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Hydration of nitriles using a metal–ligand cooperative ruthenium pincer catalyst†

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Nitrile hydration provides access to amides that are important structural elements in organic chemistry. Here we report catalytic nitrile hydration using ruthenium catalysts based on a pincer scaffold with a dearomatized pyridine backbone. These complexes catalyze the nucleophilic addition of H₂O to a wide variety of aliphatic and (hetero)aromatic nitriles in ^tBuOH as solvent. Reactions occur under mild conditions (room temperature) in the absence of additives. A mechanism for nitrile hydration is proposed that is initiated by metal–ligand cooperative binding of the nitrile.

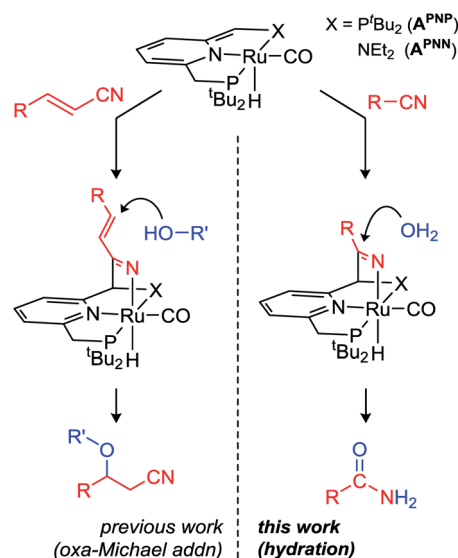
Introduction

Amides are an important class of compounds that occur in a large variety of biologically active compounds, polymers and synthetic intermediates,¹ and a variety of synthetic methods have been developed for their formation.² Nitrile hydration provides an atom-efficient synthesis of amides and is carried out on an industrial scale, for example in the production of acrylamide³ and nicotinamide.⁴ The direct nucleophilic addition of water to the C≡N bond is kinetically slow, and a variety of catalysts have been developed, but it is often difficult to prevent over-hydrolysis to the corresponding carboxylic acids.⁵ Nitrile hydratase enzymes yield amides with excellent selectivity,⁶ but the limited substrate scope prevents their widespread use. There has been significant recent interest in the development of nitrile hydration catalysts based on transition metals, but most systems reported to date still require relatively high temperatures to reach appreciable catalytic turnover.⁷ Recent progress with Rh(I),⁸ Ru(II),⁹ Pd(II),¹⁰ and Pt(II)¹¹ catalysts has allowed catalytic nitrile hydration under mild conditions, but additives are often required for high activity (*e.g.*, AgOTf in Grubbs' Pt(II)/phosphinous acid catalysts,¹¹ or Sc(OTf)₃ in Yin's Pd(II) catalysts¹⁰).

Metal complexes with pincer ligands have found widespread use in a large variety of catalytic reactions,¹² but the incorporation of a reactive fragment in the ligand backbone to enable 'bifunctional' or 'cooperative' substrate binding/activation has only recently started to emerge.¹³ Examples include the (reversible) binding of unsaturated fragments such as CO₂,¹⁴ SO₂,¹⁵ carbonyl compounds,¹⁶ or nitriles.¹⁷ Our group reported

that Milstein's dearomatized Ru PNN pincer complex **A^{PNN}** (Scheme 1, left) catalyzes the conjugate addition of alcohols to α,β-unsaturated nitriles *via* a metal–ligand cooperative (MLC) mechanism by activation of the nitrile C≡N bond.¹⁸ Very recently, similar reactivity was observed with a related Mn catalyst.¹⁹ This mode of activation reduces the bond order from 3 in the nitrile (C≡N) to 2 in the MLC intermediate (–C=N–Ru), which significantly alters the reactivity profile of the substrate.

Having established that conjugate (1,4-) addition of weak alcohol nucleophiles is enabled by metal–ligand cooperation, we hypothesized that also 1,2-additions to MLC-activated nitriles may be feasible. Herein we describe our results on catalytic nitrile hydration using Ru complexes with dearomatized pyridine-based pincer ligands, and demonstrate that a large variety of aliphatic and (hetero)aromatic nitriles is



Scheme 1 MLC activation of nitriles towards addition of O-nucleophiles.

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with this hypothesis is the observation that 4-hydroxybenzonitrile is not converted even at 80 °C, indicating that the acidic Ar-OH moiety deactivates the catalyst in an irreversible manner.^{18a,23}

Unhindered esters such as 4-acetoxycyanobenzonitrile and ethyl 4-cyanobenzoate also do not form the amide products, likely because of competing ester hydrolysis to the carboxylic acids. For more hindered esters, however, the reaction works well as demonstrated by full conversion of *tert*-butyl 4-cyanobenzoate (**1i**).

Heteroaromatic nitriles are also converted, with even shorter reaction times under the standard conditions (<1 h with 3 mol% catalyst **A^{PNP}**). Consequently, this class of substrates reaches full conversion within a day at room temperature using a catalyst loading of 0.5 mol% (TON = 200). The *ortho*-, *meta*- and *para*-isomers of cyanopyridine (**1p-r**), as well as the pyrazine derivative (**1s**) are converted to the amide products in quantitative yields. Moreover, substrates with oxygen- (furan, **1t,u**) or sulfur-containing rings (thiophenes, **1v,w**), structural motifs that are often considered catalyst poisons, underwent this ruthenium-catalyzed process with remarkable ease.

Substrates with sp³-C substituents adjacent to the C≡N group react more slowly but nevertheless are converted with complete selectivity to the amide. For example, benzyl cyanide (**1x**) and its α -methylated analogue (**1y**) show full conversion to the amide within 1 day at room temperature, and the same applies to 3-phenylpropionitrile (**1z**) and the sp²-substituted cinnamonitrile (**1aa**). While purely aliphatic nitriles also react at room temperature, these substrates require longer reaction times to reach synthetically useful conversions (65–83% after 2 days). However, gentle heating of the reaction mixtures to 50 °C for 1 day affords amide products in >99% isolated yield for primary, secondary as well as tertiary alkyl nitriles (**1ab-ae**). Substrates with two nitrile moieties were selectively converted to the corresponding diamides (**1af/1ag**).

The substrates shown in Table 1 are converted by catalysts **A** already at room temperature (and many reach full conversion within 24 h), but, with the exception of heteroaromatic nitriles, turnover numbers (TONs) for the majority of entries are only modest due to the presence of 3 mol% catalyst (maximum TON = 33). To examine whether higher turnover numbers can be obtained, we tested the reaction at elevated temperature (70 °C) for aromatic nitrile **1b** and aliphatic dinitrile **1af**. For **1b**, hydration is initially fast (63% conversion in 1 h), but then gradually slows down to reach 98% conversion in 20 h under those conditions (TON = 196). Dinitrile **1af** also gave >98% conversion of the starting material in 24 h at 70 °C to afford a mixture of mono- and diamide products in a 57 : 43 ratio, which corresponds to a total nitrile TON of 307. Moreover, with 3-cyanopyridine (**1q**) the catalyst reaches 1000 turnovers within a day at 70 °C, demonstrating the robustness of the catalyst at elevated temperature.

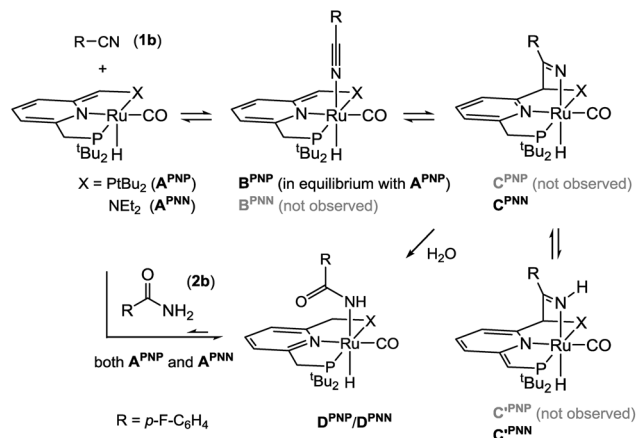
Mechanistic considerations

To obtain insight in the species present during turnover, we monitored a catalysis reaction mixture (3 mol% **A^{PNP}**, 5 eq. H₂O

in ⁴BuOH) with the fluoro-substituted benzonitrile **1b** by ¹⁹F NMR spectroscopy. This showed the presence of three distinct ¹⁹F-containing species. Two of these correspond to the nitrile starting material and the amide product by comparison to authentic samples. A third species was present throughout the course of the reaction in minor amount (¹⁹F NMR: δ –118 ppm, *ca.* 2.5% based on integration). This resonance was also observed in the ¹⁹F NMR of a catalysis mixture with **A^{PNN}**, suggesting that **A^{PNP}** and **A^{PNN}** lead to similar speciation under the reaction conditions.

To characterize this species and develop an understanding of the individual steps that might be involved in the catalysis, we carried out stoichiometric NMR scale experiments in THF-*d*₈ between the dearomatized pincer complexes **A** and the components present in the catalysis reaction mixture (see ESI for details†). Analysis of a 1 : 1 mixture of complex **A^{PNP}** and nitrile **1b** by ¹H NMR spectroscopy showed a substantial broadening of the Ru–H signal which is additionally shifted downfield by *ca.* 7 ppm, indicating a rapidly exchanging equilibrium between the starting materials and the Ru-nitrile adduct **B^{PNP}** (Scheme 2).^{18b}

The sterically less hindered PNN analogue **A^{PNN}** reacts with nitrile **1b** to an equilibrium mixture of **C^{PNN}** and **C'^{PNN}** according to NMR spectroscopy. In this mixture, a characteristic ¹H NMR resonance is observed at δ 11.65 (singlet) for the NH fragment in **C^{PNN}**. Additionally, signals at δ –10.47/–13.75 ppm (doublets, $J_{\text{PH}} = 25.7/32.8$ Hz) appear for the Ru–H groups in **C^{PNN}**/**C'^{PNN}**. The NMR features are very similar to those observed previously by us for the reaction of **A^{PNN}** and benzonitrile,^{18b} and confirm that the latter products arise from MLC binding of the C≡N bond. The different outcomes of the reaction between **1b** and **A^{PNN}** or **A^{PNP}** indicates that the divergent steric demands of the PNP *vs.* PNN ligand affects the equilibria between species **A**, **B** and **C**. Although a detailed comparison between PNN- and PNP-based catalysts is beyond the scope of this research, preliminary DFT calculations indicate that MLC-binding of benzonitrile is exergonic at the N-arm (–2.7 kcal mol^{–1}) whereas it is endergonic at the P-arm (PNN:



Scheme 2 Stoichiometric reactions between **A** and the components present in the reaction mixture.



+9.4 kcal mol⁻¹; PNP: +7.2 kcal mol⁻¹). It should be noted that, even though the relative stability of the various intermediates is sensitive to steric effects, metal–ligand cooperative reaction pathways are involved in complexes with both these ligands.²⁴

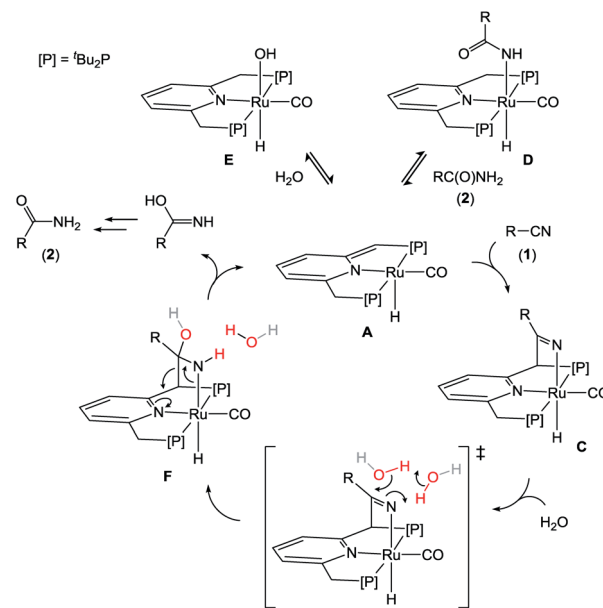
Regardless of the different equilibrium compositions, addition of H₂O (1 equiv.) to a THF-*d*₈ solution of both the PNN- or PNP-based mixture resulted in the slow appearance of a new species (**D**) with a ¹⁹F NMR shift (*ca.* δ −118 ppm) that agrees well with the one observed during catalysis in ^tBuOH. Compounds **D** were also obtained cleanly by treatment of the dearomatized Ru pincer complexes **A** with amide **2b**, and are formulated as Ru-carboxamides (**D**^{PNN}/**D**^{PNP}, Scheme 2) on the basis of their NMR spectra (see ESI†).

The observation that **D** is the dominant Ru-containing species during turnover suggests that it may be a dormant catalyst state, and that catalysis could be subject to product inhibition. Indeed, when a mixture of nitrile substrate and amide product (33 and 10 equiv., respectively, relative to **A**^{PNP}) is present at the start, the reaction rate is decreased but full conversion is still obtained (see ESI†). Also, when **D**^{PNP} is prepared independently and tested in catalysis, hydration of **1b** occurs with a rate that is qualitatively similar to that with **A**^{PNP}.

These observations indicate that the coordinatively saturated complex **D** is able to generate the active species *via* a rapid equilibrium with **A**. This was further confirmed by the observation that **D**^{PNN} reacts with nitrile **1b** to regenerate the equilibrium mixture of **C**^{PNN} and **C'**^{PNN}, a reaction that presumably involves **A**^{PNN} as an intermediate (Scheme 2). For the PNP analogue, the concentration of **D**^{PNP} decreases upon addition of H₂O, and NMR resonances attributable to the free amide **2b** appear. From these data it is clear that the catalyst speciation in this system is complex: the dearomatized pincer complexes **A** are involved in dynamic equilibria with H₂O, nitrile, as well as amide. Thus, even though under the reaction conditions most of the Ru is present in a coordinatively saturated, inactive form (**D**), access to catalytically active species is kinetically facile and allows turnover of the nitrile substrate.

Mechanistic proposals for catalytic nitrile hydration in the literature often involve Lewis acid activation of the nitrile, sometimes in conjunction with ligand-mediated ('bifunctional') deprotonation of H₂O,^{25,26} or nucleophilic attack of a reactive ligand–OH fragment (*e.g.*, in catalysts with phosphinous acid ligands).²⁷ For bimetallic complexes, a reactive metal-hydroxide group can be generated adjacent to a metal–nitrile adduct, which provides a low-energy pathway to hydration.²⁸

On the basis of the results discussed above, we propose that the mechanism of nitrile hydration by **A** is distinct and follows the steps illustrated in Scheme 3. Under the reaction conditions, complexes **A** react reversibly with amide or H₂O to form the off-cycle species **D** and **E**, respectively. Although these species are not catalytically relevant, a rapid equilibrium between these dormant states and **A** ensures an entry into the catalytic cycle. We propose that catalysis is initiated by MLC binding of the nitrile substrate (**1**) to form **C**. The MLC mode of C≡N bond activation results in a reduced CN bond order of **2**, and transfers the basicity 'stored' in the pincer framework in **A** onto the nitrile-derived Ru–N moiety in **C**. This allows



Scheme 3 Proposed catalytic cycle.

deprotonation of the pro-nucleophile H₂O and attack at the electrophilic C-atom of the activated nitrile to form **F**, either directly or facilitated by a hydrogen-bond network involving additional H₂O or ^tBuOH as proton-shuttles.^{29,27b} The coordinated hydroxyamido fragment in intermediate **F** may be liberated from the metal complex by a retro-cycloaddition to form the iminol, which tautomerizes in solution to the final amide product (**2**). Alternative pathways that directly convert **F** to the Ru-carboxamide **D** cannot be ruled out at present. The proposed reaction pathway accounts for several key experimental observations, including (i) the requirement for a reactive ligand site for catalytic activity, (ii) an optimum in catalyst activity as function of the amount of water added, (iii) and the occurrence of product inhibition.

Conclusions

In summary, this work describes an efficient homogeneous catalyst for the hydration of nitriles. Complexes based on dearomatized PNP or PNN ligands are shown to be active under very mild reaction conditions (ambient temperature, additive-free). The PNP Ru pincer catalyst is tolerant to a variety of functional groups, and allows the hydration of a broad range of aliphatic, aromatic and heteroaromatic nitriles. On the basis of stoichiometric experiments, a mechanism is proposed that involves metal–ligand cooperative activation of the nitrile C≡N bond. The results suggest that the generation of intermediates with a C=N moiety (*i.e.*, a reduced bond order in comparison to the nitrile starting material) *via* this mode of nitrile activation significantly increases its reactivity and leads to facile attack by (pro)nucleophiles as weak as H₂O. We anticipate that this strategy may be more broadly applicable and lead to novel reactivity of nitriles and other unsaturated organic compounds. A more detailed examination of the catalytic reaction



mechanism, as well as modification of the catalyst to increase productivity (e.g., by minimizing product inhibition) are ongoing in our laboratory.

Conflicts of interest

There are no conflicts to declare.

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