of 118 kJ mol^{-1} . Note that although both pathways **A** and **B** provide the formation of but-3-en-1-ylzinc, the species **V_A** and **V_B** are not identical. They are located differently within the zeolite pore system and interact with the hydrogen atoms of structurally different Si–O(H)–Al groups (Figure S8). Both **V_A** and **V_B** can be further transformed to 1-butene π -complex (**VI**) with the Zn^{2+} site, with the activation energy being equal to 30 kJ mol^{-1} and 75 kJ mol^{-1} , respectively. The energy of 1-butene (**VII**) desorption from the Zn^{2+} site is 75 kJ mol^{-1} .

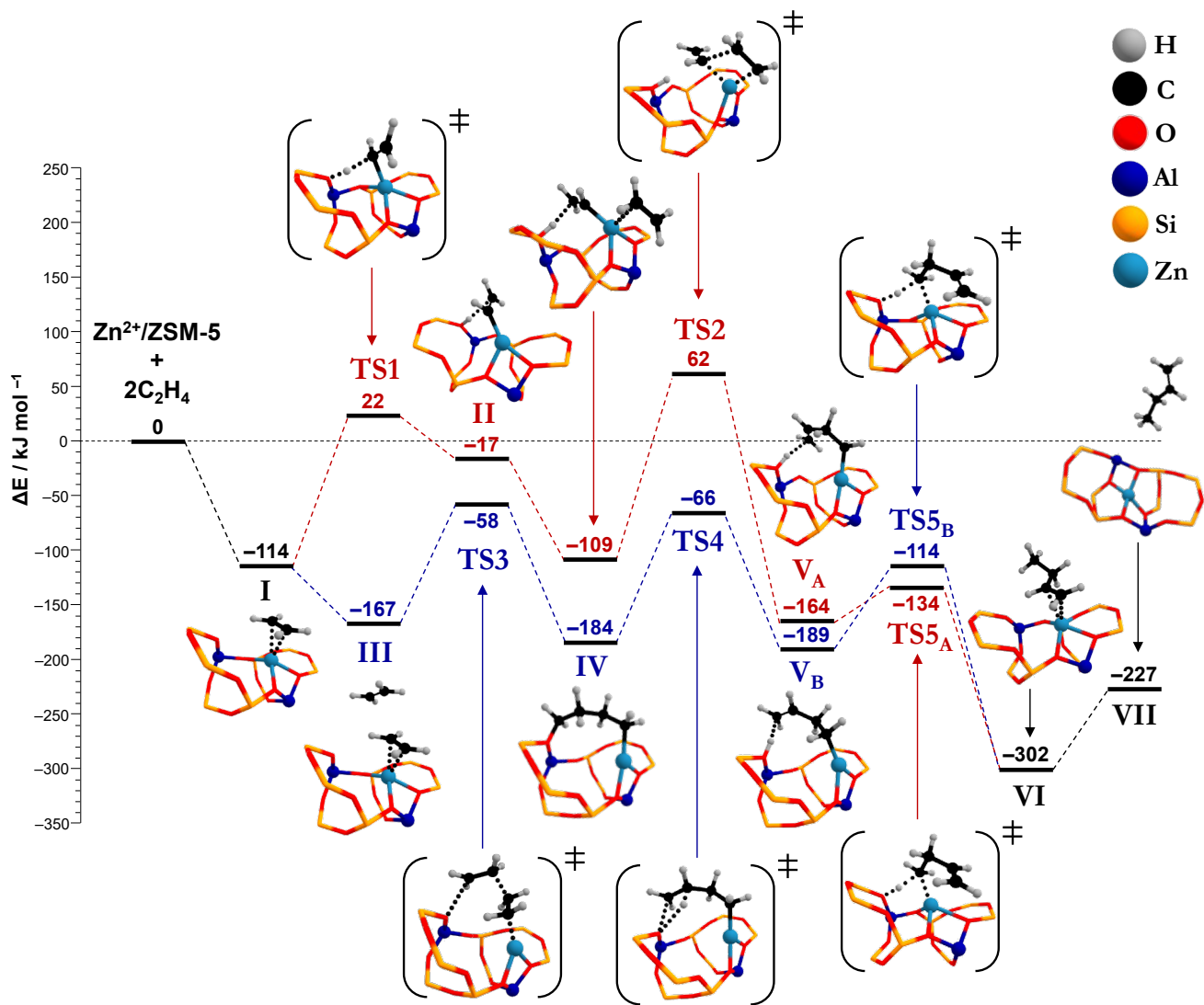


Figure 3. Calculated energy profiles and optimized structures for ethene transformation on Zn²⁺/ZSM-5 by pathways **A** (in red) and **B** (in blue) of Scheme 2. The energies in kJ mol⁻¹ have been referenced to the optimized zeolite cluster and two ethene molecules.

Therefore, theoretical investigation of ethene dimerization on Zn²⁺/ZSM-5 zeolite shows that although both pathways **A** and **B** in Scheme 2 are possible, the pathway **B** involving **IV** is more probable, because it is characterized by lower activation barriers for all steps leading to **V** formation than pathway **A**.

4. CONCLUSION

In summary, remarkably selective dimerization of ethene to 2-butene was experimentally detected on Zn²⁺/ZSM-5 zeolite with high content of Zn²⁺ sites and low quantity of BAS at 296–473 K. The intermediate species, including π -complex of ethene with the zinc sites and but-3-en-1-ylzinc, were identified with ¹³C MAS NMR and FTIR methods. Two alternative pathways of but-3-en-1-ylzinc intermediate formation were verified with DFT calculations. It is inferred that the pathway with the involvement of the saturated bridged dimeric Zn–(CH₂)₄–O species is more probable. It is shown that ethene conversion to 2-butene increases with the increase of the quantity of BAS in Zn²⁺/ZSM-5 zeolite, however, the selectivity of the reaction decreases. The results obtained are of potential interest for developing industrially relevant Zn-containing zeolite catalysts for the selective conversion of ethene to 2-butene.

ASSOCIATED CONTENT

The Supporting Information is available free of charge on ACS Publication website at DOI:

¹H MAS NMR spectra of benzene-¹³C₆ adsorbed on Zn²⁺/ZSM-5 and Zn²⁺/H-ZSM-5 zeolites. Cluster model of MFI-type zeolite and the local geometry of zeolite-confined Zn²⁺ cation used for DFT calculations. ¹H MAS NMR spectra of pure Zn²⁺/ZSM-5 zeolite and ethene-¹³C₁ adsorbed on Zn²⁺/ZSM-5. ¹H and ¹³C (CP) MAS NMR spectra of ethene-¹³C₁ adsorbed on Zn²⁺/H-ZSM-5 zeolite. FTIR spectra of pure Zn²⁺/ZSM-5 zeolite and ethene and n-butene adsorbed on Zn²⁺/ZSM-5. The optimized structures of but-3-en-1-ylzinc.

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