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Development and mechanistic investigation of the manganese(III) salen-catalyzed dehydrogenation of alcohols†

Simone V. Samuelsen,a Carola Santilli,a Märten S. G. Ahlquistb and Robert Madsen*ab

The first example of a manganese(II) catalyst for the acceptorless dehydrogenation of alcohols is presented. N,N′-Bis(salicylidene)-1,2-cyclohexanediaminomanganese(III) chloride (2) has been shown to catalyze the direct synthesis of imines from a variety of alcohols and amines with the liberation of hydrogen gas. The mechanism has been investigated experimentally with labelled substrates and theoretically with DFT calculations. The results indicate a metal–ligand bifunctional pathway in which both imine groups in the salen ligand are first reduced to form a manganese(II) amido complex as the catalytically active species. Dehydrogenation of the alcohol then takes place by a stepwise outer-sphere hydrogen transfer generating a manganese(III) salan hydride from which hydrogen gas is released.

Introduction

Metal-catalyzed dehydrogenation of alcohols gives rise to aldehydes and ketones, which can be further transformed into imines, amides, esters, carboxylic acids and various heterocycles in the same pot.1 The acceptorless dehydrogenation constitutes an attractive synthetic protocol since it does not require any stoichiometric oxidants and only releases hydrogen gas as a co-product. The dehydrogenative transformations are usually catalyzed by complexes of the platinum-group metals such as ruthenium and iridium.2 Recently, however, non-noble metal complexes based on iron, cobalt and manganese have also been shown to catalyze alcohol dehydrogenations.3 Especially manganese-catalyzed dehydrogenations have been a hot research area since the first catalyst was introduced in 2016.3 Since then several research groups have presented different manganese complexes for preparing various functional groups and heterocyclic frameworks (Fig. 1).4 These complexes have also been used to catalyze the hydrogenation of carbonyl compounds.5 The significance of the discoveries is exemplified by the large number of works published on this topic over the past two years3–5 including several reviews.3e–d

Notably, the developed complexes are all manganese(i) compounds with CO ligands and a pincer ligand. The electron-withdrawing CO ligand is necessary for stabilizing the low oxidation state of manganese. The mechanism for the dehydrogenation with these complexes is believed to involve a catalytic cycle with different manganese(i) species3,4,5b,5d and no catalytic activity is observed without the CO ligands or with the corresponding manganese(II) dihalide complexes.3e Although manganese is an Earth-abundant and cheap metal, manganese(i) complexes are not inexpensive since they are prepared in several steps from Mn2(CO)10. This carbonyl complex is quite expensive due to its difficult preparation by a carbonylation reaction.7 As a result, there is a need for identifying a new and more abundantly available class of manganese complexes for alcohol dehydrogenations. Especially, it would be attractive to catalyze the dehydrogenations by higher valent complexes where stabilization by CO ligands is not necessary.

This prompted the question whether manganese(III) complexes would be able to catalyze the same acceptorless alcohol dehydrogenations? Manganese(III) complexes such as

![Fig. 1 Manganese(i) complexes for alcohol dehydrogenation](image-url)
the Jacobsen’s catalyst are widely known for catalyzing oxidation reactions in the presence of a stoichiometric oxidant. This includes epoxidation of olefins, oxidation of alcohols as well as hydroxylation and halogenation of alkanes. The reactions proceed through catalytic cycles with different manganese(III), (IV) and (V) species.

Herein, we describe the first example of an acceptorless alcohol dehydrogenation catalyzed by a manganese(III) complex. An easily available manganese(III) salen catalyst has been shown to liberate hydrogen gas from alcohols and the transformation has been applied to the synthesis of imines from alcohols and amines. The mechanism has been investigated by experimental and theoretical methods which indicates a metal–ligand bifunctional pathway for the removal of hydrogen gas from the alcohol.

Results and discussion

The transformation was discovered while attempting to develop a more convenient in situ-formed catalyst system (Table 1). Benzyl alcohol was treated with an equimolar amount of cyclohexylamine in the presence of Mn2(CO)10 and different ligands. The most promising result was obtained with N,N'-bis(salicylidene)ethylenediamine (H2salen) as the ligand (entry 1). Since this is a common ligand for manganese(III) complexes, an experiment was also performed with the Jacobsen’s catalyst (1, Fig. 2). Interestingly, this increased the yield to 79% with some unreacted benzyl alcohol remaining. This observation shows that the acceptorless dehydrogenation of alcohols can also be catalyzed by manganese(III) complexes and a number of experiments were now performed to optimize the reaction. First, several derivatives of Jacobsen’s catalyst were prepared to investigate the influence of the aryl substituent, the axial group on manganese, the scaffold and the Schiff base functionality (Fig. 2). Essentially the same result was observed when the unsubstituted analogue 2 (entry 3) and the tert-butyl group can therefore be omitted. The axial substituent on manganese, on the other hand, needs to be chloride since both the bromide 3 and the acetate 4 gave significantly lower conversion and yield (entries 4 and 5). The same was observed when the trans-1,2-diaminocyclohexane scaffold was changed to either the cis (i.e. 5), the ethylene (i.e. 6) or the benzene (i.e. 7) analogue (entries 6–8). Notably, the corresponding salan complex 8 also furnished some conversion into the imine (entry 9) which may indicate that the Schiff base functionality in the salen ligand plays an important role in the mechanism (vide infra). In the end, trans-N,N'-bis(salicylidene)-1,2-cyclohexanediame was selected as the salen ligand for the transformation and thus complex 2 as the catalyst.

For further optimization, the influence of additives and the solvent were investigated (Table 2). A number of common additives were included in the reaction, but they all led to lower imine yields due to a moderate conversion of benzyl alcohol (results not shown). We have previously used nitride salts as a basic additive for alcohol dehydrogenations, but lower yields and conversion were also observed when Li3N and Mg3N2 were added to the reaction (entries 1 and 2). However, with Ca3N2 a complete transformation of benzyl alcohol and a high yield of the imine was now obtained (entry 3). The optimum amount of Ca3N2 was 16.7% since both higher and lower quantities decreased the yield of the imine (entries 4–7). We observed the same optimum percentage in our earlier work and 16.7% is an intriguing number since one equiv. of Ca3N2 can theoretically react as a base with 6 equiv. of either the alcohol or water.

The influence of the solvent was then investigated and a slightly improved outcome was observed upon conducting the reaction in toluene (entry 8). Dioxane and heptane gave lower

Table 1  Dehydrogenation with manganese salen/salan complexes

<table>
<thead>
<tr>
<th>Entry</th>
<th>Catalyst</th>
<th>BnOH conversion (%)</th>
<th>Imine yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>5% Mn2(CO)10, 10% H2salen</td>
<td>—</td>
<td>38</td>
</tr>
<tr>
<td>2</td>
<td>1</td>
<td>81</td>
<td>79</td>
</tr>
<tr>
<td>3</td>
<td>2</td>
<td>86</td>
<td>81</td>
</tr>
<tr>
<td>4</td>
<td>3</td>
<td>70</td>
<td>65</td>
</tr>
<tr>
<td>5</td>
<td>4</td>
<td>66</td>
<td>64</td>
</tr>
<tr>
<td>6</td>
<td>5</td>
<td>83</td>
<td>74</td>
</tr>
<tr>
<td>7</td>
<td>6</td>
<td>60</td>
<td>58</td>
</tr>
<tr>
<td>8</td>
<td>7</td>
<td>27</td>
<td>25</td>
</tr>
<tr>
<td>9</td>
<td>8</td>
<td>47</td>
<td>12</td>
</tr>
</tbody>
</table>

Conditions: BnOH (1 mmol), CyNH2 (1 mmol), catalyst (0.1 mmol), tetradeacne (0.5 mmol, internal standard), 4 Å MS (150 mg), mesitylene (4 mL), reflux, 48 h. Determined by GC using the internal standard. GC yield based on the internal standard. 6% of N-cyclohexylbenzamidine was also detected. Traces of benzyl benzoate was also detected.
solvent (4 mL), readditive, tetradecane (0.5 mmol, internal standard), 4–10% Li3N Mesitylene 66 12d
1 10 20% Li3N Mesitylene 54 48f
2 10 20% Mg3N2 Mesitylene 100 92
3 10 20% Ca3N2 Mesitylene 100 93
4 10 16.7% Ca3N2 Mesitylene 100 93
5 10 33% Ca3N2 Mesitylene 84 84
6 10 10% Ca3N2 Mesitylene 100 90
7 10 5% Ca3N2 Mesitylene 100 88
8 10 16.7% Ca3N2 Toluene 100 98
9 10 16.7% Ca3N2 Dioxane 65 62
10 10 16.7% Ca3N2 Heptane 100 96
11 5 16.7% Ca3N2 Toluene 76 73
12 2.5 16.7% Ca3N2 Toluene 69 66
13 1.25 16.7% Ca3N2 Toluene 80 77
14 5 20% MgSO4 Toluene 80 77
15 5 20% Na2SO4 Toluene 87 79
16e 5 16.7% Ca3N2 Toluene 97 81
17f 5 20% Ca(OH)2 Toluene 99 95

d Conditions: BnOH (1 mmol), CyNH2 (1 mmol), 2 (X/100 mmol), additive, tetradeane (0.5 mmol, internal standard), 4 Å MS (150 mg), solvent (4 mL), reflux, 48 h. 

The inimination could be performed with anilines where 74%, 63% and 62% yield were obtained with aniline, p-anisidine and p-trifluoromethylaniline, respectively (entries 8–10). In entries 9 and 10 as well as in entries 5 and 6 about 10–30% of unreacted benzyl alcohol was also detected.

The reaction could be extended to the synthesis of pyroles by reacting cis-but-2-ene-1,4-diol with primary amines. The transformation could be carried out with both aniline and cyclohexylamine to afford the desired product in moderate yield (Scheme 1).

The gas evolution was measured in a separate experiment under the optimized conditions in Table 2, entry 16, but without adding Ca3N2 (resulting in a slightly lower 78% imine yield). One equivalent was collected and the gas was identified as dihydrogen, which confirms the acceptorless dehydrogenative pathway. Essentially no imine was formed when the reaction was conducted in the absence of complex 2. The possibility for trace metal impurities is a serious concern in the development of reactions with new metal catalysts and studies have shown that some coupling reactions with manganese catalysts are most likely mediated by traces of other elements. For that reason, complex 2 was analyzed by inductively coupled plasma mass spectrometry (ICP-MS) for traces of other metals known to perform alcohol dehydrogenations. However, none of these elements could be detected beyond their detection limit and it is therefore highly unlikely that another metal is responsible for the observed results.
As mentioned above, complex 2 dissolves completely in toluene upon heating the mixture to reflux. However, after the imination has gone to completion and the reaction is cooled to room temperature, complex 2 precipitates out again and 95% can be recovered after addition of a small amount of hexane. Furthermore, the retrieved complex can be subjected to a new catalytic reaction which gave 82% yield of the imine with a 4.5% catalyst loading.

When the optimized reaction in Table 2, entry 16 was performed with PhCD\(_2\)OH instead of PhCH\(_2\)OH, the product was exclusively PhCD\(_2\)NC\(_6\)H\(_5\) with no hydrogen incorporation into the benzylic position. This may imply that the dehydrogenation takes place by a monohydride pathway and no metal-dihydride species is formed.

To investigate whether the hydride abstraction takes place in the rate-limiting step, the primary kinetic isotope effect (KIE) was measured. The initial rate was determined with both PhCD\(_2\)OH and PhCH\(_2\)OH in the reaction with cyclohexylamine which gave a KIE of 2.00. This somewhat modest value shows that breakage of the C–H bond is one of several slow steps in the transformation.

To gain more experimental information about the reaction pathway, the studies were also supplemented by a Hammett study where the change in charge at the benzylic position between the starting material and the transition state can be determined. We have previously used Hammett studies with para-substituted benzyl alcohols to analyze the rate-limiting step in dehydrogenations catalyzed by ruthenium and iridium complexes.\(^\text{20}\) Thus, five para-substituted benzyl alcohols (X = OCH\(_3\), CH\(_3\), F, Cl and NO\(_2\)) were allowed to compete with the parent benzyl alcohol in the imination with cyclohexylamine. The reactions were monitored by GC, which allowed for determining the consumption of each alcohol. Assuming a first order reaction in the alcohol, their relative reactivities (k\(_X\)/k\(_H\)) can be determined as the slope of the line when ln(c\(_0\)/c) for one para-substituted benzyl alcohol is plotted against the same values for benzyl alcohol. These plots gave straight lines for all five para-substituted benzyl alcohols and made it possible to use the

### Table 3 Imination of alcohols with cyclohexylamine\(^a\)

<table>
<thead>
<tr>
<th>Entry</th>
<th>Alcohol</th>
<th>Imine</th>
<th>Yield(^b) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>PhOH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>89</td>
</tr>
<tr>
<td>2</td>
<td>CH(_3)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>70</td>
</tr>
<tr>
<td>3</td>
<td>CH(_3)CH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>83</td>
</tr>
<tr>
<td>4</td>
<td>CH(_2)OCH(_3)</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>90</td>
</tr>
<tr>
<td>5</td>
<td>PhCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>60</td>
</tr>
<tr>
<td>6</td>
<td>CH(_3)SCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>60</td>
</tr>
<tr>
<td>7</td>
<td>CH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>70</td>
</tr>
<tr>
<td>8</td>
<td>FCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>73</td>
</tr>
<tr>
<td>9</td>
<td>ClCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>85</td>
</tr>
<tr>
<td>10</td>
<td>BrCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>61</td>
</tr>
<tr>
<td>11</td>
<td>4-PhCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>71</td>
</tr>
</tbody>
</table>

\(^a\) Conditions: alcohol (1 mmol), CyNH\(_2\) (1 mmol), Ca\(_3\)N\(_2\) (0.167 mmol), toluene (4 mL), reflux, 48 h. \(^b\) Isolated yield. \(^c\) Reaction time 72 h.

### Table 3 (Contd.)

<table>
<thead>
<tr>
<th>Entry</th>
<th>Alcohol</th>
<th>Imine</th>
<th>Yield(^b) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>12</td>
<td>PhCH(_2)OH</td>
<td>PhCD(_2)NC(_6)H(_5)</td>
<td>60</td>
</tr>
</tbody>
</table>

As mentioned above, complex 2 dissolves completely in toluene upon heating the mixture to reflux. However, after the imination has gone to completion and the reaction is cooled to room temperature, complex 2 precipitates out again and 95% can be recovered after addition of a small amount of hexane. Furthermore, the retrieved complex can be subjected to a new catalytic reaction which gave 82% yield of the imine with a 4.5% catalyst loading.

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Hammett equation \( \lg \left( \frac{k_2}{k_1} \right) = \sigma \rho \) to construct a Hammett plot (Fig. 3). A good correlation was obtained with the standard \( \sigma \) values\(^2\) giving a straight line with a \( \rho \) value of \(-1.24\). This negative slope shows that substrates with electron-donating groups react faster and that a partial positive charge is built up at the benzylic carbon in the rate-limiting step consistent with a hydride transfer from the alcohol. Attempts to use different sets of \( \sigma \) values\(^3\) gave a poor correlation and radical intermediates are therefore not involved in the catalytic cycle. This was also confirmed by conducting the imination in the presence of one equiv. of the radical trapping agents cyclohexa-1,4-diene and 2,4-diphenyl-4-methylpent-1-ene which in both cases had no influence on the imine yield.

To further understand the mechanism, we used density functional theory (DFT) calculations to estimate the Gibbs free energies of possible intermediates and transition states. The starting point was the Mn(salen)OBn complex \( \text{9} \), where Cl has been replaced by a benzylic alkoxide (Fig. 4). This transformation involves the elimination of HCl, which is possible under the basic conditions. In fact, an experiment in the absence of a base (complex \( \text{2} \) and benzyl alcohol) gave no conversion at all while the absence of the amine (complex \( \text{2} \), benzyl alcohol and \( \text{Ca}_3\text{N}_2 \)) resulted in 20% of benzyl benzoate.

The initial idea was that complex \( \text{9} \) undergoes \( \beta \)-hydride elimination. However, the activation energy for this process was found to be prohibitively high (at 37.9 kcal mol\(^{-1} \) relative to \( \text{9} \)) which is partly due to the lack of an available coordination site. Instead, we found an alternative reaction where the hydride is transferred from the benzylic carbon to the imine carbon of the salen ligand. The activation energy was merely 17.6 kcal mol\(^{-1} \) and the product complex \( \text{11} \) is at 6.6 kcal mol\(^{-1} \). A similar pathway has been identified in the activation of (PNNP)Fe(II) eneamido complexes with isopropanol.\(^{23} \) The product benzaldehyde is then replaced by benzyl alcohol, from which a proton is transferred to the amide nitrogen of the reduced salen ligand. The resulting complex where one imine of the salen is

---

**Table 4**  Imination of amines with benzyl alcohol\(^a\)

<table>
<thead>
<tr>
<th>Entry</th>
<th>Amine</th>
<th>Imine</th>
<th>Yield(^b) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td></td>
<td></td>
<td>90</td>
</tr>
<tr>
<td>2</td>
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<td></td>
<td>69</td>
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<tr>
<td>3</td>
<td></td>
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<td>88</td>
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<td>4</td>
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</tr>
<tr>
<td>10</td>
<td></td>
<td></td>
<td>62</td>
</tr>
</tbody>
</table>

\(^a\) Conditions: \( \text{BnOH} (1 \text{ mmol}), \text{amine} (1 \text{ mmol}), 2 (0.05 \text{ mmol}), \text{Ca}_3\text{N}_2 (0.167 \text{ mmol}), \text{toluene} (4 \text{ mL}), \text{reflux, 48 h} \). \(^b\) Isolated yield.

---

**Scheme 1**  Pyrrole synthesis from but-2-ene-1,4-diol and amines.

---

Fig. 3  Hammett plot for the imination with para-substituted benzyl alcohols.
hydrogenated is at $-5.2$ kcal mol$^{-1}$ relative to 9. The lowest activation energy found from the hydrogenated intermediate was for another hydride transfer to the second imine of the salen, with a transition state at $14.5$ kcal mol$^{-1}$ (maximum $\Delta G_{\text{act}} = 19.7$ kcal mol$^{-1}$, see ESI† for more details).

After complete dissociation of benzaldehyde a key species 12 is formed, where one imine is hydrogenated to the amine and the other is reduced to an amide ligand. This species resembles intermediates from alcohol dehydrogenations with (PNP)Ru(II), (PNP)Mn(I), (PNNP)Fe(II) and (PNP)Ir(III) catalysts which have an amide ligand that can act as a Brønsted base and a metal that can serve as a hydride acceptor. They have all been proposed to react via an outer-sphere hydrogen transfer mechanism. The main difference is the metal and the oxidation state, which in the current case is manganese(III). The same outer-sphere hydrogen transfer was identified here and the activation energy is $26.7$ kcal mol$^{-1}$ relative to 12 and $27.2$ kcal mol$^{-1}$ relative to 15, which corresponds to a TOF of $83$ h$^{-1}$ at the reaction conditions (Fig. 5). The calculated KIE of this reaction is 2.9, which is in reasonable agreement with the experimental value of 2.0. Finally, we calculated the relative rates of the para-substituted benzyl alcohols used in the experimental study. The relative rates were calculated from the prereactive complex 15 and a good agreement was found with the experimental results giving a very similar $\rho$ value (Fig. 6). After the formation of the manganese(III) hydride intermediate 17, benzaldehyde is assumed to react irreversibly with the amine. From complex 17 the formation of hydrogen gas requires an activation energy of $22.6$ kcal mol$^{-1}$, in a step that regenerates the active catalyst.

The proposed mechanism indicates that the Schiff base functionality of the salen ligand is crucial for the reactivity. This was also confirmed by isolating the salen complex after the imination reaction with PhCD$_2$OH and cyclohexylamine. The isolated complex showed the incorporation of one deuterium atom on each of the Schiff base carbons as would be expected from the proposed mechanism. The involvement of the Schiff base was also observed in the optimization where both the salen complex 2 and the salan complex 8 catalyzed the dehydrogenation although the latter in a low yield (Table 1). When the imination with $5\%$ of the salan complex 8 was repeated with KOBu as the base, a $56\%$ yield was obtained of $N$-benzylidene cyclohexylamine. This result can be explained by elimination of HCl from 8 to afford the catalytically active species 12. Similar eliminations of hydrogen halides under basic conditions have been described with (PNP)Mn(I), (PNP)Ru(II) and (PNNP)Fe(II) complexes to form the corresponding amido compounds. Notably, a post analysis by LCMS of the imination with salan complex 8 showed the formation of salen complex 2 as the main manganese species together with minor amounts of some unidentified complexes. None of the starting complex 8 could be detected after the imination. These observations show that a salan complex can be converted into the corresponding salen complex under the reaction conditions and explains why salen complex 2 is almost fully recovered after the reaction. So far, however, we have not been able to identify...
a pathway by which hydrogen is eliminated from a salan complex to afford the salan compound.

Conclusions

In summary, we have described a new catalyst for the acceptorless dehydrogenation of alcohols. The manganese(III) salen complex 2 mediates the formation of imines from alcohols and amines with the liberation of hydrogen gas. The reaction can be performed with different alcohols and amines and can be extended to the synthesis of pyroles. Complex 2 can be recovered from the reaction and used again without significantly affecting the catalytic activity. The mechanism is believed to involve a bifunctional pathway where both the metal and the ligand participates in the dehydrogenation reaction. We envision the discoveries will spur much interest in the development of new transformations with hydrogen gas and manganese(III) catalysts.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The project was supported by the Technical University of Denmark through PhD fellowships to SVS and CS. We thank Mathias Bersting for his assistance in performing the Hammett study and determining the KIE.

Notes and references


Complex 1 is commercially available, but in our hands its catalytic performance in the dehydrogenation depended on the supplier. Therefore, all manganese(II) complexes used in this work were synthesized by standard literature protocols and no commercial samples were used. The presynthesized complexes gave reproducible results and could be stored at room temperature under an inert atmosphere for a longer period without affecting the product yield.

KOH, NaOH, LiOH, KO-t-Bu, K$_3$PO$_4$, Cs$_2$CO$_3$, AgBF$_4$, Et$_3$N, MgBr$_2$, MgO, LiCl, LiBr, and LiF.


Similar yields are reported in ref. 16.


This elimination does not appear to be a very facile reaction, which may explain the lower yield of the imine with complex 8 as the catalyst. Treatment of complex 8 with one equiv. of KOt-Bu in refluxing toluene in the absence of the substrates showed no conversion into other manganese compounds after 1 h according to LCMS. Upon prolonged treatment, other manganese species were formed with complex 2 as the main compound after 48 h.