Catalytic Conversion of Anisole: Evidence of Oxygen Removal in Reactions with Hydrogen

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Abstract The conversion of anisole, a prototypical bio-oil compound, was catalyzed by Pt/Al₂O₃ in the presence of H₂ at 573 K, with the selectivity for C–O bond breaking approximately matching that for ring hydrogenation; these reactions were accompanied by methyl group transfers matching those in the conversion catalyzed by HY zeolite.

Keywords Anisole · Zeolite Y · Supported platinum · Transalkylation · Hydrodeoxygenation

1 Introduction

Plant biomass is a potentially valuable renewable source of fuels and chemicals. Lignocellulose can be rapidly transformed into liquids by pyrolysis, giving “bio-oil,” but the high oxygen content leads to immiscibility with fossil fuels, instability, and corrosion [1–3]. Bio-olys typically contain several hundred compounds [4, 5], and an understanding of the reaction networks in their catalytic upgrading is limited; even information characterizing the upgrading of individual bio-oil compounds is scarce [6–10]. Our approach to begin unravelling this chemistry, and specifically to understand oxygen removal reactions, was to investigate the conversion of a bio-oil compound, anisole, chosen as a prototype because it has both an ether linkage and an aromatic ring. Reactions were catalyzed by a solid acid (HY zeolite, Zeolyst CBV 720) and separately by a catalyst consisting of a metal dispersed on an acidic support (Pt/Al₂O₃, Sigma–Aldrich, 1% wt Pt). The goal of the research was to develop detailed reaction networks, characterized by numerous connected pathways, for the conversion of anisole with catalysts representing these two important classes of catalysts.

2 Experimental

Catalytic reactions were carried out in a once-through packed-bed flow reactor under the following conditions, with liquid anisole (methoxybenzene, Sigma–Aldrich, 99.8%) vaporized into a flowing gas stream: catalyst mass 4.1–401.5 mg, temperature 573 K, pressure 140 kPa, anisole liquid flow rate 0.03 mL min⁻¹, gas feed flow rate 100 mL min⁻¹ (as pure N₂ in experiments with HY zeolite and as 30% H₂/70% N₂ in experiments with Pt/Al₂O₃). Each catalyst powder was diluted with particles of inert, nonporous α-Al₂O₃. The product gas stream was condensed at 285–288 K. Uncondensed gases and condensate samples collected periodically were analyzed by gas chromatography and gas chromatography/mass spectrometry.

3 Results

Dozens of products were observed in the conversions with each catalyst. Significant catalyst deactivation, characterized by a 50% reduction in conversion within 1 h on stream, was observed with HY zeolite; the Pt/Al₂O₃ catalyst working in the presence of H₂ was much more stable. Selectivities to major and minor products at low initial
Table 1: Products of conversion of anisole catalyzed by Pt/Al₂O₃ and by HY zeolite (liquid product streams)

| Product       | Selectivity to product in reaction catalyzed by Pt/Al₂O₃ | Selectivity to product in reaction catalyzed by HY zeolite |
|---------------|---------------------------------------------------------|----------------------------------------------------------|
| Phenol        | 0.65                                                    | 0.60                                                     |
| 2-Methylphenol| 0.15                                                    | 0.11                                                     |
| Benzene       | 0.044                                                   | –                                                        |
| Cyclohexanone | 0.033                                                   | –                                                        |
| 2,6-Dimethylphenol | 0.024                                               | 0.011                                                    |
| 2-Methylanisole | 0.0077                                            | 0.13                                                     |
| 4-Methylphenol | 0.0022                                                | 0.045                                                    |
| 4-Methylanisole | 0.0013                                              | 0.15                                                     |

Data were extrapolated to zero time on stream, and thus represent approximate initial selectivities determined at a weight hourly space velocity of 18 (g of reactant)/(g of catalyst * h), a pressure of 140 kPa, and a temperature of 573 K. Selectivity is defined as yield [mol product formed/mol of organic reactant fed]/conversion [mol of organic reactant consumed/mol of organic reactant fed]. In the experiments with the Pt/Al₂O₃ catalyst, the feed molar ratio of H₂ to the organic reactant was 2400.

Fig. 1: Selectivity for the formation of phenol (closed circles) and 2-methylphenol (open circles) in the conversion of anisole catalyzed by HY zeolite at 573 K. Data for each product were fitted with a straight line and extrapolated to zero conversion; intercepts of regression lines significantly different from zero selectivity at zero conversion (analyzed with 95% confidence limits) indicate primary products, in this case phenol, and those not significantly different from zero (analyzed with 95% confidence limits) are considered non-primary, in this case 2-methylphenol.

Conversions (0.14 for Pt/Al₂O₃ and 0.16 for HY zeolite) are summarized in Table 1; these data represent extrapolations to zero time on stream and therefore initial catalyst performance, prior to significant deactivation.

The most abundant products of the conversion catalyzed by HY zeolite were phenol, methylanisoles, and methylphenols; less abundant products included dimethylanisoles and dimethylphenols. The substitution at the 2- and 4-positions on the aromatic ring is consistent with the kinetically determined substitution preference of aromatic ethers and aromatic alcohols [11]. Trace products included those with additional methyl-group substitutions on the aromatic ring (observed with each catalyst) and hydrogenation products, such as cyclohexane, which were also formed with Pt/Al₂O₃ in the presence of H₂.

The most abundant products of the reactions catalyzed by Pt/Al₂O₃ were phenol, 2-methylphenol, and benzene; less abundant products included water, methane, methanol, 2-methylanisole, and 4-methylanisole. The appearance of cyclohexanone, for example, indicates hydrogenation of the aromatic ring [12].

Thus, an important class of reaction, observed for both catalysts, is methyl group transfer. Hydrogenolysis, hydrogenation, and hydrodeoxygenation were also observed in the conversion catalyzed by Pt/Al₂O₃ in the presence of H₂.

Selectivity versus conversion plots (e.g., Fig. 1) were used to identify products as primary or not [13, 14] (these designations are empirical, falling short, for example, of providing information about intermediates that were too reactive to be detected). The selectivity–conversion data observed for anisole conversion catalyzed by HY zeolite (Figs. 1, 2) indicate that phenol, 2-methylanisole, 4-methylanisole, and 2,6-dimethylphenol (data not shown) were primary products and that 2-methylphenol and 4-methylphenol (data not shown) were non-primary products. Benzene and cyclohexanone were not observed.

On the basis of the data showing that methyl group transfer (transalkylation) is the predominant class of reaction and a primary reaction in the formation of phenol, 2-methylanisole, and 4-methylanisole in the conversion catalyzed by HY zeolite, we infer the qualitative reaction network shown in Fig. 3. For simplicity, the source of the methyl group is not always shown in the reaction network.

The selectivity–conversion data observed for anisole conversion catalyzed by Pt/Al₂O₃ in the presence of H₂ indicate that phenol, 2-methylphenol, and 4-methylphenol (data not shown) were primary products and that benzene, cyclohexanone, 2- and 4-methylanisole, and 2,6-dimethylphenol (data not shown) were non-primary products.

Presuming that methyl group transfer, hydrodeoxygenation, and hydrogenation are the important reaction classes, and by recognizing which compounds were primary products, we inferred the reaction network of Fig. 6 for anisole conversion catalyzed by Pt/Al₂O₃ in the presence of H₂.

The data characterizing the conversion of anisole catalyzed by Pt/Al₂O₃ are represented satisfactorily by first-order kinetics (Fig. 7). The pseudo first-order rate constant for the overall disappearance of anisole was found to be...
19 L (g of catalyst · h) under the conditions stated in the caption of Fig. 6. Selectivities to benzene relative to cyclohexanone indicate that the rate of hydrogenolysis of the C–O bond approximately matched that of hydrogenation of the aromatic ring at low conversions and at the low pressure of these experiments.

**Fig. 2** Selectivity for the formation of 2-methylanisole (closed circles) and 4-methylanisole (open circles) in the conversion of anisole catalyzed by HY zeolite; conditions are stated in the text. Data for each product were fitted with a straight line and extrapolated to zero conversion; intercepts significantly different from zero selectivity at zero conversion (analyzed with 95% confidence limits) indicate primary products, in this case both 2-methylanisole and 4-methylanisole, and those not significantly different from zero are considered to be evidence of non-primary products.

**Fig. 3** Reaction network for the conversion of anisole catalyzed by HY zeolite at 573 K. Reactions are postulated on the basis of the products identified, including trace compounds, and their formation as a result of possible methyl group transfer reactions. Bold arrows show the reactions that are kinetically most significant (with the width of the arrow denoting a rough measure of rate; the wider the arrow, the faster the reaction).

**Fig. 4** Selectivity for the formation of several products in the conversion of anisole catalyzed by Pt/Al₂O₃ in the presence of H₂ at 573 K. Data for each product were fitted with a straight line and analyzed with 95% confidence limits as indicated in the caption of Fig. 1; primary products in this case are phenol (open circles) and 2-methylphenol (closed triangles); benzene (closed circles) is a non-primary product.

**Fig. 5** Selectivity for the formation of 2-methylanisole (closed circles) and cyclohexanone (open circles) in the conversion of anisole catalyzed by Pt/Al₂O₃; conditions are stated in the text. Data for each product were fitted with a straight line and extrapolated to zero conversion; intercepts significantly different from zero selectivity at zero conversion (analyzed with 95% confidence limits) indicate primary products, and those not significantly different from zero (analyzed with 95% confidence limits) are considered to be evidence of non-primary products, in this case 2-methylanisole and cyclohexanone.

**4 Discussion**

The reaction network of Fig. 3 accounts for major, minor, and trace products; it contains more detail than the reported reaction network proposed for conversion of anisole catalyzed by the zeolite HZSM-5 [6] and is broadly consistent with it. Individual compounds produced with HZSM-5 and not identified in our work include 3-methylphenol and...
methane, possibly indicating the higher reaction temperature (673 K) or higher inverse space velocity in the reported work.

Similarly, the reaction network of Fig. 6 accounts for the major, minor, and trace products in the reactions catalyzed by Pt/Al$_2$O$_3$, and, again, the detail markedly exceeds than of any comparable reaction network including catalytic hydrodeoxygenation. A comparison of Figs. 3, 6 shows that primary products formed in the presence of one catalyst (e.g., 2-methylphenol) are not necessarily primary products formed in the presence of the other catalyst. The low selectivity to benzene relative to that of products resulting from methyl group transfers suggests that higher concentrations of H$_2$ (and high pressures) will be required to achieve acceptable yields and selectivities for oxygen removal.

5 Conclusions

In the conversion of anisole catalyzed by Pt/Al$_2$O$_3$ and by HY zeolite, methyl group transfers were found to be kinetically significant reactions, and when the catalyst was Pt/Al$_2$O$_3$ used in the presence of H$_2$, the conversion was accompanied by products of C–O bond-breaking and hydrogenation reactions. Data such as those reported here provide a starting point for predicting the conversion of bio-oils, specifically for removal of oxygen and upgrading of fuel properties.

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