Transhydrogenation of pentane and 1-hexyne over $\text{CrO}_x/\text{Al}_2\text{O}_3$ and potassium-doped $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalysts

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Abstract

The transhydrogenation of pentane (P) and 1-hexyne (1HY) was investigated over 4% $\text{CrO}_x/\text{Al}_2\text{O}_3$ and potassium-doped 4% $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalysts over a range of temperatures (523–773 K) with a 5:1 P:1HY ratio. Over the $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalyst, transhydrogenation clearly occurred at temperatures below 625 K where the yield of alkenes was higher for the co-fed system than for a combination of the individual yields. Due to the acidic nature of the alumina, many of the products were alkylated olefins and alkylated hydrocarbons formed by coincident alkylation and isomerisation. When pentane was added to a feed containing 1-hexyne, the extent of carbon deposition was reduced. By comparing transhydrogenation to limited hydrogen 1-hexyne hydrogenation at 623 K, it was shown that the processes of hydrogenation and transhydrogenation were different, with hydrogenation favouring alkanes, while transhydrogenation favoured alkenes. This may be because pentane dehydrogenation only releases two hydrogen atoms, which only allows 1-hexyne to hydrogenate to 1-hexene. Therefore, if the rate of alkene isomerisation and desorption is faster than that of pentane dehydrogenation, only alkenes will be observed. The latter proposal would suggest that the dehydrogenation/hydrogenation process is closely coupled and would be consistent with pentane influencing 1-hexyne surface chemistry. The effect of the potassium doping was to increase the yield of alkenes. The reason for this may be related to changes in the nature of the surface chromia species. The potassium also neutralised the acid sites on the alumina, reducing the extent of alkylation and hydrogenolysis, which suppressed the formation of other alkynes in the product mix.

Keywords Transhydrogenation · Dehydrogenation · Pentane · 1-Hexyne · $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalyst

Introduction

The production of olefins has been of major industrial importance since World War II using catalysts to yield high octane aviation fuel [1]. Since then, the petrochemical industries have been constantly developing new processes for improved olefin production, involving new catalyst formulations and modifications to reactor and plant design. This effort has also been extended to accommodate different hydrocarbon feeds in order to maximise production [2]. Currently, olefins are

produced via a number of processes in oil refining industries to successfully maximise the production. For example in the late 1980s, the CatofinTM process was designed for on-demand production of propylene and isobutylene using propane and isobutane, respectively [3–5].

The primary purpose of the transhydrogenation process historically has been to convert low-value cracked hydrocarbons into valued distillate products. Although transhydrogenation is not a new technology for production of olefins [6–11], there is scant scientific attention towards the invention, even though the invention has the potential to lift the equilibrium limitation of direct alkane dehydrogenation for the production of olefins by constantly removing the hydrogen produced through hydrogenation of an alkyne or alkadiene [12–14]. Hence, transhydrogenation combines the two processes (alkane dehydrogenation and alkyne hydrogenation) over a catalyst to form two alkenes. Alkane dehydrogenation is endothermic (typically around +125 kJ mol⁻¹) while alkyne/alkadiene hydrogenation is

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exothermic (typically around -165 kJ mol^{-1}); therefore, by coupling the endothermic dehydrogenation process with the exothermic hydrogenation process it is possible to generate a process, where the reaction conditions may be adjusted in order to produce a reaction that is net endothermic, net exothermic or thermally stable, which can simplify and reduce the cost involved in the process [11, 15]. For example, the transhydrogenation process between pentane and 1-hexyne is thermodynamically favoured at most temperatures.

The reaction sequence is thought to occur through dehydrogenation followed by hydrogenation with the hydrogen produced during the dehydrogenation used in the subsequent hydrogenation process:

At high temperatures, however, secondary reactions such as cracking and carbon deposition are also favoured; therefore, the reaction tends to get less selective as temperature is increased. The extent and nature of the carbon deposit will be related to the reactant and reaction temperature used.

The recent literature has revealed transhydrogenation reactions on various catalysts such as supported vanadia, chromia and doped chromia, most of which involved the use of LPG hydrocarbons. Jackson et al. [12–14] studied transhydrogenation of propane and propyne over chromia/alumina in a 1:1 ratio. The transhydrogenation reaction was shown to produce propene above the equilibrium value expected from propane dehydrogenation at 823 K. By 873 K, the conversion of propane was $\sim 80\%$, but the yield of propene was low, which was suggested to be due to secondary reaction of the propene forming methane and carbon deposits. Transhydrogenation was also performed between propyne and butane over a 1% vanadia/ θ -alumina catalyst at 823 K. Propyne and butane were co-fed, which resulted in an increase in conversion of propyne to propene compared to when it was fed singly over the catalyst. The transhydrogenation reaction was observed to deliver a $\sim 72\%$ increase in propene yield, while all the butane reacted was converted to butene isomers.

In this paper, the activity of a chromium oxide/alumina catalyst and a potassium-doped chromium oxide/alumina catalyst for the transhydrogenation of pentane/1-hexyne was evaluated primarily for the production of olefins and other valuable products. Most of the previous work has centred on the C-3 system, but in this work we were interested to see if transhydrogenation was applicable to a wider range of hydrocarbons. The C-6 hydrocarbons allowed us to study

the behaviour of conjugated (2,4-hexadiene) and non-conjugated hexadiene (1,5-hexadiene) isomers in comparison with 1-hexyne. The information on these systems will be reported in a subsequent paper. Pentane was chosen as the alkane as it was also a liquid hydrocarbon and in principle could make a full range of unsaturated species. The ratio of alkane:alkyne was chosen as $\sim 5:1$. Due to the stronger adsorption of the alkyne relative to the alkane, it was decided to have an excess of alkane. The specific ratio was chosen for experimental convenience. A wide temperature range was chosen to fully probe the transhydrogenation reaction. At temperatures below 600 K, pentane dehydrogenation has an equilibrium position that gives $< 1\%$ pentene, whereas by 773 K the equilibrium value is $\sim 20\%$. This would allow us to determine if pentane dehydrogenation was a controlling factor in transhydrogenation as the yields should follow the dehydrogenation equilibrium, even though the overall reaction is not equilibrium limited under these conditions.

Experimental

A 4% chromium oxide/alumina catalyst was prepared by incipient wetness impregnation using a γ -alumina support. The alumina support was pre-dried at 353 K for 16 h prior to impregnation. An aqueous solution of $[\text{NH}_4]_2\text{Cr}_2\text{O}_7$ (99+% Aldrich) was used to prepare the catalyst with a 4% w/w loading. The γ -alumina support (surface area $208 \text{ m}^2 \text{ g}^{-1}$, pore volume 0.52 ml g^{-1}) was supplied by Johnson Matthey, UK. After impregnation, the catalyst was dried for 2 h at 353 K and then at 393 K overnight. A potassium-doped 4% chromium oxide/alumina catalyst was prepared by incipient wetness impregnation using the above catalyst. The catalyst was treated with a solution of KOH sufficient to generate a 10% loading of potassium. This was then dried at 353 K for 2 h and then at 393 K overnight. Finally, both samples were calcined at 873 K for 6 h. After calcination, the samples were ground using a mortar and pestle and sieved to a uniform particle size of 250–425 μm .

Thermo-gravimetric analysis was performed on the catalysts using a combined TGA/DSC SDT Q600 thermal analyser coupled to an ESS evolution mass spectrometer for evolved gas analysis. The samples (~ 10 – 30 mg) were heated from ambient temperature to 1273 K using a heating ramp of 15 K min^{-1} in 2% H_2/N_2 or 2% O_2/Ar . Relevant mass fragments were followed by online mass spectrometry.

Elemental analysis was carried out using PerkinElmer analyst 400, atomic absorption spectrometer, with winlab 32. Typically, 0.1 g of catalyst was accurately measured to 0.001 g and digested with *aqua regia* reagent (1:3 HNO_3/HCl). CHN analysis was performed by combustion using a CE-440 elemental analyser.

Colorimetric analysis was conducted to determine the oxidation state of the chromium on the catalyst (dry catalyst, un-calcined, calcined and the reduced catalyst). Cr(VI) reacts with diphenylcarbazide in acid solution to produce a red–violet colour of unknown composition which can be measured at 540 nm. The procedure is nearly specific for Cr(VI). The acid range is critical such that the final acidity must be between 0.7 and 1.3 M H⁺. The process was calibrated by preparing chromium standards before samples containing no more than 25 µg Cr were pipetted into a 50-ml volumetric flask, diluted to approximately 40 ml and mixed well. To this, 5 ml 0.5 M sulphuric acid and 1 ml diphenylcarbazide solution were added, made up to the mark and mixed well. This was then allowed to stand 5–10 min for colour development and the absorbance measured at 540 nm.

The activity and selectivity of the catalyst were determined using an all-glass, atmospheric, flow, microreactor (Fig. 1). The catalyst (0.5 g) was reduced in situ with pure hydrogen (40 ml min^{−1}) by heating to 823 K and holding at this temperature for 2 h. After reduction was completed, the flow was switched to argon (40 ml min^{−1}) and the system purged for 30 min.

Pentane (99+% Aldrich) and 1-hexyne (97+% Aldrich) were introduced to the argon carrier gas (30 ml min^{−1}) using temperature controlled vaporisers. A molar feed ratio of 5.3:1 for the pentane:1-hexyne system was obtained using a temperature of 273 K for each vaporiser. The gas was then put through a static mixer before continuing to the reactor. The reaction was studied with each component individually and as a mixed feed, in the temperature range 523–773 K at atmospheric pressure. No hydrogen was co-fed in the process. The

reactor temperature was controlled using a West controller and the reaction temperature monitored via a thermocouple placed in the catalyst bed. The weight hourly space velocity (WHSV) was 2.8 h^{−1} for pentane and 0.6 h^{−1} for 1-hexyne. The reaction products were condensed and collected in heptane at 253 K and then analysed by gas chromatography (Agilent series, FID detector) fitted with a 150-m Petrocol column. An online ESS mass spectrometer was also coupled to the reactor to continuously follow any non-condensed eluted products over the 2-h time period of the reactions. The conversion for each reactant feed was calculated as:

$$\frac{\text{moles of reactant converted}}{\text{total moles fed}} \times 100$$

yield as:

$$P \text{ alone} = \frac{\text{moles of C for each product formed}}{\text{Total moles of C for P fed}} \times 100$$

$$1\text{HY alone} = \frac{\text{moles of C for each product formed}}{\text{Total moles of C for 1HY fed}} \times 100$$

$$P + 1\text{HY} = \frac{\text{moles of C for each product formed}}{\text{Total moles of C for [P + 1HY] fed}} \times 100$$

Results

The alumina support and both catalysts (chromia/alumina and K-doped chromia/alumina) were analysed by XRD, and only the pattern for alumina was observed (Supplementary

Fig. 1 Block schematic of flow microreactor

information). No chromia species were detected. This was expected as previous studies [16–18] have shown that crystalline chromium oxide is not observed at these loadings.

In order to determine the efficacy of the transhydrogenation process, the chromium oxide/alumina and potassium-doped chromium oxide/alumina catalysts were subjected to pentane and 1-hexyne individually and as a molar mixture of pentane:1-hexyne in the ratio (5.3:1) over the temperature range 523–773 K. The reactions were initially performed with the alumina support to evaluate its effect on the various reactants. The results were mainly cracking and alkylation products with ~50% and ~1% conversion at 623 K for the 1-hexyne and pentane, respectively. There was only a slight difference observed when 1-hexyne and pentane were co-fed over the alumina, with conversion of 1-hexyne still ~50%, but the pentane conversion was ~2% showing an increase of 1% compared to when it was run alone. The product distribution was very similar to that seen when 1-hexyne was run alone. This behaviour of alumina has been reported elsewhere in the literature [13, 14].

CrO_x/Al₂O₃ catalyst

A standard TGA-TPR from ambient temperature to 1273 K on the chromium oxide/alumina gave a single weight loss/hydrogen uptake centred at ~650 K and is presented in Fig. 2.

This is in keeping with the literature where reduction was detected at 643 K [19]. The catalyst was analysed by AAS analysis and colorimetric analysis to determine the amounts of Cr³⁺ and Cr⁶⁺ in the catalyst at various stages in the preparation. The results are presented in Table 1. Appreciable amounts of both Cr(III) and Cr(VI) oxidation states were observed at various treatment stages of the catalyst preparation. As expected, the dried catalyst contained a high percentage of Cr(VI) (3.21% c.f. the total loading 3.32%), while the reduced catalyst contained the lowest percentage of Cr(VI) at 0.22%.

Table 1 Percentage of Cr(III) and Cr(VI) during various catalyst treatments

Catalyst sample	Total %Cr	%Cr(III)	%Cr(VI)
As prepared ^a	3.32	0.11	3.21
Dried at 393 K	3.21	1.21	2.00
Calcined at 873 K	2.92	1.20	1.72
Reduced at 823 K	3.31	3.09	0.22

^a Air-dried before the oven overnight drying

These results confirm that the chromium in the catalyst has a mixed oxidation state even after reduction; therefore, the catalysts have been designated CrO_x/alumina (and K-CrO_x/alumina) to signify that the chromium species are non-stoichiometric oxides. This is in keeping with the literature [16, 18, 19].

The reactant conversions were followed during the individually fed and co-fed reactions. An increase in the conversion of pentane was observed at all temperatures when pentane and 1-hexyne were co-fed compared to pentane dehydrogenation, while for 1-hexyne the conversion at the higher temperatures (> 573 K) showed a decrease when pentane was co-fed relative to 1-hexyne with no added pentane. The results are presented in Fig. 3.

The product yields were calculated and are presented in Figs. 4, 5, 6, 7 and 8 (detailed analysis of the products is reported in Supplementary information, Tables S1–S5). As the temperature was decreased, the yield of olefin also decreased; however, the selectivity to olefins, which decreased as temperature decreased from 773 to 623 K, subsequently increased at the lower temperatures (573 K and 523 K).

Gaseous products were analysed by online mass spectrometry. In all cases, the evolution of hydrogen was observed at the start of the reaction but later declined, and after ~40 min no hydrogen was detected. Methane and ethene were observed to co-evolve with hydrogen but did

Fig. 2 TGA-TPR of chromium oxide/alumina catalyst

Fig. 3 Conversion comparison of pentane (P), 1-hexyne (1HY) and a 5.3:1 P:1HY mixture over $\text{CrO}_x/\text{Al}_2\text{O}_3$

Fig. 4 Yield of products at 773 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over $\text{CrO}_x/\text{alumina}$

Fig. 5 Yield of products at 673 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne over $\text{CrO}_x/\text{alumina}$

Fig. 6 Yield of products at 623 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne over $\text{CrO}_x/\text{alumina}$

Fig. 7 Yield of products at 573 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne over $\text{CrO}_x/\text{alumina}$

not decline and were maintained to the end of the reaction. This was observed for reaction temperatures between 773 and 623 K, while only pulses of these gases were observed with 573 and 523 K temperatures (shown in Supplementary material, Figs. S1–S3). The results obtained at 773 K are presented in Figs. 9, 10 and 11.

After use, the catalysts were analysed to determine whether there had been any carbon laydown. The results are shown in Fig. 12, where it can be seen that the co-feed has a lower carbon deposition relative to 1-hexyne.

Hydrogenation of 1-hexyne was also performed to determine to compare the catalyst efficacy between hydrogenation and transhydrogenation. The reaction was performed using a limited hydrogen feed (2% H_2/N_2) in the ratio 2:1 hexyne:2% H_2/N_2 . The results are shown in Fig. 13. Although there are

Fig. 8 Yield of products at 523 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne over $\text{CrO}_x/\text{alumina}$

some similarities in the yield of alkynes and alkadienes relative to transhydrogenation, there are significant differences, e.g. the yield of alkanes. Clearly, even in a hydrogen lean environment direct hydrogenation gives a significant alkane yield. This result may suggest that the hydrogen transfer in transhydrogenation does not simply go via an adsorbed hydrogen species.

K- $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalyst

The catalyst was characterised by XRD and TPR. No XRD reflections associated with chromia were detected. TPR (shown in Supplementary material) gave hydrogen uptakes

at 540 K and 760 K, which are consistent with the literature [20].

In general, increased conversions were obtained with pentane over the K-doped chromia catalyst compared to the non-doped catalyst. This was most marked with the pentane/1-hexyne co-feed, where higher conversions were obtained at all reaction temperatures compared to conversions obtained using the chromia catalyst. The results are presented in Fig. 14. The 1-hexyne conversions change very little between single feed and pentane co-feed.

The product distributions obtained over the doped catalyst had significant differences from those observed over the chromia catalyst. For example, 3-methylpentyne, one of the major products formed over the chromia catalyst, was not observed, and instead, an increased yield of 3-methyl-1-hexene and traces of benzene were observed. The distribution of the products (selectivity) changes with temperature and the % yield observed for individual products was also temperature dependent. The results are presented in Figs. 15, 16, 17, 18 and 19 (full detailed analysis is shown in Supplementary material, Tables S6–S10).

The evolved gases observed with the doped catalyst are the same (H_2 , CH_4 and C_2H_4) as observed over the chromia catalyst. However, with the pentane feed (Fig. 20), C_2H_4 was the major evolved gas. It was initially observed at ~10 min of the reaction, but later completely declined by ~60 min. Methane followed a similar pattern but with a much lower yield. This is in contrast to the gas evolution from the chromia catalyst. Hydrogen evolution was at a very low level, considerably reduced from that seen over the chromia catalyst. The results from 723 K are presented in Figs. 20, 21 and 22.

Fig. 9 Profile of the evolved gases over $\text{CrO}_x/\text{Al}_2\text{O}_3$ using pentane only at 773 K

Fig. 10 Profile of the evolved gases over $\text{CrO}_x/\text{Al}_2\text{O}_3$ using 1-hexyne only at 773 K

Fig. 11 Profile of the evolved gases over $\text{CrO}_x/\text{Al}_2\text{O}_3$ using P/1HY at 773 K

After use, the catalysts were analysed to determine whether there had been any carbon laydown. The results are shown in Fig. 23, where it can be seen that overall the extent of carbon deposition is less than that found with the non-doped catalyst. The general trend is also similar in that the co-feed has a lower carbon deposition relative to 1-hexyne except at 523 K where the opposite is true.

Discussion

$\text{CrO}_x/\text{alumina}$

On the basis of the characterisation results, the $\text{CrO}_x/\text{alumina}$ catalyst after reduction is similar to other chromia

Fig. 12 Carbon laydown on the spent $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalysts as a function of temperature

Fig. 13 Comparison of hydrogenation and transhydrogenation $\text{CrO}_x/\text{alumina}$ at 623 K

Fig. 14 Conversion comparison of P, 1HY and P/1HY mixed feeds over $\text{K-CrO}_x/\text{Al}_2\text{O}_3$

Fig. 15 Yield of products at 773 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over $\text{K-CrO}_x/\text{alumina}$

Fig. 16 Yield of products at 723 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over $\text{K-CrO}_x/\text{alumina}$

Fig. 17 Yield of products at 623 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over $\text{K-CrO}_x/\text{alumina}$

catalysts used for dehydrogenation reported in the literature [13, 16–19, 21]. The amount of Cr(VI) found in the sample is consistent with the literature for similar catalysts

Fig. 18 Yield of products at 573 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over K-CrO_x/alumina

Fig. 19 Yield of products at 523 K with feeds of pentane, hexyne and 5.3:1 pentane:hexyne feed over K-CrO_x/alumina

Fig. 20 Profile of the evolved gases over K-CrO_x/Al₂O₃ using P only at 723 K

[16–19] as is the absence of any XRD detectable crystalline chromium oxide. At this loading of chromia, a mixture of surface species such as dimers and polymers of various lengths have been shown to be present by Raman spectroscopy [21].

From the conversion data presented in Fig. 3 and the yield data presented in Figs. 4, 5, 6, 7 and 8, it is clear that when pentane and 1-hexyne are co-fed, they each alter the others reaction chemistry. Behaviour such as this has been observed previously with reactions between alkenes and alkynes [22] but not between alkane and alkyne. Examining the results when pentane was fed over the catalyst, the yield of olefins drops rapidly with temperature as expected from thermodynamics. At high temperatures (> 773 K), pentane can dehydrogenate to give pentenes; however, at lower temperatures the equilibrium position is such that only trace levels of olefin can be formed (~0.8% conversion at 600 K). Indeed, below 623 K pentane reactivity is very low. There is no such limitation for the hydrogenation of 1-hexyne to hexenes, but there must be a source of hydrogen. When 1-hexyne was passed over the catalyst at 773 K, a significant yield of alkene was observed and this can be associated with molecular breakdown of 1-hexyne, depositing carbon and liberating hydrogen that can be available for hydrogenation of other hexyne molecules. At lower temperatures, the extent of olefin formation from 1-hexyne is much less as carbon deposition decreases and the nature of the deposit will be hydrocarbonaceous, limiting available hydrogen. Therefore, it is easier to detect transhydrogenation at lower temperatures when other routes to olefins are less favoured. It should be remembered that reactions such

Fig. 21 Profile of the evolved gases over K-CrO_x/Al₂O₃ using 1HY only at 723 K

Fig. 22 Profile of the evolved gases over K-CrO_x/Al₂O₃ using P/1HY at 723 K

Fig. 23 Carbon laydown on the spent K-CrO_x/Al₂O₃ catalyst as a function of temperature

Fig. 24 Alkene selectivity in C₅–C₇ products from co-feed of pentane and 1-hexyne

as alkylation and isomerisation, as well as dehydrogenation and hydrogenation, are also taking place over the catalyst. The effect of transhydrogenation can be seen by examining the selectivity to alkenes (Fig. 24).

Over the CrO_x/alumina catalyst, the alkene selectivity and yield increase between 623 and 773 K as would be expected from thermodynamics (and kinetics). However, the alkene selectivity also increases as the temperature is decreased from 623 to 523 K. Indeed, the yield of alkenes is slightly greater at 573 K than 623 K (Figs. 6, 7). At 573 K, the yield of *n*-C₆ alkenes is over seven times the yield obtained from 1-hexyne alone. This enhancement in alkene yield can be assigned to transhydrogenation between pentane and 1-hexyne, which has no thermodynamic limitations at these temperatures ($\Delta G = -47 \text{ kJ mol}^{-1}$). However, as stated above, transhydrogenation is not the only reaction taking place. Although the total alkene yield is observed to have generally increased during the transhydrogenation process, most of the alkenes are isomers of the primary target products as well as alkylated alkenes. Indeed, many of the alkylated products are alkenes, which suggest that in essence, rapid isomerisation and alkylation are occurring coincidentally with the transhydrogenation process, e.g. 3-methylpentene and 1,4-hexadiene (both isomers of 1-hexyne) are common significant products from both the 1-hexyne feed and pentane/1-hexyne co-feed (Supplementary information). It is also noticeable that there is very little alkane formation.

Gas phase products are also produced with all three reaction feeds: pentane, 1-hexyne and pentane/1-hexyne mix (Figs. 9, 10, 11). Only hydrogen, methane and ethene were detected no C₃ or C₄ gases were observed. These three gas phase products are typical of cracking and hydrogenolysis. It is noticeable that the hydrogen evolution is not maintained, suggesting that the sites responsible for this reaction are deactivated during the course of the reaction. The results

presented in Fig. 12 confirm that there is carbon deposition, which may result in the loss of activity. It is noticeable that the hydrogen evolution maximises at different times when 1-hexyne is the feed and when there is a pentane/1-hexyne co-feed. The co-feed evolution maximises ~ 10 min, whereas the 1-hexyne feed gives a maximum hydrogen evolution at ~ 20 min. The pentane feed system shows a broad evolution of hydrogen that would be consistent with an evolution at ~ 10 min and ~ 20 min. This suggests that there may be two sites with 1-hexyne only able to access one, whereas pentane may access both. This would invoke competitive adsorption, which would be a route by which pentane may influence 1-hexyne surface chemistry.

To better understand the transhydrogenation process, a limited hydrogen hydrogenation of 1-hexyne was performed over the $\text{CrO}_x/\text{alumina}$ catalyst (Fig. 13) and it is clear that hydrogenation and transhydrogenation are not similar processes. Hydrogenation, even with a limited hydrogen feed, produces a significant yield of n-alkanes. This catalyst is not optimised for alkene selectivity from hydrogenation and so produces the thermodynamically favoured alkane. In contrast, virtually no n-alkane is produced from the transhydrogenation reaction, and even the yield of iso-alkane is lowest during transhydrogenation, suggesting that the hydrogen transfer process is much more controlled than in hydrogenation. This has mechanistic implications for our understanding of transhydrogenation. The hydrogenation reaction reveals that if atomic hydrogen is present on the surface, then alkane is the favoured product. For the transhydrogenation reaction to favour alkenes, it must control the hydrogen supply or there are two types of hydrogen on the surface. It has been shown using isotopic labelling that different hydrogen populations do exist on the surface [13]. Therefore, the difference between transhydrogenation and hydrogenation may relate to 1-hexyne sampling different hydrogen populations. The other possibility is that as pentane dehydrogenation only releases two hydrogen atoms, this only allows 1-hexyne to hydrogenate to 1-hexene. Therefore, if the rate of alkene isomerisation and desorption is faster than that of pentane dehydrogenation, only alkenes will be observed as there is never free hydrogen for further hydrogenation.

K-CrO_x/alumina

The addition of potassium to catalysts is a standard method to change activity, change selectivity and reduce by-product formation [23]. Detailed spectroscopic and mechanistic analysis of $\text{CrO}_x/\text{alumina}$ has shown that the addition of potassium does not affect alkane dehydrogenation but significantly affects side reactions [21]. Therefore, the addition of potassium has a noticeable effect on product selectivity. For example, when pentane and 1-hexyne are co-fed at 623 K, the yield of alkenes

was 30%, whereas over the $\text{CrO}_x/\text{alumina}$ catalyst under the same conditions the alkene yield was only 9%. Most strikingly, no alkynes are formed over the K-CrO_x/alumina, specifically 3-methylpentyn, which was a major product via structural isomerisation over $\text{CrO}_x/\text{alumina}$. Much of this change can be understood as a neutralisation of the acid sites on the alumina; indeed, it has been shown in a number of reports that the role of potassium is to remove acid sites [20, 21, 24]. Nevertheless, there is still considerable alkylation with significant yields of iso-alkanes (including methylcyclohexane) although this could be base catalysed [25, 26]. Alkylated alkenes such as 3-methyl-2-pentene and 3-methyl-1-hexene were also major products. Figure 24 shows the alkene selectivity as a function of temperature, and it can be seen that the alkene selectivity does not significantly change with temperature over the K-CrO_x/alumina in contrast to the $\text{CrO}_x/\text{alumina}$ catalyst. This lack of variation in alkene selectivity across the temperature range would be consistent with transhydrogenation and is consistent with the detailed product analysis (Supplementary information), where hexene isomers are specifically enhanced. It is also clear that the K-CrO_x/alumina catalyst gives an enhanced yield of alkenes compared to $\text{CrO}_x/\text{alumina}$. A possible explanation for this difference may be related to the fact that addition of potassium to $\text{CrO}_x/\text{alumina}$ is known to change the nature of the surface chromia species [20]. In $\text{CrO}_x/\text{alumina}$, at least five surface chromia species have been identified [16]; when potassium is added to a $\text{CrO}_x/\text{alumina}$, the surface chromia species alter tending to form isolated and oligomeric chromia [20]. It is possible that these species facilitate transhydrogenation giving an increased alkene yield.

As with the $\text{CrO}_x/\text{alumina}$ catalyst, light gases were evolved from the reaction of the three feeds over K-CrO_x/alumina (Figs. 20, 21, 22). However, the manner of their evolution is markedly different. For all three feeds, methane and ethene are evolved as pulses, typically over 0.3 h, whereas hydrogen is evolved over most of the time on stream. However, when the gas evolutions are examined in more detail, further differences between the two catalysts are found. For all the feeds, the major gas produced is ethene. This is in contrast to the gas evolution over the $\text{CrO}_x/\text{alumina}$, where hydrogen and ethene were always the main gaseous products. The short pulse of the hydrocarbon gases produced over the K-CrO_x/alumina suggests that a small number of sites catalysing cracking reactions such as

were rapidly deactivated. The extent of carbon deposition over the K-CrO_x/alumina catalyst is noticeably less than over the $\text{CrO}_x/\text{alumina}$ (Fig. 23). This is in keeping with the

literature [12, 18, 19] where potassium doping reduces the extent of carbon deposition. The pentane/1-hexyne co-feed shows lower carbon laydown than expected from a sum of each species individually, except for the reaction at 523 K where the extent of carbon laydown is similar to the sum of the individual feeds. Interestingly, if the effect of pentane on 1-hexyne surface chemistry was due to competitive adsorption, it may be expected that at lower temperatures where the adsorption of pentane is limited, the effect would be less and this would explain why at 523 K the carbon deposition is purely the sum of the pentane deposition and 1-hexyne deposition.

Conclusions

The transhydrogenation of pentane and 1-hexyne was investigated over $\text{CrO}_x/\text{Al}_2\text{O}_3$ and $\text{K-CrO}_x/\text{Al}_2\text{O}_3$ catalysts. Over the $\text{CrO}_x/\text{Al}_2\text{O}_3$ catalyst, transhydrogenation clearly occurred at temperatures below 625 K, where the yield of alkenes was higher for the co-fed system than for a combination of the individual yields. Due to the acidic nature of the alumina, many of the products were alkylated olefins and alkylated hydrocarbons due to coincident alkylation and isomerisation. Pentane was shown to affect the chemistry of 1-hexyne over and above the transhydrogenation reaction, e.g. the extent of carbon deposition was reduced. By comparing transhydrogenation to limited hydrogen 1-hexyne hydrogenation at 623 K, it was shown that the processes of hydrogenation and transhydrogenation were different, with hydrogenation favouring alkanes, while transhydrogenation favoured alkenes. It was suggested that this may be due to different hydrogen pools that influence the final product or that as pentane dehydrogenation only releases two hydrogen atoms, this only allows 1-hexyne to hydrogenate to 1-hexene. Therefore, if the rate of alkene isomerisation and desorption is faster than that of pentane dehydrogenation, only alkenes will be observed. The latter proposal would suggest that the dehydrogenation/hydrogenation process is closely coupled and would be consistent with pentane influencing 1-hexyne surface chemistry. The effect of potassium doping was to increase the yield of alkenes. The reason for this may be related to changes in the nature of the surface chromia species suggesting site specificity for transhydrogenation. The potassium also neutralised the acid sites on the alumina, reducing the extent of alkylation and hydrogenolysis, which suppressed the formation of other alkynes in the product mix.

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