Small-molecule activation at Au(III): metallacycle construction from ethylene, water, and acetonitrile†

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Incorporation of the simple, readily available, building blocks ethylene, water and acetonitrile into Au(tpy)(OCOCF3)2 (tpy = 2-(p-tolyl)pyridine) in a one-step reaction leads to high yields of a new 6-membered ring gold(III) metallacycle complex. The metallacycle has been characterized spectroscopically and crystallographically, and the mechanism of its formation has been investigated with the aid of DFT calculations.

Introduction

The functionalization of alkenes has great practical and economic value in catalysis, and as the simplest of such building blocks, ethylene is of particular interest.1 Gold is known for its ability to π-coordinate and thereby activate alkenes and alkynes towards nucleophilic attack, which is considered to be a key step in gold catalysis.2–8 However, there are only scarce reports of Au(III)-mediated functionalization of the simplest olefin, ethylene. One important example was provided by Atwood and co-workers (Scheme 1, top),9 who demonstrated that ethylene and propylene could be functionalized stoichiometrically at the Au(III) complex Au(bipy)Cl2+ (bipy = 2,2′-bipyridine) in water to furnish Au(III) hydroxyalkyl products that were observed in solution by 1H NMR spectroscopy but not isolated. Reactions of ethylene and propylene with HAuCl3 and AuCl3(tppts) (tppts = 3,3′,3″-phosphanetriyltris(benzenesulfonic acid) trisodium salt) led to Au(III) hydroxyalkyl species, which upon heating underwent gold reduction to the metal with concomitant formation of organic oxygenated products (alcohols, aldehyde or ketone).9 More recently, Bochmann and co-workers (Scheme 1, bottom)10 showed that ethylene slowly undergoes a formal insertion into the Au–OAcF bond trans to the pyridine-N in a diarylpyridine CNC pincer complex to yield an Au(III) acetoxyalkyl complex (OAcF = OCOCF3).

Our group recently reported a combined experimental and computational study of the formal insertion of ethylene into an Au–O bond of Au(tpy)(OAcF)2 (1; tpy = 2-(p-tolyl)pyridine; Scheme 2, top).11 Selective insertion into the Au–O bond trans to N of the chelating tpy ligand furnished Au(tpy)–(CH2CH2OAcF)(OAcF) (2). Although the trans to C coordination site is kinetically more accessible, the trans to N insertion product is thermodynamically favoured and is formed by nucleophilic attack by free “OAcF” at the coordinated ethylene. This discovery led to further investigations, and herein we report that the coordinated and inserted ethylene molecule can undergo further functionalization due to the availability of the trans to C coordination site at Au.
of water. However, involvement of reagents arising from a (possibly gold) catalyzed hydrolysis of acetonitrile, or from impurities inadvertently present in this solvent, must also be considered.

By positive-ion high resolution mass spectrometry (HRMS), the parent ion at \( m/z = 451.1068 \) established the elemental composition \( C_{16}H_{18}AuN_2O \) for the cation (calcd \( m/z = 451.1084 \)), in accord with the presence of one N and one O atom in the chelate ring. This composition, which eliminates alternative 5, was corroborated by elemental analysis. Support for the formulation of the \( N,O \)-containing chelate 3 was obtained by \( ^1\text{H} \)–\( ^{15}\text{N} \) HMBC NMR spectroscopy. In the spectrum, \( ^1\text{H} \) signals arising from two different N atoms are clearly observed. One signal at \( \delta(^{15}\text{N}{^1\text{H}}) = -212 \) arises from the chelate NH and correlates with the broadened \( ^1\text{H} \) signal of the NH group at \( \delta(^1\text{H}) = 10.28 \) and with the signal from the metal-lacyle methyl group at \( \delta(^1\text{H}) = 2.51 \), but not with the other protons of the metallacycle. The other \( ^{15}\text{N} \) signal at \( \delta(^{15}\text{N}{^1\text{H}}) = -129 \) arises from the N atom in the tpy ligand and has a correlation with the signal arising from the CH proton \( \alpha \) to the pyridyl-N atom at \( \delta(^1\text{H}) = 9.33 \). The \( ^{19}\text{F} \) NMR spectrum shows only one signal, arising from the \( -\text{OAc}^\cdot \) anion. So as to further unambiguously distinguish between alternatives 3 and 4, a NOE experiment was performed. The \( ^1\text{H} \)–\( ^1\text{H} \) NOESY spectrum shows a clear NOE interaction between the NH proton at \( \delta \) 10.28 and the CH proton at \( \delta 9.33 \). These H atoms are therefore in close proximity, supporting the crystallographic analysis and verifying that 3 does indeed represent the structure of the metallacycle.

An ORTEP plot of the molecular structure of 3 is shown in Fig. 1, with selected bond distances and angles. The \( \text{Au}^{\text{III}} \) centre has the expected square planar geometry. The deviation of the C9–Au–N1 bond angle in the tpy chelate from 90 to 80.60(10)\(^\circ\) is typical, and the \( \text{Au} \)–ligand distances \( \text{Au}1–\text{N1} \), \( \text{Au}1–\text{C9} \), and \( \text{Au}1–\text{C4} \) (sp\(^3\)) are within the ranges that have previously been observed for the related complexes \( \text{Au}(\text{tpy})(\text{Me}_2) \), \( \text{Au}(\text{tpy})(\text{CH}_2\text{CH}_2\text{OAc})_2 \), and \( \text{Au}(\text{tpy})(\text{CH}_2\text{CH}_2\text{OCH}_2\text{CF}_3) \) \( (\text{OAc}^\cdot) \). The N5–C3 (1.284(4) \( \text{Å} \)) and C3–O2 (1.332(3) \( \text{Å} \)) bond distances are consistent with the corresponding average N–C (1.282 \( \text{Å} \)) and O–C (1.325 \( \text{Å} \)) in \( \text{Pt}(\text{n}) \) imino ether complexes.\(^{15–23} \) These distances are, however, distinctly different from the distances in a [(tpa)Rh(II)]\(^{2+}\) (tpa = \( N,N,N \)-tri (2-pyridylmethyl)amino) analogue to 3, in which the corresponding N–C and O–C bond distances are 1.245(19) and 1.374 (17) \( \text{Å} \), respectively.\(^{23} \) The directionality of the N5–O1A interaction between the NH of the 6-membered ring and the nearest O (O1A) of the trifluoroacetate anion suggests the presence of a hydrogen bond; in fact, the final refinement shows a donor–acceptor distance (N5–O1A) of 2.8708(2) \( \text{Å} \). This is in excellent agreement with the proposed structure 3. The crystal structure of 3 is monoclinic, but with \( \beta = 90^\circ \). In addition, it is twinned \( via \) a twofold rotation about the crystallographic a

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See ESI pages S5, S13, and S14† for details.

† No structural examples of \( \text{Au} \) imino ethers were found \( via \) SciFinder.
Mechanistic studies by experiment and DFT calculations

Initial mechanistic experiments probed whether a gold-mediated or gold-catalyzed hydrolysis of acetonitrile to form acetamide might be involved in the process (Scheme 4). First, no formation of 3 was seen when 2 was treated with only acetamide in CH₂Cl₂ (Scheme 4A). Second, the reaction of 2 with wet acetonitrile-d₃ in the presence of acetamide led to the generation of 3-d₃, and a minor amount of 3 as inferred by integration of the metallacycle–methyl resonance at δ 2.51 in the ¹H NMR spectrum (Scheme 4B). These experiments suggest that acetonitrile, and not acetamide, is the main reactant in the formation of the metallacycle.

Scheme 5 shows a possible mechanism for the formation of 3. The unobserved cationic ethylene complex 6 is generated from either 1 or 2, in agreement with our previous findings. The hydroxyethyl complex 7 is formed from 2 after nucleophilic attack by water, by analogy with Atwood’s study, and parallel to reaction of trifluoroethanol with 2, which proceeds via 6 yielding 10. Acetonitrile substitution for the OAcF anion of 7 at the kinetically most accessible position trans to tpy-C furnishes a crucial intermediate 8, which undergoes cyclization by an intramolecular nucleophilic attack of the hydroxyethyl-O atom on the nitrile-C atom of acetonitrile, analogous to the alcoholysis of Pt(n)-coordinated nitriles to provide Pt(n) imino ethers. An OAcF anion assisted proton transfer from O to N finally provides 3. Close analogues of the putative intermediates 7 and 8 exist and support their mechanistic viability. The hydroxyethyl complex 7 is an analogue of Au(tpy)Me(OTf) and Au(tpy)Me(OAc); both have been structurally characterized as the pertinent stereoisomers with Me trans to tpy-C.

The relative stability of all intermediates shown in Scheme 5 has been assessed by means of DFT calculations, which show that they are all accessible under the experimental conditions. The TS from 8 to 11 has also been computed along with the pertinent stereoisomers with Me trans to tpy-C.

§ For an estimation of the transition state energies involved in the transformation of 1 to 7, see ref. 11.
Fig. 3) and it shows that cyclization is triggered by the deprotonation of the hydroxyl group by an external \(\text{OAc}^-\). This transition state is 13.0 kcal mol\(^{-1}\) higher than \(\text{1}\) and involves an effective energy barrier of 21.8 kcal mol\(^{-1}\) relative to \(\text{2}\). The long experimental reaction time required for this transformation may be due to the existence of many equilibria, which are dependent on the different reactant concentrations.

Concluding remarks

An eventual reductive elimination from \(\text{3}\) would generate 2-methyl-2-oxazoline, with an important heterocyclic ring structure. Previously, 2-oxazolines have been prepared by Sawamura and coworkers\(^{26-28}\) by Au(III) catalyzed asymmetric aldol reactions of isocyanides and aldehydes. Hashmi and coworkers more recently\(^{29,30}\) prepared oxazolines by Au(l)-catalyzed cycloisomerizations of propargyl amides. Interestingly, it has been recently reported that cyclometallated Au(III) aryl–pyridine complexes may act as efficient catalysts for the three-component synthesis of propargylic amines from aldehydes, secondary amines, and alkynes, and of substituted oxazoles from \(N\)-benzyl imines, alkynes, and acyl chlorides.\(^{31,32}\) The \([\text{tpa}]\text{Rh}^{\text{III}}\) analogue to \(\text{3}\) was synthesized in a two-step process by \(\text{H}_2\text{O}_2\) oxidation of a coordinated ethylene, followed by reaction with acetonitrile and \(\text{NH}_4\text{PF}_6\).\(^{33}\) In this context, our one-step assembly of an oxazoline from an alkene, a nitrile, and water represents a new strategy with obvious possibilities in organic synthesis.\(^{33}\)

In conclusion, the convenient and high-yield one-pot, four-component assembly of a new cationic metallacyclic Au(III) complex has been described. In the process, the three small, readily available building blocks ethylene, water, and acetonitrile have been incorporated into one product. The transformation demonstrates that both of the potentially labile coordination sites in \(\text{Au(tpy)(OAc)}^3\) can be utilized – a fact that may be of importance for catalytic applications. The metallacycle is constructed in such a fashion that the high \(\text{trans}\) (alkyl) end of the assembled chelate ligand occupies the position that is \(\text{trans}\) to the low \(\text{trans}\)-effect (N donor) of the tpy supporting ligand, and \text{vice versa.}

Experimental and computational section

General experimental methods

Gold(III) complexes \(\text{1}\) and \(\text{2}\) were prepared by previously reported procedures.\(^{11,14}\) Distilled water was used. \(\text{CH}_3\text{CN}\) and \(\text{CH}_2\text{Cl}_2\) were purified using a MB SPS-800 solvent purifying system from MBraun. The gold(III) complexes studied here are not sensitive to air, so inert atmosphere was not utilized, except for the synthesis of \(\text{9}\) which is described in the ESI.\(^{\dagger}\) NMR spectra were recorded on Bruker Avance D PX200, AVII400, DRX500, AVII600 and AV600 instruments at ambient temperature. \(^1\text{H}\) and \(^{13}\text{C}\) spectra have been referenced relative to the residual solvent signals. \(^{19}\text{F}\) has been referenced to \(\text{CFCl}_3\) by using \(\text{C}_6\text{F}_6\) (\(-164.9\) ppm with respect to \(\text{CFCl}_3\) at 0 ppm) as an internal standard. The \(^{15}\text{N}\) chemical shifts have been calibrated using \(\text{MeNO}_2\) as an external standard at 0 ppm. The peaks in the \(^1\text{H}\) NMR and \(^{13}\text{C}\) NMR spectra were assigned by the aid of 2D NMR techniques such as HSQC, HMBC, COSY, NOESY, and \(^1\text{H}\)–\(^{15}\text{N}\) HMBC according to the numbering scheme shown below. Mass spectra (ESI) were obtained on a Micromass QTOF II spectrometer and a Bruker Daltonics maXisII spectrometer. Elemental analysis was performed by Mikroanalytisches Laboratorium Kolbe, Mülheim an der Ruhr, Germany.
Preparation of [Au(tpy)(C–N)]^+ [OOCF_3] (3) from 1. Au(tpy) (OOCF_3)_2 (50.1 mg, 0.0847 mmol, 1.0 equiv.) was dissolved in acetonitrile (3 mL). Water (10 µL, 0.55 mmol, 6.5 equiv.) was added. Ethylene was bubbled through the solution at ambient pressure for two minutes, and the flask was sealed with a glass stopper. The reaction mixture was stirred at ambient temperature in the absence of light for 7 days. The reaction mixture was filtered and the volatiles were removed under reduced pressure furnishing 3 as a white solid (38.1 mg, 0.0675 mmol, 80%).

Preparation of [Au(tpy)(C–N)]^+ [OOCF_3] (3) from 2. Au(tpy)- (CH_2CH_2O)OCOCF_3 (OCOCF_3) (50.1 mg, 0.0809 mmol, 1.0 equiv.) was dissolved in acetonitrile (3 mL). Water (10 µL, 0.55 mmol, 6.8 equiv.) was added. Ethylene was bubbled through the solution at ambient pressure for two minutes, and the flask was sealed with a glass stopper. The reaction mixture was stirred at ambient temperature in the absence of light for 5 days. The reaction mixture was filtered and the volatiles were removed under reduced pressure furnishing 3 as a white solid (35.8 mg, 0.0634 mmol, 78%).

1H NMR (600 MHz, CD_2Cl_2): δ 10.28 (bs, 1H, NH), 9.33 (d, 1H, J = 5.4 Hz, H^6), 8.08 (ddd, 1H, J = 7.5, 1.4 Hz, H^3), 7.96 (d, 1H, J = 8.0 Hz, H^6), 7.71 (d, 1H, J = 7.7 Hz, H^4), 7.58 (ddd, 1H, J = 7.5, 5.6, 1.1 Hz, H^3), 7.23–7.25 (m, 2H, H^4 and H^5), 4.34 (m, 2H, OCH_2), 2.73 (m, 2H, AuCH_2), 2.51 (s, 3H, OCH_3), 2.44 (s, 3H, ArCH_3).

13C NMR (150 MHz, CD_2Cl_2): δ 175.2, 161.7, 149.8, 142.4, 142.3, 142.0, 141.9, 131.2, 129.4, 126.0, 124.9, 120.2, 66.5, 28.6, 22.0, 11.4 Hz. 15N{1H} NMR (600 MHz, CD_2Cl_2): δ −29 (NH). 1HF NMR (188 MHz, CD_2Cl_2): δ −78.1 (bs, CF). MS (ESI^+, MeCN): m/z (rel.%) 451 [(M – OOCF_3)_2^+], 100). MS (ESI^+, MeCN): m/z (rel.%) 113 [(OOCF_3)_2^+], 100) was observed among other unidentified peaks. HRMS (ESI^+, MeCN): Found: 451.1068; calcd for C_{16}H_{12}AuN_2O_6: 451.1084 (+0.0016).

Elemental analysis: Anal. Calcd for C_{16}H_{12}AuN_2O_6: C, 38.31; H, 3.22; N, 4.96. Found: C, 37.81; H, 3.19; N, 4.59.

Computational details

Calculations were carried out at the DFT level as implemented in the Gaussian09 software package. Including Grimme’s model for dispersion forces was used to optimize all geometries. This methodology was selected because previous studies have proven its solid performance in the modeling of Au(n) alkene complexes. Geometries were fully optimized without any constraint. Vibrational frequencies were computed analytically to verify that the stationary points found were energy minima or transition states. All optimizations needed for the mechanism proposal were carried out in solvent (acetonitrile) using the SMD solvation model.

5 were optimized in gas phase with the aim of comparing their geometries with an X-ray crystal structure. Gibbs energies were obtained for T = 298.15 K and p = 1 atm. In the bimolecular steps, these energies were corrected for the 1 M standard state (T = 298.15 K and p = 24.465 atm). Further computational details are given in the ESI.

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Notes and references


The synthesis, characterization, and X-ray structure of Au(tpy)(Me)(NCMe)⁺BF₄⁻ is given in the ESL.†


