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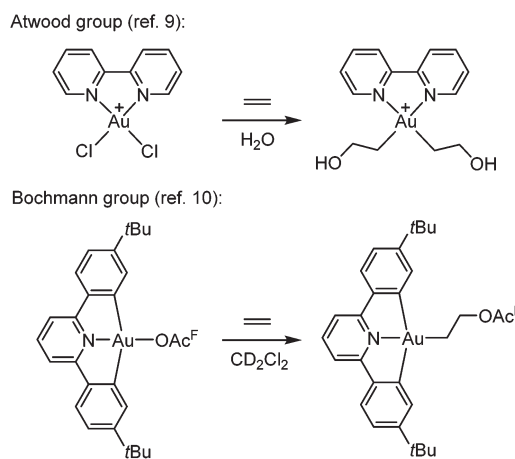
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Small-molecule activation at Au(III): metallacycle construction from ethylene, water, and acetonitrile†

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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

Scheme 1 Reactions of ethylene at Au(III) complexes (OAc^F = OCOCF₃).

Au–OAc^F bond *trans* to the pyridine-N in a diarylpyridine CNC pincer complex to yield an Au(III) acetoxyalkyl complex (OAc^F = OCOCF₃).

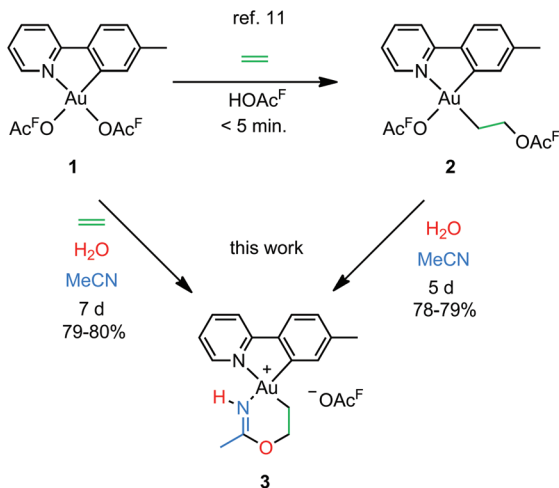
Our group recently reported a combined experimental and computational study of the formal insertion of ethylene into an Au–O bond of Au(tpy)(OAc^F)₂ (**1**; tpy = 2-(*p*-tolyl)pyridine; Scheme 2, top).¹¹ Selective insertion into the Au–O bond *trans* to N of the chelating tpy ligand furnished Au(tpy)-(CH₂CH₂OAc^F)(OAc^F) (**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 [–]OAc^F 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.

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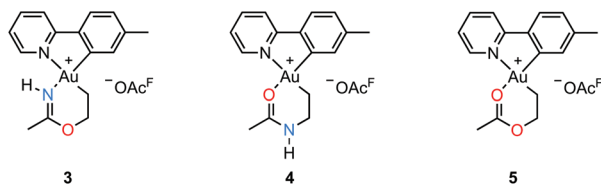
Scheme 2 Formal insertion of ethylene into an Au–O bond of Au(tpy)(OAc^F)₂ (**1**)¹¹ and preparation of the metallacyclic Au(III) complex **3** from **1** and **2** (this work); (OAc^F = OCOCF₃).

Results and discussion

Synthesis and characterization

An attempt to crystallize a product mixture containing **2** from acetonitrile led instead to the isolation of small quantities of a new crystalline complex. Examination of these crystals by single-crystal X-ray diffraction revealed that the new complex was a Au(III) metallacycle **3** (Scheme 2, lower part). Synthesis of larger quantities of **3** in about 80% yield was possible simply by stirring **2** in wet acetonitrile for 5 days at ambient temperature. Alternatively, **3** can be obtained in comparable yields by dissolving **1** in acetonitrile and adding water and ethylene. Thus, an efficient synthetic protocol has been developed wherein the three small, readily available building blocks ethylene, water, and acetonitrile are incorporated into the Au(III) metallacycle complex **3** in just one step.

The molecular structure, based on the single-crystal X-ray analysis, clearly indicated a complex which contained a new 6-membered chelate ring. Although the X-ray data and the refined structure were of excellent quality, we were concerned about the even slightest possibility of a NH *vs.* O misassignment,^{12,13} and sought additional and independent verification of the connectivity seen in the structure. Thus, the three alternatives **3**, **4**, and **5** in Scheme 3 were initially considered. The 6-ring chelates in these alternatives are formally derived from Au, ethylene, acetonitrile, and one (**3**, **4**) or two (**5**) equiv.



Scheme 3 Possible structures of the metallacyclic Au(III) complex.

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 C₁₆H₁₈AuN₂O 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 ¹H–¹⁵N HMBC NMR spectroscopy. In the spectrum,[¶] 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 ¹H signal of the NH group at $\delta(^1\text{H}) = 10.28$ and with the signal from the metal-lacycle methyl group at $\delta(^1\text{H}) = 2.51$, but not with the other protons of the metallacycle. The other ¹⁵N 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 α to the pyridyl-N atom at $\delta(^1\text{H}) = 9.33$. The ¹⁹F NMR spectrum shows only one signal, arising from the [−]OAc^F anion. So as to further unambiguously distinguish between alternatives **3** and **4**, a NOE experiment was performed. The ¹H–¹H 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 Au(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)° is typical, and the Au–ligand distances Au1–N1, Au1–C9, and Au1–C4 (sp³) are within the ranges that have previously been observed for the related complexes Au(tpy)Me₂,¹⁴ Au(tpy)(CH₂CH₂OAc^F)(OAc^F),¹¹ and Au(tpy)(CH₂CH₂OCH₂CF₃)(OAc^F).¹¹ The N5–C3 (1.284(4) Å) and C3–O2 (1.332(3) Å) bond distances are consistent with the corresponding average N=C (1.282 Å) and C–O (1.325 Å) in Pt(II) imino ether complexes.^{15–22} These distances are, however, distinctly different from the distances in a [(tpa)Rh^{III}]²⁺ (tpa = *N,N,N*-tri(2-pyridylmethyl)amine) analogue to **3**, in which the corresponding N–C and C–O bond distances are 1.245(19) and 1.374(17) 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) 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*

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

^{¶¶}No structural examples of Au imino ethers were found *via* SciFinder.



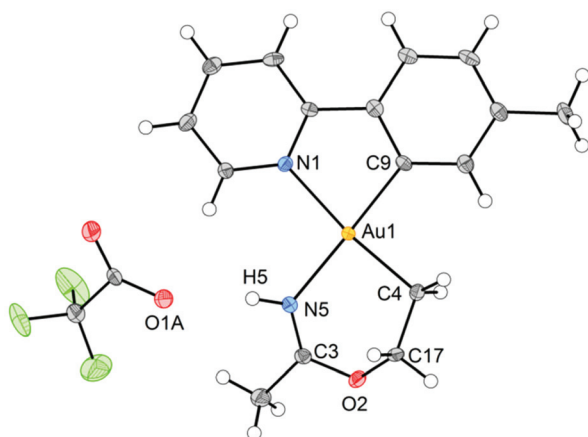


Fig. 1 ORTEP drawing of Au(III) complex **3**, with 50% probability ellipsoids. Selected bond distances (Å), angles, and torsional angles (°): Au1–C4, 2.042(3); Au1–C9, 2.019(3); Au1–N1, 2.126(2); Au1–N5, 2.092(3); C3–N5, 1.284(4); C3–O2, 1.332(3); C17–O2, 1.449(4); C4–C17, 1.510(4); O1A...N5, 2.8708(2); C4–Au1–C9, 92.39(12); C4–Au1–N5, 89.54(11); C9–Au1–N1, 80.60(10); N1–Au1–N5, 97.43(9); C4–Au1–N1, 172.99(11); C9–Au1–N5, 174.24(12); C3–O2–C17–C4, –70.8(4); Au1–C4–C17–O2, 60.2(3); Au1–N5–C3–O2, 26.5(5); C17–O2–C3–N5, 21.8(5).

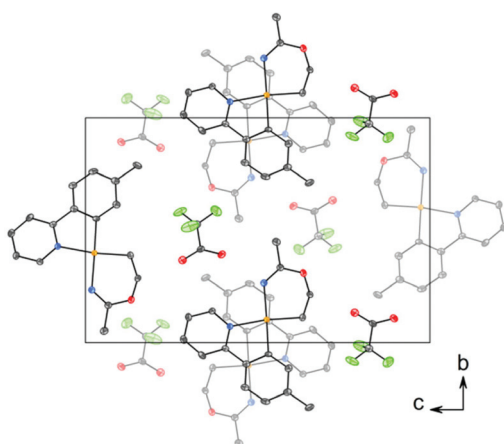
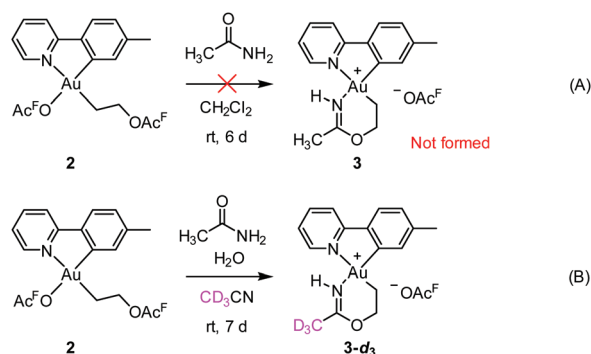


Fig. 2 Packing diagram of structure **3** viewed along the *a* axis, showing stacking between the tpy ligands. The thermal ellipsoids are displayed at 50% probability, and H atoms are omitted for clarity.

axis. The structure displays parallel displaced π - π stacking along the *a* direction, and the distance between the intersecting planes of the tpy ligands alternates between 3.42 and 3.44 Å (Fig. 2).

DFT calculations were performed on the cations of **3**, **4**, and **5** in order to further assist with the verification of the structure. Gratifyingly, comparisons of the crystallographic data with the optimized structures showed that the average r.m.s. deviations of the atomic positions were considerably smaller for **3** (average deviation 0.015 Å) than for **4** (0.022 Å) and **5** (0.032 Å), lending further support for the structural assignment.



Scheme 4 Potential role of acetamide in the formation of **3**.

Mechanistic studies by experiment and DFT calculations

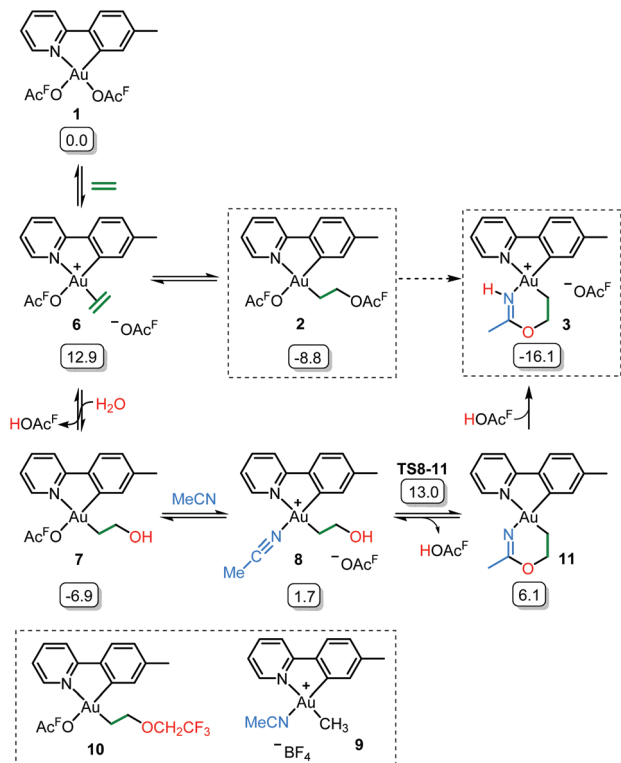
Initial mechanistic experiments probed whether a gold-mediated or gold-catalyzed hydrolysis of acetonitrile to form acetamide might be involved in the process (Scheme 4). First, no formation of **3** was seen when **2** was treated with only acetamide in CH_2Cl_2 (Scheme 4A). Second, the reaction of **2** with wet acetonitrile- d_3 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.¹¹ The hydroxyethyl complex **7** is formed from **2** after nucleophilic attack by water, by analogy with Atwood's study,⁹ and parallel to reaction of trifluoroethanol with **2**, which proceeds *via* **6** yielding **10**.¹¹ Acetonitrile substitution for the OAc^{F} anion of **7** at the kinetically most accessible position¹¹ *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(II)-coordinated nitriles to provide Pt(II) imino ethers.^{20–22} An OAc^{F} 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 N.²⁴ Complex **8** is closely related to [Au(tpy)Me(NCMe)] $[\text{BF}_4]$ (**9**) which has been prepared with the relevant stereochemistry at Au by bromide abstraction from AuBrMe(tpy) with AgBF_4 in acetonitrile.²⁵

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 (TS8-11,

§ For an estimation of the transition state energies involved in the transformation of **1** to **7**, see ref. 11.





Scheme 5 Postulated reaction mechanism for the formation of **3** ($\text{OAc}^{\text{F}} = \text{OCOCF}_3$). Framed species have been characterized by X-ray diffraction in either this or previous (ref. 11) work. Free energies obtained from DFT calculations in acetonitrile are given in kcal mol^{-1} .

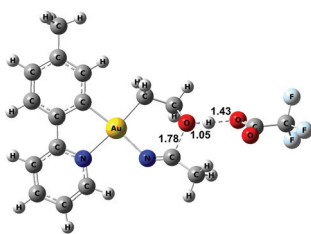


Fig. 3 Optimized geometry for TS8-11 with selected bond distances (Å).

Fig. 3) and it shows that cyclization is triggered by the deprotonation of the hydroxyl group by an external OAc^{F} . This transition state is $13.0 \text{ kcal mol}^{-1}$ higher than **1** and involves an effective energy barrier of $21.8 \text{ kcal mol}^{-1}$ relative to **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 **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(I) catalyzed asymmetric aldol reactions

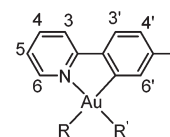
of isocyanides and aldehydes. Hashmi and coworkers more recently^{29,30} prepared oxazolines by Au(I)-catalyzed cycloisomerizations of propargyl amides. Interestingly, it has been recently reported that cyclometallated Au(III) aryl-pyridine complexes may act as efficient catalysts for the three-component 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 $[(\text{tpa})\text{Rh}^{\text{III}}]^{2+}$ analogue to **3** was synthesized in a two-step process by H_2O_2 oxidation of a coordinated ethylene, followed by reaction with acetonitrile and NH_4PF_6 .²³ 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 $\text{Au}(\text{tpy})(\text{OAc}^{\text{F}})_2$ can be utilized – a fact that may be of importance for catalytic applications. The metallacycle is constructed in such a fashion that the high *trans*-effect (alkyl) end of the assembled chelate ligand occupies the position that is *trans* to the low *trans*-effect (N donor) of the tpy supporting ligand, and *vice versa*.

Experimental and computational section

General experimental methods

Gold(III) complexes **1** and **2** were prepared by previously reported procedures.^{11,14} Distilled water was used. CH_3CN and CH_2Cl_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 **9** which is described in the ESI.† NMR spectra were recorded on Bruker Avance DPX200, AVII400, DRX500, AVII600 and AV600 instruments at ambient temperature. ^1H and ^{13}C spectra have been referenced relative to the residual solvent signals. ^{19}F has been referenced to CFCl_3 by using C_6F_6 (-164.9 ppm with respect to CFCl_3 at 0 ppm) as an internal standard. The ^{15}N chemical shifts have been calibrated using MeNO_2 as an external standard at 0 ppm . The peaks in the ^1H NMR and ^{13}C NMR spectra were assigned by the aid of 2D NMR techniques such as HSQC, HMBC, COSY, NOESY, and ^1H - ^{15}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 Microanalytisches Laboratorium Kolbe, Mülheim an der Ruhr, Germany.



Preparation of $[\text{Au}(\text{tpy})(\text{C-N})^+][\text{OCOCF}_3]^-$ (3) from 1. $\text{Au}(\text{tpy})(\text{OCOCF}_3)_2$ (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 $[\text{Au}(\text{tpy})(\text{C-N})^+][\text{OCOCF}_3]^-$ (3) from 2. $\text{Au}(\text{tpy})(\text{CH}_2\text{CH}_2\text{OCOCF}_3)(\text{OCOCF}_3)$ (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%). $^1\text{H NMR}$ (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, $J = 8.0, 7.5, 1.4$ Hz, H^4), 7.96 (d, 1H, $J = 8.0$ Hz, H^3), 7.71 (d, 1H, $J = 7.7$ Hz, H^3), 7.58 (ddd, 1H, $J = 7.5, 5.6, 1.1$ Hz, H^5), 7.23–7.25 (m, 2H, H^4 and H^6), 4.34 (m, 2H, OCH_2), 2.73 (m, 2H, AuCH_2), 2.51 (s, 3H, OCCH_3), 2.44 (s, 3H, ArCH_3). $^{13}\text{C NMR}$ (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, 21.6. $^-\text{OCOCF}_3$ was not observed by $^{13}\text{C NMR}$. $^{15}\text{N}\{^1\text{H}\}$ NMR (600 MHz, CD_2Cl_2): δ -129 (N(tpy)), -212 (NH, $|J| = 78$ Hz), as observed by ^1H - ^{15}N HMBC (see Fig. S12†). $^{19}\text{F NMR}$ (188 MHz, CD_2Cl_2): δ -78.1 (bs, CF_3). **MS** (ESI^+ , MeCN): m/z (rel.%): 451 ($[\text{M} - \text{OCOCF}_3]^+$, 100). **MS** (ESI^- , MeCN): m/z (rel.%): 113 ($[\text{OCOCF}_3]^-$, 100) was observed among other unidentified peaks. **HRMS** (ESI^+ , MeCN): Found: 451.1068; calcd for $\text{C}_{16}\text{H}_{18}\text{AuN}_2\text{O}$: 451.1084 (+0.0016). **HRMS** (ESI^- , MeCN): Found: 112.9859; calcd for $\text{C}_2\text{F}_3\text{O}_2$: 112.9856 (-0.0003). **Elemental analysis**: Anal. Calcd for $\text{C}_{18}\text{H}_{18}\text{AuF}_3\text{N}_2\text{O}_3$: 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.³⁴ The hybrid PBE0+GD3 functional^{35,36} 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(III) alkene complexes.^{8,11,37} C, H, N and O were described with the all-electron triple- ζ 6-311+G** basis set,^{38,39} whereas Au was described with the Stuttgart-Köln basis set including a small-core quasi-relativistic pseudopotential.^{40,41} 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.⁴² Complexes 3, 4 and

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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