The catalytic properties of Al2O3-supported vanadia with a wide range of VOx surface density (1.4–34.2 V/nm2) and structure were examined for the oxidative dehydrogenation of ethane and propane. UV-visible and Raman spectra showed that vanadia is dispersed predominantly as isolated monovanadate species below ~2.3 V/nm2. As surface densities increase, two-dimensional polyvanadates appear (2.3–7.0 V/nm2), along with increasing amounts of V2O5 crystallites at surface densities above 7.0 V/nm2. The rate constant for oxidative dehydrogenation (k1) and its ratio with alkane and alkene combustion (k2/k1 and k3/k1, respectively) were compared for both alkane reactants as a function of vanadia surface density. Propene formation rates (per V atom) are approximately eight times higher than ethene formation rates at a given reaction temperature, but the apparent ODH activation energies (E1) are similar for the two reactants and relatively insensitive to vanadia surface density. Ethene and propene formation rates (per V atom) are strongly influenced by vanadia surface density and reach a maximum value at intermediate surface densities (~8 V/nm2). The ratio of k2/k1 depends weakly on reaction temperature, indicating that activation energies for alkane combustion and ODH reactions are similar. The ratio of k3/k1 is independent of surface density for ethane but increases slightly with vanadia surface density for propane, suggesting that isolated structures prevalent at low surface densities are slightly more selective for alkane dehydrogenation reactions. The ratio of k3/k1 decreases markedly with increasing reaction temperature for both ethane and propane ODH. Thus, the apparent activation energy for alkene combustion (E3) is much lower than that for alkane dehydrogenation (E1) and the difference between these two activation energies decreases with increasing surface density. The lower alkene selectivities observed at high vanadia surface densities are attributed to an increase in alkene adsorption enthalpies with increasing vanadia surface density. The highest yield of alkene is obtained for catalysts containing predominantly isolated monovanadate species and operated at high temperatures that avoid homogeneous reactions (<~800 K). © 2002 Elsevier Science (USA)

INTRODUCTION

Low-molecular-weight alkenes, such as ethene and propene, can be formed via nonoxidative dehydrogenation of the corresponding alkane. Nonoxidative dehydrogenation reactions are endothermic and lead to the concurrent formation of carbon and of lower molecular weight alkanes, both of which decrease alkene yields. Oxidative dehydrogenation (ODH) of light alkanes offers a potentially attractive route to alkenes, since the reaction is exothermic and avoids the thermodynamic constraints of nonoxidative routes by forming water as a by-product. In addition, carbon deposition during ODH is eliminated, leading to stable catalytic activity. However, the yield of alkenes obtained by ODH on most catalysts is limited by alkene combustion to CO and CO2 (CO2).

Previous studies have shown that supported vanadia is the most active and selective simple metal oxide for alkane ODH, because its reducible nature leads to the rapid redox cycles required for a catalytic turnover (1–5). Other studies (6–18) have probed the kinetics and selectivity of ethane and propane ODH on VOx/Al2O3 and on VOx species supported on other metal oxides. These studies have shown that ODH rates per gram of catalyst and alkene selectivities are higher on dispersed VOx than on V2O5. The local structure of supported VOx domains strongly influences ODH reaction rates and selectivity (12–14). The effects of oxide domain size and of support on rates and selectivity have been attributed to the acid–base properties of the VOx and support surfaces (1–5, 7–18).

Alkane ODH reactions proceed via the reaction network shown in Scheme 1 (4, 10, 12–14, 19, 20), in which alkanes react with lattice oxygen to form alkenes, with a rate constant k1, or COx, with a rate constant k2. The alkenes formed undergo subsequent oxidation to COx, with a rate constant k3. Reaction rates are nearly zero order in O2 and show a weak inhibition by H2O formed in ODH reactions (14, 21). A pseudo-first-order kinetic analysis of Scheme 1 is sufficiently accurate to provide all three rate constants from rate and selectivity data as a function of...
residence times require high values of \( k_1 \) and low values of \( k_2/k_1 \) and \( k_3/k_1 \).

The effects of VO\(_x\) surface density and structure on rates and selectivities for ethane and propane ODH reactions have been previously reported (7–18), with somewhat inconsistent results. For example, for ethane ODH on VO\(_x\)/γ-Al\(_2\)O\(_3\) catalysts, the vanadia surface density that produces the highest ethane and ethene rates per V has been variously reported to occur at \( \sim 33 \) (9), \( \sim 50 \) (7), or \( \sim 70\% \) (2) of the polyvanadate monolayer coverage. As another example, the variation of propene selectivity with vanadia surface density over a range of 1.7–3.9 V/nm\(^2\) has been reported to be insignificant (2, 17) and quite significant, varying by \( \sim 20\% \) (13), for propane ODH on VO\(_x\)/γ-Al\(_2\)O\(_3\) catalysts. The lack of detailed structural characterization and rigorous kinetic analysis of integral (high conversion) rate and selectivity data, and the temperature gradients and possible homogeneous reactions prevalent at the high temperatures and conversions of many previous studies, have contributed to the persistent controversies.

Here, we report a detailed kinetic analysis of ethane ODH reactions on VO\(_x\)/Al\(_2\)O\(_3\) samples with a wide range of VO\(_x\) surface densities (1.4–34 VO\(_x\)/nm\(^2\)) and contrast the results with those obtained using propane as the reactant. We have combined this analysis with detailed structural characterization of these catalysts in order to relate their local structure to the values of the rate constants, which characterize the function of these catalytic materials. In addition, we contrast the reaction rates of ethane and propane molecules with different C–H bond energies and different adsorption energies of the respective alkenes.

**EXPERIMENTAL METHODS**

VO\(_x\)/Al\(_2\)O\(_3\) catalysts were prepared by incipient wetness impregnation of fumed γ-alumina (Degussa, AG, 100 m\(^2\)/g) with an aqueous solution of ammonium metavanadate (99%, Aldrich, Inc.) and oxalic acid (Mallinckrodt analytical grade) in a 1:2 weight ratio (12) with a pH of \( \sim 2 \). The impregnated samples were dried in air at 398 K for \( \sim 16 \) h, crushed, treated in 1.67 cm\(^3\)/s\(^{-1}\) (STP) dry air (Airgas zero grade) at 773 K for 3 h, and sieved to retain particles with diameters of 180–355 μm.

Surface areas were measured by N\(_2\) physisorption (Airgas, 99.999%) at its normal boiling point using a Quantasorb surface area analyzer and standard multipoint BET analysis methods. The catalyst samples were treated in flowing He (Airgas, 99.999%) at 393 K for 2 h before N\(_2\) physisorption measurements.

Diffuse reflectance UV–visible spectra were measured with a Varian–Cary 4 spectrophotometer equipped with a Harrick diffuse reflectance attachment. Samples were dehydrated in 20% O\(_2/He\) (1.67 cm\(^3\)/s\(^{-1}\) (STP), Praxair, 99.99%) at 723 K before measuring spectra at ambient temperature between 1 and 5 eV. The Kubelka–Munk function \( F(R_\infty) \) was used to convert diffuse reflectance data into absorption spectra using MgO as a standard (22). The absorption edge energy was estimated from the \( x \)-intercept of a linear fit in the near-edge region in a plot of \( F(R_\infty)h\nu \)\(^{1/2} \) as a function of \( h\nu \) (22).

Raman spectra were obtained using a Hololab Series 5000 Raman spectrometer (Kaiser Optical) with a frequency-doubled 75-mW Nd-YAG laser at a wavelength of 532 nm. VO\(_x\)/Al\(_2\)O\(_3\) samples (\( \sim 0.05 \) g) were pressed into wafers (0.9-cm diameter, \( \sim 0.1 \)-cm thickness) at 40 MPa and Raman spectra were measured at ambient temperatures before and after dehydration treatments. Dehydration was carried out at 723 K in flowing 20% O\(_2\) (Airgas, 99.99%) in He (Scott Specialty Gases, Inc., 99.999%) at a total flow rate of 1.0 cm\(^3\)/s\(^{-1}\) (STP) for 0.75 h. A rotating Raman cell (20 Hz) was used to prevent laser heating of the samples (23).

Reaction rate and selectivity data were measured in a packed-bed flow reactor with plug-flow hydrodynamics (24). VO\(_x\)/Al\(_2\)O\(_3\) catalysts (0.01–0.3 g) were diluted with quartz granules (180- to 355-μm diameter, 0.01–0.3 g) in order to prevent bed temperature gradients. Reaction measurements with Al\(_2\)O\(_3\), quartz chips, or empty reactors did not lead to detectable products in the temperature range of the study. Typical propane and O\(_2\) conversion ranges on VO\(_x\)/Al\(_2\)O\(_3\) catalysts were 1–2 and 10–20%, respectively. A Hewlett–Packard 6890 gas chromatograph equipped with packed (Supelco Carboxen 1004) and capillary (HP-1, 50 m, 320 μm) columns and thermal conductivity and flame ionization detectors was used to measure the concentrations of reactants and products in the effluent stream.

Ethane ODH reactions were examined at 663–743 K, while the faster propane ODH reactions were examined at lower temperatures (603–663 K). No homogeneous ethane or propane reactions were detected below 800 K. Reactor residence times were adjusted by varying reactant flow rates between 0.083 and 4.2 cm\(^3\)/s\(^{-1}\) (STP). Reactant mixtures consisted of ethane (14 kPa, Scott Specialty Gases, Inc., 99.999%) or propane (14 kPa, Airgas, 99.9%) with O\(_2\) (1.7 kPa, Praxair, 99.99%) and He as an inert diluent (Airgas, 99.999%).
Conversion and selectivity were calculated from the measured GC compositions. These data were used to calculate reaction rates from the reactant and product molar flow rates. The reaction rates and selectivities as a function of residence time were used to obtain initial alkane dehydrogenation and combustion reaction rates and the rate constants shown in Scheme 1. The sequence in Scheme 1 can be used to obtain the $k_1$ and $k_2$ rate constants from the initial alkene selectivity,

$$S_{C_{x}H_{2x}}^{0} = k_1/(k_1 + k_2),$$

and $k_3$ from the observed changes in alkene selectivity with residence time,

$$S_{C_{x}H_{2x}} = S_{C_{x}H_{2x}}^{0}[1 - (k_1 + k_2 + k_3)C_V\tau/2],$$

where $\tau$ is residence time and $C_V$ is the concentration of V atoms per reactor volume, as described in detail elsewhere (12).

RESULTS AND DISCUSSION

Catalyst Characterization

BET surface areas and apparent VO$_x$ surface densities, estimated from the V content and the BET surface areas, are shown in Table 1 for all VO$_x$/Al$_2$O$_3$ samples. Surface areas (per gram of sample) decrease with increasing VO$_x$ content. This trend corresponds to the low surface area of the added VO$_x$ because the surface area normalized per alumina content, listed in the last column of Table 1, is approximately constant. However, the 30 wt% sample shows a decrease in surface area per gram of alumina, suggesting that the formation of larger V$_2$O$_5$ crystallites blocks some support pores.

The UV–visible spectra and the absorption edge energies for all VO$_x$/Al$_2$O$_3$ samples are shown in Figs. 1a and 1b, respectively. VO$_x$/Al$_2$O$_3$ samples with 8.0, 12.5, 16.6, and 34.2 V/m$^2$ exhibit two linear regions in the near-edge region, indicating the presence of two distinct VO$_x$ structures. For those samples, two edge energies are measured and both are included in Fig. 1b. The low-energy absorption edge decreases in energy with increasing surface density, approaching values typical of bulk V$_2$O$_5$ (2 eV) (25). The edge energies based on the second linear portion of the near-edge region are higher than those based on the first linear portion and appear to be independent of the vanadia surface density. Absorption edge energies above 2.5 eV are attributable to V$^{5+}$ in distorted tetrahedral

### TABLE 1

<table>
<thead>
<tr>
<th>V$_2$O$_5$ loading (wt%)</th>
<th>Surface area (m$^2$/g)</th>
<th>Nominal VO$_x$ surface density (V/m$^2$)</th>
<th>Surface area normalized by amount of support (m$^2$/g Al$_2$O$_3$)</th>
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FIG. 1. (a) Diffuse reflectance UV–visible spectra and (b) absorption edge energies for VO$_x$/Al$_2$O$_3$ catalysts.
FIG. 2. Raman spectra for VO\textsubscript{x}/Al\textsubscript{2}O\textsubscript{3} catalysts (obtained at 298 K in flowing dry 20% O\textsubscript{2}/He after treatment at 723 K for 0.75 h).

coordination (25). Thus, UV–visible spectroscopy suggests that V\textsuperscript{5+} is present in distorted tetrahedrally coordinated species at low vanadia surface densities and that some of these tetrahedrally coordinated cations remain at higher surface densities, coexisting with an increasing fraction of V\textsuperscript{3+} present in distorted tetragonal pyramidal coordination, similar to that present in V\textsubscript{2}O\textsubscript{5} (25).

Raman spectra, obtained at ambient temperature after treatment at 723 K in 20% O\textsubscript{2}/He, for the VO\textsubscript{x}/Al\textsubscript{2}O\textsubscript{3} samples are shown in Fig. 2. V=O stretches in isolated tetrahedral monovanadates appear at 1033 cm\textsuperscript{-1}, while V–O–V stretches in two-dimensional polyvanadates have been proposed to lead to broad Raman features at 750–1000 cm\textsuperscript{-1} (25–27). The remaining bands, appearing at 998, 706, 530, 489, 410, 305, 289, 203, and 150 cm\textsuperscript{-1}, correspond to bulk V\textsubscript{2}O\textsubscript{5} crystals (1, 25). The V\textsubscript{2}O\textsubscript{5} bands in some of these spectra are intense, but the Raman cross section for V\textsubscript{2}O\textsubscript{5} crystals is about 10 times larger than for monovanadate species (23). Using this value, the ratio of V atoms in V\textsubscript{2}O\textsubscript{5} to those in monovanadate increases from 0 to 2.5 as VO\textsubscript{x} surface densities increase from 1.4 to 34.2 V/nm\textsuperscript{2}. Geometric calculations, assuming flat, ordered support surfaces, predict that a theoretical monolayer of monovanadate species occurs at 2.3 V/nm\textsuperscript{2}, and at 7.0 V/nm\textsuperscript{2} if polyvanadate species are present (12, 28). Over this range of VO\textsubscript{x} surface densities, the ratio of V\textsubscript{2}O\textsubscript{5} to monovanadate species increases from ~0.02 to ~0.1. Thus, under either definition of a monolayer, most of the surface is covered by vanadate species other than V\textsubscript{2}O\textsubscript{5}. Specifically, the V\textsubscript{2}O\textsubscript{5}:monovanadate ratio increases from 0.085 for the catalyst with a vanadia surface density of 3.6 V/nm\textsuperscript{2} to 0.115 for the catalyst with a vanadia surface density of 8.0 V/nm\textsuperscript{2}. This represents an increase of 35% in the ratio of V\textsubscript{2}O\textsubscript{5} to monovanadate species at the same time that the vanadia concentration doubled, indicating that the additional vanadia formed structures that are neither isolated monovanadates nor V\textsubscript{2}O\textsubscript{5} crystallites. The V\textsubscript{2}O\textsubscript{5}:monovanadate ratio increases to 0.63, 1.6, and 2.5 for the catalysts with vanadia surface densities of 12.5, 16.6, and 32.4 V/nm\textsuperscript{2}, which have vanadia concentrations of 15, 20, and 30 wt%, respectively. The increase in the V\textsubscript{2}O\textsubscript{5}:monovanadate ratio is proportionally larger than the additional vanadia content of these catalysts, suggesting that all added vanadia forms V\textsubscript{2}O\textsubscript{5} structures while simultaneously incorporating some monovanadates.

The spectroscopic results presented in Figs. 1 and 2 suggest that distorted tetrahedral monovanadate structures connected to the support via V–O–Al bonds are prevalent at low surface densities. As V surface densities increase, two-dimensional polyvanadates form via reactions leading to V–O–V bonds connecting neighboring VO\textsubscript{x} species. VO\textsubscript{x} species exceeding monolayer coverages react with polyvanadates structures to form three-dimensional structures
that ultimately crystallize into bulk V$_2$O$_5$, with some evidence for residual monovanadate species.

**Oxidative Dehydrogenation Rates and Selectivity**

C$_2$H$_4$, CO, and CO$_2$ formation rates, C$_2$H$_6$ conversion level, and C$_2$H$_4$, CO, and CO$_2$ selectivities are shown as a function of nominal reactor residence time in Figs. 3a and 3b for the 8.0 V/nm$^2$ sample. The lines shown were used to extrapolate rates and selectivity to zero reactor residence time or reactant conversion and to estimate the rates of secondary oxidation of C$_2$H$_4$. The decrease in C$_2$H$_6$ conversion rates with increasing residence time (Fig. 3a) reflects a slight inhibition of ethane dehydrogenation rates by the water formed during reaction (21), as reported previously for propane reactions (14, 29, 30). The decrease in ethene selectivity with increasing residence time (Fig. 3b) arises from secondary ethene oxidation reactions (Scheme 1). Data acquisition methods similar to those described for ethane ODH were used to obtain the propane ODH data. The conversion and selectivity trends displayed in Fig. 3 for ethane ODH are similar to those observed for propane ODH, as described in more detail elsewhere (13, 14).

Figures 4a and 4b show the effect of vanadia surface density on the initial rate of alkene formation per V atom (extrapolated to zero residence time). For both ethane and propane reactants, these apparent turnover rates reach a maximum at intermediate vanadia surface densities. The maximum rate is reached at a surface VO$_x$ density of $\sim$8 V/nm$^2$ for both reactants, and this maximum rate is about four times greater than that obtained on the VO$_x$/Al$_2$O$_3$ sample with the lowest surface density (1.4 V/nm$^2$). Well-dispersed monovanadate species prevalent in the latter sample are significantly less active than the oligomeric VO$_x$ structures that appear as the vanadia surface density increases. The rate per V atom increases with surface densities, even though V atoms become increasingly unavailable for catalysis as three-dimensional structures form. Thus, the surface reactivity of VO$_x$ increases even more markedly with surface density than is suggested by the data in Fig. 4. For a given VO$_x$ surface density, the apparent turnover rates for propane reactants is approximately eight times higher than for ethane, but the effects of vanadia surface density and the higher surface reactivity of the VO$_x$ oligomers are observed for both alkane reactants.

Figures 4a and 4b show that apparent turnover rates for both ethane and propane decrease at VO$_x$ surface densities above $\sim$8 V/nm$^2$, as also found previously for propane ODH on VO$_x$/ZrO$_2$ (31). These trends reflect the introduction of additional vanadia species within the inaccessible bulk of three-dimensional structures. The picture of the sample surfaces evolving from isolated and sparsely distributed monovanadates, to nearly complete polyvanadate monolayers, followed by the gradual building of additional layers would lead to low areal ODH rates at low surface densities, because most of the surface area is inactive Al$_2$O$_3$. The increasing coverage of the Al$_2$O$_3$ with monovanadate and with more reactive polyvanadates would increase the measured areal rates, until the entire surface was covered with VO$_x$ species. If the surface reactivity of such species becomes independent of the presence and scale of the third dimension, the areal rates would reach a constant value. These
trends are confirmed by the areal rates of alkene formation from ethane or propane, plotted as a function of vanadia surface density in Figs. 5a and 5b. These areal rates increase with increasing surface density and approach nearly constant values for vanadia surface densities higher than 10 V/nm², as also found for propane ODH on VOₓ/ZrO₂ (31). The constant areal rate obtained after completion of a polyvanadate monolayer suggests that the surface reac-

FIG. 4. Initial alkene formation rates as a function of vanadia surface density on VOₓ/Al₂O₃ catalysts: (a) ethane ODH, (b) propane ODH (663 K, 14 kPa C₂H₆ or C₃H₈, 1.7 kPa O₂, balance He).

FIG. 5. Initial alkene formation rates normalized per unit catalyst surface area as a function of vanadia surface density: (a) ethane ODH, (b) propane ODH (663 K, 14 kPa C₂H₆ or C₃H₈, 1.7 kPa O₂, balance He).

FIG. 6 shows activation energies (E₁) obtained from Arrhenius plots for ethene and propene formation rates from the corresponding alkanes as a function of VOₓ surface density. Ethane and propane dehydrogenation show similar activation energies, and the value decreases slightly for both reactants with increasing VOₓ surface density. Similar and even lower values of E₁ than those presented in Fig. 6 have been reported for ethane ODH on VOₓ/Al₂O₃.

Activity of VOₓ species becomes independent of domain size once the predominant structures involve V–O–V bonds.
The similar values of alkane, 1.7 kPa O2, balance He). It is the major by-product and only a small amount of CO2 is formed. For propane ODH reactions, initial propene selectivities are ~85% at the lowest vanadia surface densities, but they decrease to values similar to those for ethane reactants with increasing surface density (Fig. 7b).

The ratios of rate constants for alkane direct combustion and dehydrogenation (k2/k1) are shown in Fig. 8 as a function of VOx surface density. For ethane, k2/k1 is essentially independent of VOx surface density, but this ratio increases with increasing VOx surface density for propane reactants. For ethane, k3/k1 is weakly dependent on temperature, while for propane, k2/k1 is essentially independent of temperature on all samples, indicating that the activation energies for alkene and COx formation directly from alkanes (E1 and E2) are very similar (Fig. 9). The differences between these two activation energies (ΔE12 = E1 - E2) are small, positive, and independent of VOx surface density for ethane, but change from positive to negative with increasing VOx surface density for propane ODH. These trends are consistent with the observed trends in k2/k1 shown in Fig. 8.

Figure 10 shows ratios of rate coefficients for alkene combustion and alkane dehydrogenation (k3/k1). For both reactants, k3/k1 increases with increasing VOx surface density and decreases with increasing temperature on all samples. This is consistent with a lower activation energy for alkene reactions (E3) than for alkane reactions (E1).
FIG. 7. ODH selectivities as a function of vanadia surface density on VOₓ/Al₂O₃ catalysts: (a) ethane, (b) propane (663 K, 14 kPa alkane, 1.7 kPa O₂, balance He).

activation energy difference (\(\Delta E_{13} = E_1 - E_3\)) decreases for both alkanes with increasing VOₓ surface density (Fig. 11), consistent with trends shown in Fig. 10. \(E_1\) values decrease very slightly with increasing surface density; thus, the observed changes in \(\Delta E_{13}\) arise almost exclusively from an increase in the value of \(E_3\) with increasing VOₓ surface density.

Changes in alkene adsorption enthalpies may account for the observed increase in alkene combustion activation energies (\(E_3\)) with increasing vanadia surface density. Theoretical estimates of enthalpies for ethene adsorption on supported polyvanadate monolayers (\(V_2O_5(001)/TiO_2(001)\) (anatase)) and bulk \(V_2O_5(001)\) surfaces (32) give values of −79.6 and −39.7 kJ/mol, respectively. These density functional theory calculations are consistent with a higher activation energy on the bulk \(V_2O_5\) surface, which binds ethene much more weakly than two-dimensional \(VO_\chi\) structures. This is apparent from the definition of \(k_3\) as \(K_3k_{act}^3\), where \(K_3\) is the equilibrium constant for ethene adsorption and \(k_{act}^3\) is the rate constant for the activation of a C–H bond in ethene. Thus, the apparent activation energy is given by \(\Delta H_3 + E_{act}^3\), where \(\Delta H_3\) is the ethene adsorption enthalpy.

FIG. 8. Rate constant ratio \(k_2/k_1\) as a function of vanadia surface density on VOₓ/Al₂O₃ catalysts: (a) ethane (14 kPa C₂H₆, 1.7 kPa O₂, balance He), (b) propane (14 kPa C₃H₈, 1.7 kPa O₂, balance He).
and $E_{\text{act}}^{\text{C-H}}$ is the activation energy for the kinetically relevant C–H bond activation step in adsorbed ethene (29). The enthalpy of adsorption of ethane is not very sensitive to VO$_x$ surface structure ($-27.4$ kJ/mol on V$_2$O$_5$(001)/TiO$_2$ and $-25.4$ kJ/mol on V$_2$O$_5$(001) (32)). Taken together with the nearly constant values of $E_1$ (Fig. 6) on VO$_x$/Al$_2$O$_3$ samples with a wide range of surface density, this suggests that the energy for C–H bond activation in ethane ($E_{\text{act}}^{\text{C-H}}$) is not influenced by vanadia surface density. The mechanistic and kinetic resemblance between C–H bond activation steps in alkanes and alkenes and the different observed sensitivities of $E_1$ and $E_3$ on surface density suggest that only the adsorption enthalpy of alkenes, not that of alkanes and not the values of C–H bond activation energies ($E_{\text{act}}^{\text{C-H}}$, $E_{\text{act}}^{\text{C-H}}$), change with VO$_x$ surface density.

The sample with the lowest vanadia surface density (1.4 V/nm$^2$) and containing predominately monovanadates shows the highest initial alkene selectivity for both ethane and propane ODH. Increasing vanadia surface density either has no effect or decreases initial alkene selectivities.

![Figure 9](image_url)

**FIG. 9.** Difference between ODH activation energy and alkane combustion activation energy ($\Delta E_{12} = E_1 - E_2$) on VO$_x$/Al$_2$O$_3$ catalysts: (a) ethane (663 K, 14 kPa C$_2$H$_6$, 1.7 kPa O$_2$, balance He), (b) propane (663 K, 14 kPa C$_3$H$_8$, 1.7 kPa O$_2$, balance He).

![Figure 10](image_url)

**FIG. 10.** Rate constant ratio, $k_3/k_1$, as a function of vanadia surface density on VO$_x$/Al$_2$O$_3$ catalysts: (a) ethane reactant (14 kPa C$_2$H$_6$, 1.7 kPa O$_2$, balance He), (b) propane reactant (14 kPa C$_3$H$_8$, 1.7 kPa O$_2$, balance He).
The ratio of $k_2/k_1$ is independent of surface density for ethane and increases with increasing vanadia surface density for propane (Fig. 8). The ratio of $k_3/k_1$ increases with increasing vanadia surface density for both reactants (Fig. 10). Smaller values of $k_3/k_1$ and $k_2/k_1$ lead to higher alkene selectivity at all reactant conversion levels. As a result, the 1.4 V/nm² catalyst is the most selective for both alkanes. However, the alkene formation rates (per V atom) for the 1.4 V/nm² catalyst are approximately four times lower at a given reaction temperature compared to the 8.0 V/nm² catalyst (Fig. 4), but the rate difference can be overcome by increasing reaction temperature. The ratio of $k_2/k_1$ is relatively independent of reaction temperature, and the value of $k_3/k_1$ actually decreases with increasing temperature for both reactants, leading to an increase in alkene selectivity with increasing reaction temperature. Therefore, the alkene yield can be maximized using a catalyst covered with dispersed monovanadate species, operated at high reaction temperature. The maximum operating temperature is determined by the onset of homogeneous reactions.

**CONCLUSIONS**

The rate and selectivity of ethane and propane ODH on VOₓ/Al₂O₃ catalysts depend on the vanadia surface densities and reaction temperature. Ethane ODH at zero conversion is ~75% selective to ethene and insensitive to vanadia surface density, while initial propene selectivities during propane ODH approach 90% at low vanadia surface densities, but decrease with increasing vanadia surface density. The ratio of $k_2/k_1$ is relatively independent of both reaction temperature and vanadia surface density for ethane ODH. The ratio of $k_2/k_1$ is independent of reaction temperature, but increases with increasing vanadia surface density for propane ODH. However, $k_3/k_1$ is generally higher for propane than for ethane ODH, indicating that the propene is more reactive relative to propane than ethene is to ethane. The activation energies for ethane and propane ODH, $E_1$, and ethane and propane combustion, $E_2$, are similar in magnitude at ~115 ± 20 kJ/mol and show weak dependence on vanadia surface density. The propene formation rate is approximately eight times larger than the ethene formation rate at 663 K, suggesting that the stabilization energy for ethoxide species may be higher relative to isopropoxide in the activated state. The apparent activation energy for alkene combustion, $E_3$, is ~60–90 kJ/mol and increases with vanadia surface density, leading to a decrease in $\Delta E_{13}$ with increasing vanadia surface density. The trends in $E_3$ and $\Delta E_{13}$ appear to be caused by decreasingly exothermic alkene adsorption as vanadia surface density increases. Alkene selectivity generally increases with increasing reaction temperature and with decreasing vanadia surface density. Therefore, the highest olefin yields are obtained on catalysts containing isolated monovanadates and operated at the highest possible reaction temperature consistent with the avoidance of homogeneous reactions.

**ACKNOWLEDGMENT**

This work was supported by the Director, Office of Basic Energy Sciences, Chemical Sciences Division of the U.S. Department of Energy under Contract DE-AC03-76SF00098.
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