Catalytic Reactions of Guaiacol: Reaction Network and Evidence of Oxygen Removal in Reactions with Hydrogen

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Abstract The conversion of guaiacol, a prototypical compound representative of lignin-derived pyrolysis bio-oils, was catalyzed by Pt/Al$_2$O$_3$ in the presence of H$_2$ at 573 K. The conversion took place with a high selectivity for aromatic carbon–oxygen bond cleavage relative to the accompanying methyl group transfer reactions. This oxygen removal was not observed in the absence of H$_2$ as a co-reactant. Products that were formed by methyl-group transfer match those produced in the conversion catalyzed by zeolite HY, which was not active for oxygen removal reactions.

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

1 Introduction

Lignin constitutes as much as 30 wt% of lignocellulosic biomass [1, 2] and offers excellent potential as a feedstock for conversion to fuels and chemicals by pyrolysis, giving “bio-oils.” The results of earlier investigations [3–7] show that catalytic upgrading can be used to remove oxygen from these liquids to minimize the instability and corrosion issues. In comparison with the widely investigated cellulose-derived components of bio-oils (including carbohydrates) [8–10], the lignin-derived components have lacked attention, and there is limited understanding of the reaction networks and kinetics [11–13]. Our approach was to begin investigating the reaction network for a characteristic lignin-derived compound found in pyrolysis bio-oils—guaiacol. A specific goal was to elucidate the oxygen-removal reactions in the conversion in the presence of an important class of catalysts: a metal dispersed on an acidic support (Pt/Al$_2$O$_3$). Data were also collected with a solid acid catalyst (zeolite HY) to identify the reactions catalyzed by an acidic function (such as the Al$_2$O$_3$ support) and distinguish them from the reactions catalyzed by the metal.

2 Experimental

Powers of Pt/Al$_2$O$_3$ (1 wt% Pt, Sigma-Aldrich) and zeolite HY (Zeolyst CBV 720) were pre-treated in flowing streams of H$_2$/N$_2$ and N$_2$, respectively, at 573 K. Catalytic reactions were conducted in a once-through packed-bed flow reactor under the following conditions, with liquid reactant (99.5 wt% guaiacol) vaporized into a flowing gas stream (30% H$_2$/70% N$_2$ in experiments with Pt/Al$_2$O$_3$ and N$_2$ in experiments with HY zeolite): catalyst mass, 0.010–0.100 g (catalyst diluted with particles of inert, nonporous z-Al$_2$O$_3$); temperature, 573 K; pressure, 140 kPa; liquid reactant feed rate, 0.015 mL/min; and gas feed flow rate, 100 mL/min. 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.
Results

Dozens of products were detected and identified in the reactions observed with each catalyst. As in recent work on anisole [14], more rapid catalyst deactivation was observed with the zeolite (≈50% reduction in conversion within 1 h on stream) than with the Pt/Al₂O₃ catalyst operating in the presence of H₂ (Fig. 1). Selectivities to the most abundant products formed at low initial conversions (0.08 for Pt/Al₂O₃ and 0.11 for HY zeolite) are summarized in Table 1. Initial conversions, obtained by extrapolating conversion data to zero time on stream, were used to determine catalyst activities in the absence of catalyst deactivation. Because the conditions were chosen to allow determination of initial catalyst performance, only minimal amounts of coke were formed in our experiments; the changes in mass of our packed beds containing catalyst in the various experiments were less than 0.1%.

The predominant products of the reactions catalyzed by Pt/Al₂O₃ were phenol, benzene-1,2-diol (catechol), and 3-methylbenzene-1,2-diol (3-methylcatechol); less abundant products were water, methane, methanol, benzene, toluene, anisole, cyclohexanone, 2-methylphenol (o-cresol), 2-methoxy-3-methylphenol (3-methylguaiacol), 6-methylguaiacol, and 1,2-dimethoxybenzene. Several trace products were also detected, including cyclohexane, cyclohexene, methoxycyclohexane, xylenes, cyclohexanol, methylcyclohexanones, methylanisoles, 4-methylguaiacol, 5-methylguaiacol, dimethylenphenols, trimethylphenols, and dimethylhydroquinones. These products were presumably formed by hydrogenation, methyl group transfer (transalkylation, which could be intra- or intermolecular) and isomerization reactions.

In contrast, none of the deoxygenated products such as anisole and phenol was observed in the reactions catalyzed by the zeolite. The most abundant products observed with this catalyst were catechol, 1,2-dimethoxybenzene, 3-methylguaiacol, 4-methylguaiacol, 5-methylguaiacol, and 6-methylguaiacol; a less abundant product was 3-methylcatechol. Trace products included dimethoxytoluenes.

### Table 1 Products of conversion of guaiacol catalyzed by Pt/Al₂O₃ and by HY zeolite (liquid product streams)

| Product                  | Selectivity to product in reaction catalyzed by Pt/Al₂O₃ with H₂<sup>a</sup> | Selectivity to product in reaction catalyzed by HY zeolite<sup>b</sup> | Selectivity to product in reaction catalyzed by Pt/Al₂O₃ without H₂<sup>b</sup> |
|--------------------------|--------------------------------------------------------------------------|------------------------------------------------------------------|------------------------------------------------------------------|
| Benzene                  | 0.002                                                                    | –                                                                | –                                                                |
| Toluene                  | 0.0004                                                                   | –                                                                | –                                                                |
| Anisole                  | 0.009                                                                    | –                                                                | –                                                                |
| Cyclohexanone            | 0.026                                                                    | –                                                                | –                                                                |
| Phenol                   | 0.316                                                                    | –                                                                | –                                                                |
| o-Cresol                 | 0.031                                                                    | –                                                                | –                                                                |
| 1,2-Dimethoxybenzene     | 0.021                                                                    | 0.300                                                           | 0.242                                                           |
| 3-Methylguaiacol         | 0.012                                                                    | 0.015                                                           | 0.031                                                           |
| 4-Methylguaiacol         | Trace                                                                    | 0.071                                                           | Trace                                                           |
| 5-Methylguaiacol         | Trace                                                                    | 0.043                                                           | Trace                                                           |
| 6-Methylguaiacol         | 0.031                                                                    | 0.049                                                           | 0.105                                                           |
| Catechol                 | 0.412                                                                    | 0.49                                                            | 0.560                                                           |
| 3-Methylcatechol         | 0.125                                                                    | 0.032                                                           | 0.055                                                           |

Data were extrapolated to zero time on stream, and thus represent approximate initial selectivities determined at a weight hourly space velocity of 20 (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]

<sup>a</sup> 100 mL/min gas feed rate, 30% H₂/70% N₂; feed molar ratio of H₂ to the organic reactant was 10

<sup>b</sup> 100 mL/min gas feed rate, 100% N₂
4-methylcatechol, dimethylphenols, trimethylphenols, and tetramethylphenols. The traces of alkylated phenols were possibly produced from the impurity phenol in the feed guaiacol.

These data show that methyl-group transfer is an important class of reaction observed for both catalysts. Reactions with H₂ leading to oxygen removal as water or methanol (hydrodeoxygenation) and hydrogenation, on the other hand, were kinetically significant only with the Pt/Al₂O₃ catalyst used in the presence of H₂. In the absence of H₂, the conversion in the presence of this catalyst was nearly zero (<0.01 under our conditions, data not shown), and no deoxygenated products were observed. Then, as shown in Table 1, only products similar to those formed with the HY zeolite catalyst were observed.

To determine an approximate reaction network for the conversion of guaiacol catalyzed by Pt/Al₂O₃ in the presence of H₂, selectivity versus conversion plots (e.g., Fig. 2) and Figs. 3–5 are used to identify primary products and non-primary products.
were used to identify products as primary or not \[15, 16\] (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 guaiacol conversion catalyzed by Pt/Al\(_2\)O\(_3\) (Figs. 2, 3, 4, 5) in the presence of H\(_2\) indicate that anisole, phenol, catechol, 3-methylcatechol, 3-methylguaiacol, 6-methylguaiacol, and 1,2-dimethoxybenzene were primary products, whereas benzene, toluene, cyclohexanone (data not shown), and \(\alpha\)-cresol (data not shown) were non-primary products.

Presuming that transalkylation, hydrodeoxygenation, and hydrogenation are the important reaction classes, by recognizing which compounds were primary and which were non-primary products, and by using our chemical judgment of the most likely classes of reaction for formation of minor and trace products, we inferred the reaction network of Fig. 6 for guaiacol conversion catalyzed by Pt/Al\(_2\)O\(_3\) in the presence of H\(_2\).

### 4 Discussion

The reaction network of Fig. 6 accounts for major, minor, and trace products in the reactions catalyzed by Pt/Al\(_2\)O\(_3\); it contains more detail than previously reported reaction networks proposed for the conversion of guaiacol catalyzed by platinum or other supported metals. In most of these reports, no reaction networks were presented; the catalysts were Ru/C [17], Pd/C [17], Pt/ZrO\(_2\) [18], and V/Al\(_2\)O\(_3\) [19].

Our results are broadly consistent with these reports in terms of the classes of reactions observed, but different products are reported in our work because of the different catalysts and operating conditions. In the comparable work with the Pt/ZrO\(_2\) catalyst [18], the reported products not observed in our work include 2-methoxycyclohexanone, 1-methyl-1,2-cyclohexanediol, and 1,2-dimethoxycyclohexane; we suggest that the differences are associated with the higher pressure (8 MPa) and the higher H\(_2\) to guaiacol molar feed ratio (~25:1) in the reported work.

In the reactions catalyzed by Pt/Al\(_2\)O\(_3\), anisole and phenol were formed as primary products, presumably by hydrodeoxygenation as the C–O bond \(\alpha\) to the aromatic ring in guaiacol was broken. Higher selectivity for phenol formation implies that deoxygenation by removal of the methoxy group is more favorable than deoxygenation by removal of the hydroxyl group. The reaction network also suggests that unimolecular and bimolecular transalkylation reactions occur in parallel—as 3-methylcatechol and...
methylguaiacols are both primary products. These methylgroup transfers likely take place on the slightly acidic Al₂O₃ support. In the methylation of guaiacol, only 3-methylguaiacol and 6-methylguaiacol were formed in substantial yields (4-methylguaiacol and 5-methylguaiacol were observed only as trace products). We suggest that the high selectivities for the former products may indicate that guaiacol interacts with the Al₂O₃ support through the methoxy or hydroxy group to form methoxyphenate and hydroxyphenate species, respectively [20], facilitating methylation preferentially at the position next to these groups and leading to the formation of 3-methylguaiacol and 6-methylguaiacol.

Consistent with recent reports [11, 21], cyclohexanone was observed as a partially hydrogenated product of phenol conversion. Without H₂ in the feed, hydrodeoxygenation and hydrogenation routes, shown by the dotted lines in Fig. 6, were evidently minimized. The results emphasize the significance of H₂ for oxygen removal from aromatic oxygenates (such as guaiacol, phenol, and anisole) and suggest the need for high H₂ partial pressures in practical processes for processing of lignin-derived liquids.

Lacking the metal function, the HY zeolite catalyzed transalkylation as the only kinetically significant reactions, leading to products similar to those observed in the conversion catalyzed by the Pt/Al₂O₃ in the absence of H₂. This result is consistent with a report of the conversion of anisole catalyzed by HZSM-5 [13], and it highlights the importance of a metal function to catalyze hydrodeoxygenation.

5 Conclusions

The catalytic conversion of guaiacol catalyzed by Pt/Al₂O₃ in the presence of H₂ involves three major classes of reactions: hydrodeoxygenation, hydrogenation, and transalkylation. Selectivities to hydrodeoxygenation products were found to be comparable to selectivities to the accompanying transalkylation products. The acidic HY zeolite is not active for oxygen removal reactions. The data presented here provide a step towards predicting the conversion of lignin-derived bio-oils for removal of oxygen and upgrading of fuels.

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