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Facilitating the reduction of V-O bonds on VO_x/ZrO_2 catalysts for non-oxidative propane dehydrogenation†

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Supported vanadium oxide is a promising catalyst in propane dehydrogenation due to its competitive performance and low cost. Nevertheless, it remains a grand challenge to understand the structure–performance correlation due to the structural complexity of VO_x -based catalysts in a reduced state. This paper describes the structure and catalytic properties of the VO_x/ZrO_2 catalyst. When using ZrO_2 as the support, the catalyst shows six times higher turnover frequency (TOF) than using commercial γ -Al₂O₃. Combining H₂-temperature programmed reduction, in situ Raman spectroscopy, X-ray photoelectron spectroscopy and theoretical studies, we find that the interaction between VO_x and ZrO_2 can facilitate the reduction of V–O bonds, including V=O, V–O–V and V–O–Zr. The promoting effect could be attributed to the formation of low coordinated V species in VO_x/ZrO_2 which is more active in C–H activation. Our work provides a new insight into understanding the structure–performance correlation in VO_x -based catalysts for non-oxidative propane dehydrogenation.

Introduction

Propylene is one of the most important chemical building blocks and its high-value derivatives are of great demand in the chemical industry. Due to the large-scale exploitation of shale gas, direct propane dehydrogenation becomes a particularly important method to produce propylene using propane as feedstock. Commercialized propane dehydrogenation plants generally utilize Pt-based and CrO_x-based catalysts, however these two kinds of catalysts are either expensive or toxic.¹ Alternatively, supported vanadium oxide is a promising catalyst compared with Pt and CrO_x for its competitive performance, low cost and low toxicity.²,³

It is well established that VO_x -based catalysts can be utilized in propane oxidative dehydrogenation (ODH). However, there is a trade-off effect between conversion and selectivity associated with over oxidation to CO_x , which prevents the ODH process from achieving high propylene yield. In contrast, better reactivity and selectivity, especially excellent regeneration stability can be achieved when supported VO_x catalysts are used in the non-oxidative propane dehydrogenation (PDH) process. Propagate 1. Nevertheless, the structure-performance

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correlation of VO_x -based catalysts is still unclear due to the structural complexity of the supported VO_x catalytic system. Previous studies have confirmed that several factors could influence the performance of VO_x -based catalysts such as polymerization forms of VO_x , chemical states of V and the identity of the support.¹⁴⁻¹⁸

The support effect is an essential parameter because VO_x can be bonded to a support through the V–O–support interaction. The variety of supports could lead to significant changes in the catalytic properties of VO_x . It is generally accepted that the support could affect reactivity by stabilizing the active sites and/or altering the electron state of active sites. ^{19–22} The support effect has been extensively studied over VO_x -based catalysts in the ODH process. Researchers have concluded that the lattice oxygen in V=O and V–O–support (V–O–S) bonds is consumed in C–H activation and H₂O is formed. Thus, support identities could affect the reaction by tuning the oxygen vacancy formation energy, which is confirmed by experiments and DFT calculations. ^{23–25}

However, the support effect works in an entirely different way in the PDH process because not only active sites but also reaction mechanisms are distinct from those of the ODH process. V–O bonds directly catalyze C–H activation forming $\rm H_2$ rather than $\rm H_2O$. Previous studies found that the reactivity of PDH is dependent on the bond strength of V–O–S.²² It is proposed that V–O–S bonds are active sites. In addition, the support identity also influences the behavior of carbon deposition on $\rm VO_x$ based catalysts, which inversely affects activity and on-stream stability.^{12,18} Nevertheless, the effect of a support

is still not clear because the structure of VO_x and the VO_x -support interface in a reduced state is still ambiguous.

Herein, we explore the support effect on the catalytic performance of VO_r-based catalysts for PDH and provide a new insight into understanding how support identity matters. We discovered that VO_x/ZrO₂, a well-known catalyst for ODH, ²⁶⁻²⁸ has a much more superior PDH performance than commonly used VO_r/Al₂O₃. The turnover frequency (TOF) is six times higher by loading VO_x on ZrO₂ than on Al₂O₃. The remarkable improvement of reactivity has not been reported in previous studies. Rate measurement of catalysts with gradient V loadings was employed to identify the active site of the VO_r/ZrO₂ catalyst. In situ Raman spectroscopic measurements, X-ray photoelectron spectroscopy (XPS) and density functional theory (DFT) calculations were used to determine the structure evolution of VO_x under a reducing atmosphere, and they show that many more V-O bonds on ZrO2 (V=O, V-O-V and V-O-Zr) are consumed during reduction. On this basis, we proposed that the facile reducing nature of V-O bonds promotes the formation of lower coordinated V species which accounts for C-H activation enhancement.

Results and discussion

Catalyst structure

A series of characterization techniques were used to determine the bulk and surface structure of the catalysts (VO_x loaded on ZrO_2 and Al_2O_3 are denoted as xVZr and xVAl where x represents the mass percentage of V, metal base). The bare ZrO_2 support, 1VZr and 1VAl were characterized by X-ray diffraction (XRD). As shown in Fig. 1a, ZrO_2 is mainly composed of monoclinic phase (m- ZrO_2 , JCPDS 72-1669) with small amount of a tetragonal phase (t- ZrO_2 , JCPDS 79-1771), which is consistent with the previous report using the same preparation method. The same XRD pattern of 1VZr and pure ZrO_2 indicates that VO_x does not change the crystalline structure of ZrO_2 . In addition, for VO_x loaded catalysts, only diffraction lines of the support could be detected (Fig. 1a and $S1a\dagger$), which means amorphous VO_x is well dispersed on these catalysts. 30,31

Vis-Raman spectroscopic measurements, which are sensitive to the presence of V2O5, were performed over the catalysts to gain insight into the kind of vanadium species. Raman spectra of the ZrO2 support, 1VZr and 1VAl are displayed in Fig. 1b. A band at 1011 cm⁻¹ is attributed to vanadyl stretching of surfacedispersed VO_x. No sharp band at 995 cm⁻¹, assigned to the stretching vibration of V=O in crystal V₂O₅, is detected. With the V loading increasing, a band characteristic of V₂O₅ at 995 cm⁻¹ appears after the loading reached 3 wt% (Fig. S1b†). This demonstrates the existence of crystal V2O5 in 3VZr and 4VZr. From the enlarged region on the left from 75 to 160 cm⁻¹, a small band which corresponds to V₂O₅ at around 150 cm⁻¹ can be seen in 2.5VZr. This implies that VO_x begins to crystalize at 2.5VZr. Besides, the absence of a peak at around 770 cm⁻¹ suggests no ZrV₂O₇ is formed in our samples.³² The Raman spectra demonstrate that VO_x species are well dispersed as oligomeric species when the loading is less than 2 wt% and V₂O₅ crystals are formed after the loading reaches 2.5 wt%.

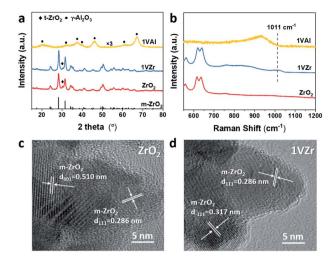


Fig. 1 (a) XRD patterns of the ZrO_2 support and VO_x supported on ZrO_2 and Al_2O_3 . (b) Raman spectra (532 nm excitation) of the ZrO_2 support and VO_x supported on ZrO_2 and Al_2O_3 . TEM images of (c) ZrO_2 and (d) VO_x/ZrO_2 .

Surface density of V was calculated through the BET surface area and V loading. The detailed results are listed in Table S1.† 2.5VZr has a V density of 6.9 $\rm nm^{-2}$ which is a bit lower than the theoretical monolayer coverage. It has been proved that the presence of $\rm V_2O_5$ is inevitable below theoretical monolayer coverage by a simple incipient wetness impregnation method. This agrees with the observation of the $\rm V_2O_5$ peak in 2.5VZr through Raman spectroscopic measurements.

The morphologies of ZrO₂ and 1VZr were also characterized by TEM (Fig. 1c and d). ZrO₂ is in the form of a nanoparticle with a relevant uniform size of around 30 nm and its morphology does not change after loaded with VO₂. Lattice

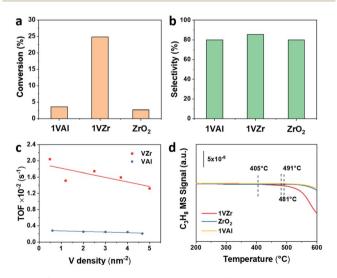


Fig. 2 (a) Initial propane conversion and (b) propylene selectivity (based on all products) of 1VAI, 1VZr and ZrO₂. Reaction conditions: $m_{\rm cat}=0.4~{\rm g}$; $C_3H_8:N_2:H_2=7:36:7;T=550~{\rm C}$; inlet flow = 50 mL min⁻¹. (c) Comparison of TOF values between VZr and VAI with different V densities. (d) C_3H_8 -TPSR of 1VZr, ZrO₂ and 1VAI.

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fringes of V₂O₅ could not be found because amorphous VO_x is well dispersed on the surface of ZrO2. The TEM results are in good agreement with the observations of XRD and Raman results.

Catalytic performance

Catalytic performance of VO_x supported on ZrO₂ and γ-Al₂O₃ for the PDH reaction was studied. The initial propane conversion and propylene selectivity based on all products are illustrated in Fig. 2a and b. The initial conversion of 1VZr is approximately five times higher than that of 1VAl and pure ZrO₂, which shows a significant support effect. The total selectivity towards propylene is around 80% and gas-phase selectivity towards it is >90% (Fig. S2a†) for all catalysts, which are at a similar level with respect to those of the VO_x-based catalysts in previous studies. 10,14,18,33 In addition, we performed 120 min of the on stream reaction and studied the reaction-regeneration cycles over 1VZr (Fig. S2b and S3†). The propane conversion and propylene selectivity exhibit little change during six cycles, implying outstanding regeneration stability.

Because the surface areas of ZrO₂ and Al₂O₃ are different, when the V loadings of VZr and VAl are identical, they have different V densities. Considering the activity is also influenced by V density, the TOFs of VZr and VAl were also compared based on V density (Fig. 2c and Table S2†). The calculated TOF of VZr is almost 6-fold higher than that of VAI and only slightly decreases with the V surface density, which implies that the activity is strongly support-dependent. This phenomenon is similar to that of the GaO_x catalytic system, where the activity is not a consequence of Ga nuclearities, but of the Ga-O-support interaction.34

To further evaluate the intrinsic activity in C-H activation, propane temperature-programmed surface reaction (C₃H₈-TPSR) experiments were carried out over 1VZr, ZrO2 and 1VAl. The temperature where the C₃H₈ signal begins to drop is determined as the C-H activation temperature. As shown in Fig. 2d, the initial C-H activation temperature of 1VZr is around 405 °C, which is almost 80 and 90 °C lower than that of ZrO₂ (481 °C) and 1VAl (491 °C). Moreover, the C-H activation temperature of 3VAl (similar V density to 1VZr, Fig. S4†) was also tested, which is 100 °C higher than that of 1VZr. Along with the TOF difference between VZr and VAl, we can conclude that the intrinsic C-H activation ability of 1VZr is distinguishable.

Active phase identification

It has been proved that pure ZrO2 can also catalyze propane dehydrogenation through coordinatively unsaturated Zr (Zr_{cus}).³⁵⁻³⁷ In addition, Jeon et al. prepared V-Zr mix oxide and found that V incorporation into the ZrO2 bulk phase would promote generation of more Zr_{cus}.38 Although in our VO_x/ZrO₂ system, VO_x is supported on the surface of a support, not in the bulk phase, there still remains different possibilities for this superior performance of the VO_x/ZrO₂ catalyst: ZrO₂ enhances the activity of VO_x , VO_x enhances the activity of ZrO_2 (Zr_{cus}), or a combination of the two. Therefore, it is essential to verify if the active phase is VO_x or Zr_{cus} in order to understand the origin of

this dramatic performance improvement. To this end, various experiments were performed to figure out whether the catalytic performance is due to VO_x or Zr_{cus} or both of them.

The propane consumption rate versus V loading is shown in Fig. 3. The reaction rate rises linearly as the V loading increases until the V loading reaches 2 wt%, indicating VO_r should be the active component. After this linearly increasing period, the C₃H₈ consumption rate does not change with increasing V loading because full VOx overlayers are formed at 2 wt% and crystalized V2O5 appears afterward as confirmed by the Raman spectra (Fig. S1b†). In addition, previous articles reported a shift of XPS Zr 3d peaks and reduction peaks of ZrO₂ seen in the H₂ temperature programmed reduction (H₂-TPR) profile if Zr_{cus} is the active site.37,39,40 However, these phenomena cannot be observed in the VO_x/ZrO₂ system. No ZrO₂ reduction peak can be observed up to 600 °C (Fig. 4a), while our reactions were conducted at 550 °C. In addition, the XPS test also confirms that the binding energy of Zr 3d in ZrO₂ and 1VZr has no difference (Fig. S5†). These results provide evidence that ZrO2 is not the

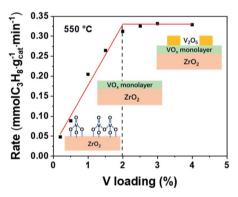


Fig. 3 C₃H₈ reaction rate versus different V loadings on VZr catalysts at 550 °C.

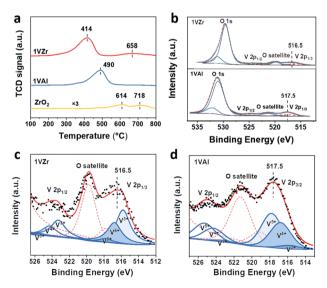


Fig. 4 (a) H₂-TPR profiles of 1VZr, 1VAl and pure ZrO₂. (b) XPS O 1s and V 2p peaks for 1VZr and 1VAl. Deconvolution results of V 2p_{3/2} for (c) 1VZr and (d) 1VAl after reduction for 30 minutes

active phase in VZr catalysts. Along with catalytic activity being linearly related to the amount of V, which directly proves VO_x is responsible for the activity, we suggest VO_x rather than Zr_{cus} to be the main active phase. However, it is a pity that these experiments cannot identify which kind of V–O bond (V=O, V–O–V, V–O–Zr or all of them) is responsible for C–H activation. A potential method to solve this problem is Surface Organometallic Chemistry (SOMC), 41–43 which provides access to producing well-defined isolated VO_x only possessing V=O and V–O–support. This direction still needs further investigation.

Structure-performance correlation

H₂-TPR experiments were performed to test the reducibility and VO_r-support interaction of the catalysts (Fig. 4a). The main peak at around 400-500 °C is attributed to the reduction of VO_x and peaks above 600 °C are ascribed to a partial reduction of ${\rm ZrO_2.^{36,44-46}}$ The reduction temperature of ${\rm VO}_x$ for 1VZr is lower than that for 1VAl. Considering that the reduction temperature is influenced by the dispersion of VO_x, 15,47 we performed H₂-TPR on a series of VAI and VZr catalysts (Fig. S6†). The reduction temperatures of VZr catalysts are at about 400 °C while those of VAl catalysts are at about 500 °C. Thus, the low reduction temperature of the VZr catalyst implies that the interaction between VOx and ZrO2 is weaker than that between VOx and Al₂O₃. This phenomenon is further discussed in the in situ Raman spectroscopy section below. Alternatively, the peak area of 1VZr is larger than that of 1VAl. We calculated the H: V ratio through the peak area and estimated the average oxidation state (AOS) of V. As listed in Table S3,† the AOS of V in 1VZr and 1VAl corresponds to 3.5 and 4.0, which indicates that VO_x can be more readily and deeply reduced on ZrO2.

To further identify the valance state of V, XPS investigation was carried out over 1VZr and 1VAl after $\rm H_2$ reduction for 30 min. V 2p core levels of 1VZr and 1VAl are displayed in Fig. 4b. We compare the binding energy of V in different catalysts qualitatively. The binding energy of V 2p_{2/3} centers at 516.5 eV in 1VZr and 517.5 eV in 1VAl. This indicates that the valance state of V in 1VZr is lower than that in 1VAl. According to the deconvolution results of V 2p shown in Fig. 4c, d and Table S4,† the oxidation state of V is a mixture of V⁵⁺, V⁴⁺ and V³⁺, which is consistent with previous results.^{33,44,48} The fraction of V³⁺ is 48.6% in 1VZr, which is higher than 6.6% in 1VAl.

It has been elucidated in the former articles that V^{3+} is more active in the PDH reaction. However, seldom detailed analyzes have been conducted to identify the structure of VO_x in a reduced state and the structure-performance correlation is not clear. Therefore, *in situ* UV-Raman spectroscopy, which is more sensitive to the surface VO_x structure, was performed to monitor the evolution of V=O, V-O-V, and the V-O-support.

The evolution of V–O–V, V–O–Zr and V=O in 1VZr during $\rm H_2$ reduction is shown in Fig. 5a and b. A broad band at 750–950 cm⁻¹ is ascribed to V–O–V and V–O–Zr bonds.^{32,50} The band at around 1020 cm⁻¹ is assigned to V=O in dispersed $\rm VO_x$.^{27,32,51} The number of V–O–V and V–O–Zr bonds decreases substantially which implies that plenty of these bonds are consumed during reduction. In addition, nearly 60% of V=O in 1VZr disappeared after reduction

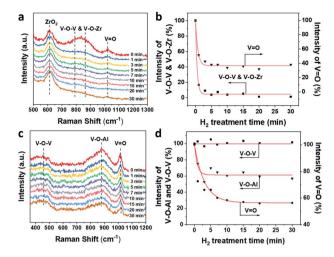


Fig. 5 (a) In situ UV-Raman (325 nm excitation) spectra of VZr during H_2 reduction at 550 °C. (b) Intensities of V-O-V, V-O-Zr, and V=O versus H_2 treatment time. (c) In situ UV-Raman (325 nm excitation) spectra of 1VAl during H_2 reduction at 550 °C. (d) Intensities of V-O-V, V-O-Al and V=O versus H_2 treatment time.

for 30 min. The comparative experiment of 1VAl is displayed in Fig. 5c and d. The bands attributed to V–O–V and V–O–Al on the Al₂O₃ support are located at 400–600 cm⁻¹ and 910 cm⁻¹ respectively.⁴⁹ A sharp band at 1018 cm⁻¹ is ascribed to V=O.⁴⁹ In contrast with V–O–Zr and V–O–V in the 1VZr sample, the V–O–Al band only decreases to 60% of that for fresh 1VZr and no change of the V–O–V band is detected which implies V–O–V bonds on the Al₂O₃ support cannot be reduced under H₂ treatment. Meanwhile, 40% of V=O in 1VAl is reduced, which is less than that in 1VZr. The result of *in situ* Raman spectroscopy confirms that not only V=O and V–O–Zr, but also V–O–V can be deeply reduced in 1VZr which results in the deeper reduction degree of 1VZr observed in H₂-TPR and XPS studies.

The facile reduction nature of V–O bonds in the VZr catalyst can be interpreted by the lower electronegativity of Zr compared with Al, which causes the difference of the interaction between VO_x and the support. It has been shown that a lower support cation electronegativity can result in a higher electron density of the V–O–S bond, which lead to these bonds being readily reduced. ^{52,53} Along with the lower reduction temperature of VZr shown in the H_2 -TPR profile, it can be deduced that the interaction between VO_x and ZrO_2 weakens the V–O bonds thus resulting in the facile reduction nature of the VZr catalyst.

Note that when O atoms are removed during the reduction period, the chemical environment of V changes, which leads to the formation of coordinatively unsaturated V. It is established that metal cations $(Ga^{3+}, Zn^{2+}, Zr^{4+}, etc.)$ with a low coordination number are more active for the reaction. As more V–O bonds are reduced in the VZr catalyst, we could deduce that lower coordinated V sites are more active for the C–H activation. Our recent DFT calculations also proved that coordinatively unsaturated VO_x was more active in C–H activation while this work gives experimental evidence.

Table 1 Oxygen vacancy formation energies of vanadium supported on m-ZrO₂($\bar{1}11$) and γ-Al₂O₃(100)

	Oxygen vacancy formation energies (eV)				
	v=o	V-O-V	V-O-support		
V_2O_5/m - $ZrO_2(\bar{1}11)$ V_2O_5/γ - $Al_2O_3(100)$	0.13 -0.07	$-0.13 \\ 0.44$	0.03 0.36	0.14 0.35	$0.34 \\ -0.25$

DFT calculations

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DFT calculations were conducted to gain further insight into the structure of the active site and its influence on catalytic performance. Noting that the TOF difference between VZr and VAl exists with all the V densities, we constructed dimeric vanadium oxide species on the m-ZrO₂($\bar{1}11$) and γ -Al₂O₃(100) surface as representatives^{36,55} (Fig. S7†).

We calculated oxygen vacancy formation energy of different oxygen atoms, with the energy of $H_2O(g)$ and $H_2(g)$ as the reference, in these vanadium oxide dimers to confirm the different reducibilities they showed in our experiment results. As listed in Table 1, oxygen atoms are removed from V=O, V-O-V and the V-O-support. Four of five V-O bonds in VZr (V=O, V-O-V and two V-O-Zr) have relatively low oxygen formation energies while only two of five in VZr do (V=O, V-O-Al). The calculated oxygen vacancy formation energies are in good agreement with the Raman spectroscopy result that V=O, V-O-V and V-O-Zr bonds in VZr can be reduced under a H_2 atmosphere while only V=O and part of V-O-Al are reduced in VAl.

Since VZr and VAl have different degrees of reduction, we constructed V_2O_2/m -Zr $O_2(\bar{1}11)$ and V_2O_3/γ -Al $_2O_3(100)$ models to represent the catalyst structure in a reduced state (Fig. 6a and b). In terms of the *in situ* Raman results and calculated oxygen vacancy formation energies, three (one in each V=O, V-O-Zr and V-O-V) and two (one V=O and one V-O-Al) V-O bonds with the lowest oxygen vacancy formation energies were removed from the initial dimeric V_2O_5 structure respectively. The propane dehydrogenation barriers over partially reduced VZr and VAl were computed and transition state (TS) geometries are shown in Fig. 6c. The first and second step C-H activation

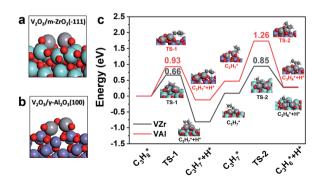
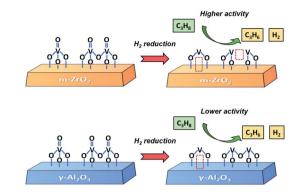


Fig. 6 Reduced catalyst models of (a) V_2O_2/m -Zr O_2 (111) and (b) V_2O_3/γ -Al $_2O_3$ (100). (c) Calculated potential energy diagrams of the first and second propane dehydrogenation step.



Scheme 1 Propane dehydrogenation on VZr and VAl catalysts with different V structures and VO_x reduction degrees.

barriers in VZr is 0.66 eV and 0.85 eV, respectively, while in VAl they are 0.99 eV and 1.26 eV, respectively. DFT calculations also confirm that low coordinated V species are more active in C–H activation, which is consistent with the highest TOF and the lowest C–H activation temperature for VZr observed in the catalytic performance and C_3H_8 -TPSR tests.

Based on the experimental and theoretical results, we propose a structure–performance correlation of VZr and VAl catalysts, as shown in Scheme 1. VO_x can be readily reduced on ZrO_2 because the interaction between VO_x and ZrO_2 facilitates the reduction of V=O, V-O-V and V-O-Zr bonds. However, for the VAl catalyst, only V=O and some V-O-Al bonds can be reduced. Thus, more low coordinated V species form in the VZr catalyst during reduction, and these low coordinated V species exhibit a better performance in PDH.

Conclusions

In summary, we prepared VO_x loaded on ZrO_2 through a simple incipient wetness impregnation method, which exhibits a dramatically improved performance compared with VO_x loaded on Al_2O_3 for propane dehydrogenation. The $TOF_{C_3H_8}$ of 1VZr is determined to be 0.0161 s⁻¹, which is almost six times higher than that of 1VAl.

We further prove that the remarkable reactivity of the 1VZr catalyst was attributed to the promotion of C–H activation over VO_x species rather than it over ZrO_2 . Besides, combining in situ Raman and XPS spectroscopy results, we propose the enhanced C–H activation on VZr results from the facile reduction of V=O, V–O–V and V–O–Zr bonds, thus producing deeply reduced and lower coordinated V species. DFT calculations also confirm that the C–H rupture energy barrier is lower for partially reduced VZr with low coordinated V species. Considering the dramatic performance achieved through the interaction between VO_x and ZrO_2 , our work provides a new insight into high-performance VO_x -based catalysts for propane dehydrogenation.

Conflicts of interest

There are no conflicts to declare.

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