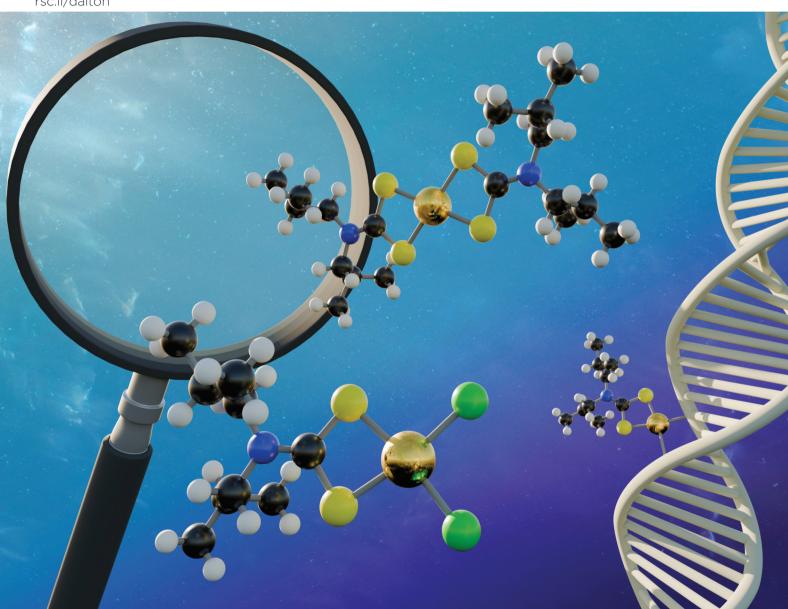
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dithiocarbamate complexes in solution†

A revised understanding of the speciation of gold(III)

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The cytotoxic series of cis-platin mimics "[AuX2(dtc)]" (X = Cl, Br; dtc = dithiocarbamate) were recently patented as promising anticancer metallotherapeutics. Using a range of dialkyl-, cyclic alkyl- and diaryldithiocarbamate ligands, we have discovered that "[AuX2(dtc)]" actually exist in solution as a mixture containing neutral [AuX₂(dtc)] and cationic [Au(dtc)₂]⁺. For the latter, single crystal X-ray crystallography proved that a variety of halide-containing anions such as $[AuX_4]^-$, $[AuX_2]^-$ and even X^- balanced the charge. Based on a thorough investigation into the synthesis of these compounds, we discovered that literature syntheses which claim to produce pure material in fact generate mixtures. In some cases the major component of the mixture is actually the cationic [Au(dtc)₂]⁺ rather than the claimed neutral [AuX₂(dtc)]. Refinement of the synthetic conditions led to a mixture where the neutral [AuX₂(dtc)] was the dominant component, from which pure solid $[AuX_2(dtc)]$ could be obtained by fractional crystallisation. However, the isomerisation process immediately restarted upon dissolution of the crystalline material, thus it is not possible to obtain pure $[AuX_2(dtc)]$ in solution. This discovery has important ramifications for any future use of these compounds, especially as therapeutics since the solution-phase speciation means that "pure" [AuX2(dtc)] cannot exist under biologically relevant conditions. A critical reinterpretation of existing literature data demonstrates that there is already significant uncertainty surrounding which component(s) of this mixture are biologically active.

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Introduction

Since the Thalidomide disaster it has been discovered that many pharmaceutically active molecules have isomers which are highly toxic, and there are plenty of documented cases where a licensed pharmaceutical can convert to a toxic isomer *in vivo*. Thus, a thorough understanding of the solution-phase speciation of therapeutic molecules is required to ensure that no toxic isomers are formed under biologically relevant conditions. Recently, compounds of the general formulation $[AuX_2(dtc)]$ (X = Cl, Br; dtc = dithiocarbamate; Fig. 1) have gained a lot of attention as potential new metallopharmaceuticals. Structurally these compounds are very similar to cisplatin and they are considered extremely active antitumour

The established syntheses of "[AuX₂(dtc)]" are supposedly straightforward and have been known since the 1960s (Scheme 1).¹⁰ The generally accepted understanding of the reactivity, across a vast range of dtc ligands, posits that the

$$\begin{array}{c} X_{MM}, \\ X \end{array} \qquad \begin{array}{c} X_{MM}, \\ X \end{array} \qquad \begin{array}{c} X_{MM}, \\ X_{MM},$$

Fig. 1 A selection of " $[AuX_2(dtc)]$ " complexes which have demonstrated anti-cancer activity (X = Cl, Br).

agents against cultured human cancer cell lines.²⁻⁶ Although this family of compounds has been known for decades, their applications have only just started to emerge: in addition to metallopharmaceuticals, they have also found potential uses as ¹¹C-radiolabelled PET tracers,⁷ homogeneous catalysts,⁸ and molecular precursors for the deposition of gold monolayers by Atomic Layer Deposition.⁹

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$$Na \longrightarrow NR_{2} \longrightarrow NR_{2} \longrightarrow NR_{2} \longrightarrow NR_{2} \longrightarrow NR_{2}$$

$$S \longrightarrow Au \longrightarrow S$$

$$S \longrightarrow Au \longrightarrow S$$

$$S \longrightarrow NR_{2} \longrightarrow NR_{2}$$

$$X \longrightarrow Au \longrightarrow S$$

Scheme 1 The two most common synthetic routes towards "[Au X_2 (dtc)]" by salt metathesis (top) and oxidative addition (bottom). X =Cl or Br; R₂ = dialkyl, cyclic alkyl, diaryl.

addition of one molar equivalent of M(dtc) (M = Li⁺, Na⁺, etc.) to a solution of "AuX₃" (X = Cl, Br)§ leads to the sole formation of neutral "[AuX2(dtc)]", whereas addition of two molar equivalents of M(dtc) results in formation of the cationic "[Au(dtc)₂]⁺", supposedly with X⁻ as anion. 11 Alternatively, oxidation of the Au(1) species $[Au(dtc)]_n$ with elemental X_2 is claimed to generate "[AuX2(dtc)]" as a single product.10 Reported yields are almost quantitative but characterisation data are often sparse, especially in the older literature where high resolution NMR spectroscopy was not routinely available.

Although the existing data are generally consistent with the proposed product formulations, solution-phase anomalies have been noted in some recent publications. For example, "[AuBr₂(MSDT)]" (MSDT = methylsarcosinedithiocarbamate) clearly showed two sets of resonances in its 1H and ¹³C{¹H} NMR spectra. ¹² The second set of resonances were initially misassigned as a "hypothesized isomerism in solution" caused by restricted rotation around the C-N bond of the dtc ligand, 13 but were later reassigned as the putative dimeric species [AuBr₂(µ²-dtc)]₂ on the basis of infra-red spectroscopic evidence.¹⁴ Similarly, the reported synthesis of "[AuCl₂(Bz₂dtc)]" (Bz₂dtc = dibenzyldithiocarbamate) clearly showed two sets of dtc-based resonances in the 1H and ¹³C{¹H} NMR spectra (see Fig. S5–S7 from the ESI of ref. 8) although no explanation was given. Finally, [Au(dtc)₂] was observed in the mass spectra of some [AuX2(dtc)] species; this was explained as decomposition in solution, although no NMR data was presented to corroborate this.15

We were seeking a high-yielding synthesis of pure [AuX2(dtc)] using dialkyl-, cyclic alkyl-, and diaryl-dithiocarbamate ligands (Fig. 2). Following the published synthetic procedures we also observed anomalies with the generally understood reactivity of this family of compounds. Hence, we conducted a detailed study into the synthesis of "[AuX₂(dtc)]", the results of which are reported herein. The solution-phase behaviour of this long-established series of supposedly pure

§ Where the formula of a compound is given inside quotation marks, it indicates that the exact speciation of this compound is not known. Gold dithiocarbamate complexes are discussed extensively in the main text, but "AuX3" indicates that the exact nature of the gold(III) halide reagent is not clear. Many publications interchangeably refer to anhydrous AuX3, hydrated AuX3, AuX3 dissolved in aqueous HX, H[AuX4], and even [AuX4] salts, and it is not always clear which of these Au(III) compounds have been used. To avoid lengthy discourse, these have been simplified to "AuX₃" as a generic term for a source of gold(III) halide.

"[AuX₂(dtc)]" complexes synthesized in this paper (X = Cl, Br).

compounds is not as simple or as well-understood as previously thought, and our findings have important implications for any future application of these compounds, especially as metallopharmaceuticals.

Results and discussion

Identifying the mixture

There are many reported syntheses of "[AuX2(dtc)]" using a range of reaction conditions. Chlorinated solvents e.g. CH₂Cl₂, CHCl₃, 10 and CCl₄, 8 can be employed (sometimes in combination with other solvents such as acetone), but addition of an antisolvent is required to precipitate the "[AuX₂(dtc)]" product. Alternatively, water can be used as solvent: a yellow powder, claimed to be "[AuX2(dtc)]", precipitated upon addition of an aqueous solution of Na(dtc) to an aqueous solution of "AuX₃". 12-14

Thus, following the method of Nilakantan et al., 16 addition of an aqueous solution of Na(Et2dtc) to aqueous K[AuCl4] immediately led to the precipitation of a vellow solid (Scheme 2). However, close analysis of the ¹H NMR spectrum of this solid clearly showed the presence of two overlapping Et2dtc-containing products at very similar chemical shift values (Fig. 3). The major set of peaks matched the data reported by Nilakantan et al. for "[AuCl2(Et2dtc)]"16 but the minor set did not match any residual starting material (Na(Et₂dtc)), or plausible byproducts such as tetraethylthiuram disulfide. Repeated syntheses varying conditions such as concentration and temperature always resulted in a mixture of products being formed (albeit in slightly differing ratios), and the same product mixture was also observed in the NMR spectra obtained in a variety of deuterated solvents, confirming this mixture was the outcome of the synthetic procedure.

Scheme 2 Synthetic route to "[AuCl₂(Et₂dtc)]" in aqueous conditions.

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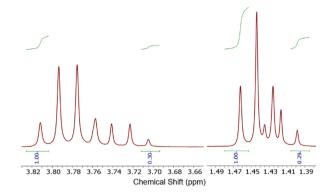


Fig. 3 The ¹H NMR spectrum (in CDCl₃) of the as-synthesized [AuCl₂(Et₂dtc)] mixture focused on the CH₂ and CH₃ regions of the spectrum.

Attempts to purify this mixture by column chromatography were partially successful. The components were clearly separable on silica, forming distinct orange and yellow bands which eluted at different retention times. However, ¹H NMR spectroscopic analysis of the "pure" fractions obtained from the column revealed that a mixture was still present, albeit the ratio of products had altered significantly to favour one component of the mixture. When these NMR sample solutions were rerun after standing for several days, the ratio of products had changed again. Repeated separations on silica resulted in the same outcome. This indicated that "[AuCl₂(Et₂dtc)]" isomerises in solution, rather than undergoing a decomposition process.

To investigate this anomaly further, syntheses of some analogous complexes were conducted under the same reaction conditions. A yellow solid was isolated from the synthesis of "[AuCl₂(Me₂dtc)]" in water, and the ¹H NMR spectrum (DMSO-d₆) contained two resonances which overlapped with the residual HDO peak. In other deuterated solvents (acetone-d₆, CD₃CN) it was clear that two distinct products at very similar chemical shift values were present (see ESI, Fig. S5-S7†). Similarly, the aqueous synthesis of "[AuCl₂(iPr₂dtc)]" and "[AuCl₂(iBu₂dtc)]" both yielded products with ¹H NMR spectra clearly showing two distinct components. For "[AuCl₂(ⁱPr₂dtc)]", comparison of our data to that reported by Angeloski et al. suggested that the mixture contained [AuCl₂(ⁱPr₂dtc)] and [Au(ⁱPr₂dtc)₂][AuCl₄],¹⁷ and an X-ray crystal structure of the former (see ESI, Fig. S1†) was obtained. 18 Fractional crystallization of the "[AuCl₂(ⁱBu₂dtc)]" product mixture afforded two distinct coloured crystals with different morphologies (orange rods and yellow blades) as viewed under an optical microscope. The orange rods corresponded to [AuCl₂(¹Bu₂dtc)] (Fig. 4a) whereas some of the yellow blades were [Au(¹Bu₂dtc)₂][AuCl₂] (Fig. 4b).

[AuCl₂(¹Bu₂dtc)] is a 4-coordinate square planar species with two symmetry-independent molecules in the asymmetric unit. The Au-S and Au-Cl distances are consistent with analogous distances in other [AuCl₂(dtc)] structures, ¹⁵ but there are no aurophilic interactions. 19 Instead, there are short CH···Au

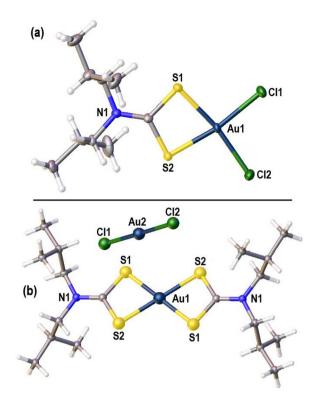


Fig. 4 (a) Solid-state molecular structure of [AuCl₂(iBu₂dtc)] showing the Au1-centred molecule. Selected bond lengths (Å): Au-S1 2.2946(14); Au-S2 2.2948(13); Au-Cl1 2.3240(14); Au-Cl2 2.3092(14); (b) solid-state molecular structure of [Au(iBu2dtc)2][AuCl2]. Range of Au-S bond lengths (A): 2.3236(6)-2.3414(6).

intermolecular contacts between neighbouring pairs of molecules. The approximate CH···Au distances of 2.951 and 3.223 Å are 0.3-0.6 Å less than the sum of van der Waals radii for Au and H,20 and are consistent with stated literature examples of CH···Au(III) hydrogen bonding.21 The concept of CH···Au(III) hydrogen bonding is still controversial, 22-24 but the approximate CH···Au angles in [AuCl₂(ⁱBu₂dtc)] of 153.16° and 149.50° indicate significant directionality to this interaction.

[Au(¹Bu₂dtc)₂][AuCl₂] contains two symmetry-independent ion pairs in the asymmetric unit, with distinct square planar Au(III) and linear Au(I) environments. The Au-S bonds are slightly longer than those found in [AuCl₂(¹Bu₂dtc)], but are consistent with other Au-S bonds from various [Au(dtc)₂] cations. 15 No CH...Au interactions or aurophilic interactions are present in this structure. Both compounds are unknown in the literature, but the related species [Au(¹Bu₂dtc)₂][AuCl₄] was isolated by Rodina et al. from the reaction of [Cd(¹Bu₂dtc)₂] with "AuCl3" and we were able to match the unit cell parameters from other yellow blade crystals in the same batch to this structure.²⁵ The [AuCl₂] anion likely originated from a background redox process where the dithiocarbamate ligand is oxidized to a thiuram disulfide with concomitant reduction of Au(III) to Au(I).

Similar experiments with other types of dtc ligand were conducted: "[AuCl₂(pyrr-dtc)]" and "[AuCl₂(pip-dtc)]", with cyclic

dtc ligands based on pyrrolidine and piperidine respectively, were synthesized by reacting K[AuCl₄] with the respective Na(dtc) salts in water.²⁶ The complex multiplets observed for the methylene protons in the ¹H NMR spectra of the products precluded the identification of a mixture, although they were consistent with literature data for "[AuCl₂(dtc)]". ²⁶ However, the ¹³C{¹H}NMR spectra of both mixtures clearly showed the presence of two distinct environments for each of the methylene carbons in the ring (see ESI, Fig. S16 and S18†). It was also possible to obtain crystals from both reaction products and the synthesis of [AuCl₂(dtc)] was confirmed by matching our unit cell parameters to literature data. ^{27,28} Poor-quality crystals of the corresponding [Au(dtc)₂]⁺ cations with anions such as [AuCl₄]⁻, [AuCl₂]⁻ and Cl⁻ were also grown. These datasets were not of great quality, but the atomic connectivity was unambiguous and we were able to match our unit cell parameters to existing data for some cationic structures.²⁸ Therefore "[AuCl2(dtc)]" with cyclic dtc ligands also exist as mixtures of products rather than a single species.

Diaryl dithiocarbamates are much less widely studied than their dialkyl analogues, although they are of interest as ligands in coordination chemistry, notably as single-source precursors for metal sulfide materials.^{29–31} The reaction of Li(*p*-tolyl₂dtc) with K[AuCl₄] afforded a brown solid with a ¹H NMR spectrum clearly showing two distinct compounds. From this mixture two crystal morphologies, orange blocks and yellow plates, were grown. The orange blocks corresponded to [AuCl₂(p-tolyl₂dtc)], whereas the yellow plates gave the structure of [Au(*p*-tolyl₂dtc)₂][AuCl₂] (Fig. 5).

 $[AuCl_2(p-tolyl_2dtc)]$ exists as a square planar Au(III)species with Au-S and Au-Cl bond lengths consistent with bond lengths from other [AuCl₂(dtc)] structures.¹⁵ [Au(p-tolyl₂dtc)₂][AuCl₂] contains two symmetry-independent square planar Au(III) cations and two linear Au(I) anions in the asymmetric unit, with Au-S bond lengths being consistent with other examples of Au-S bonds in gold dialkyldithiocarbamate complexes. Neither complex exhibits aurophilic interactions, but more interestingly the Au(1) anions appear to be acting as hydrogen bond acceptors to an aromatic CH on a neighbouring cation. The approximate CH···Au distance of 2.760 Å is substantially within the sum of van der Waals radii for H and Au (3.52 Å), 20 albeit the approximate CH···Au angle of 148.64° is not consistent with a strongly directional interaction. It is, however, a significantly more linear interaction than the majority of CH···Au(1) hydrogen bonds which have been noted in the literature.²²

A similar reaction between AuBr₃ and Li(p-tolyl₂dtc) also afforded a brown precipitate of "[AuBr₂(p-tolyl₂dtc)]" with two distinct compounds in its ¹H NMR spectrum. Two crystalline products were isolated, orange rods and yellow plates, with both morphologies giving viable structure solutions. Further crystallographic discussion of these compounds is in the ESI (Fig. S3†).

In light of the X-ray diffraction results, it is clear that both the neutral (orange) [AuX₂(dtc)] and the cationic (yellow) [Au(dtc)₂]⁺ can be co-crystallized from the same reaction mixture with a variety of different dtc and halide ligands. To the best of our knowledge, both series of compounds are indefinitely stable in the solid phase.

Assigning the components of the mixture

With these results in mind, we re-examined the ¹H NMR spectroscopic data of the "[AuCl₂(Et₂dtc)]" mixture (Fig. 3). Although ¹H NMR spectroscopic data for [Au(Et₂dtc)₂]Cl had been reported in the literature, 15 to avoid any potential isomerisation issues we synthesized [Au(Et2dtc)2][BF4] as a single pure compound, including its crystal structure (see ESI, Fig. S4†), by addition of two equivalents of Na(Et2dtc) to K [AuCl₄] in the presence of Na[BF₄] as a halide abstractor (Scheme 3).

We monitored [Au(Et₂dtc)₂][BF₄] in solution and discovered that it is indefinitely stable as a single species (i.e. it does not

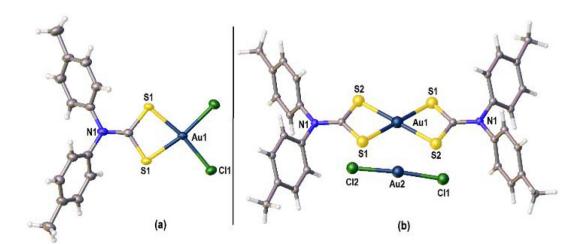


Fig. 5 Solid-state molecular structure of (a) [AuCl₂(p-tolyl₂dtc)₂]. Selected bond lengths: Au-Cl 2.3167(11), Au-S 2.2979(12) Å; (b) [Au(p-tolyl₂dtc)₂][AuCl₂]. Range of Au–S bond lengths (Å): 2.3326(6)–2.3399(6).

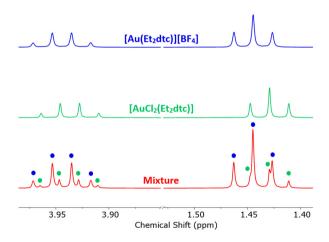
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$$2 \begin{bmatrix} Na & S \\ S & ---- NEt_2 \end{bmatrix} \underbrace{Na[BF_4]}_{-3NaCl} = \underbrace{Et_2N - --- S}_{S} \underbrace{Nu_1 & \oplus \\ Na[BF_4] & ---- NEt_2 \end{bmatrix}}_{-3NaCl} = \underbrace{Na[BF_4]}_{-3NaCl} = \underbrace{Na[BF_4]}_$$

Scheme 3 Synthesis of [Au(Et₂dtc)₂][BF₄] (reaction conducted in acetone)

isomerise in solution), but to our surprise we matched its ¹H NMR spectrum to the *major* peaks in the "[AuCl₂(dtc)]" mixture (Fig. 6). This indicated that - under the reaction conditions reported by Nilakantan et al. - the major product which precipitated from aqueous solution is actually cationic [Au(Et₂dtc)₂]⁺ rather than the stated *neutral* product [AuCl₂(Et₂dtc)].¹⁶ It is worth noting that the solid product which precipitated from water was yellow (both in our hands and also as reported by Nilakantan et al.), and all crystals of cationic [Au(dtc)₂]⁺ that we obtained were also yellow. This contrasts to the neutral species which formed orange crystals. Partial integration of the non-overlapping peaks suggested that the ratio of products was roughly 5:3 $[Au(Et_2dtc)_2]^+$: $[AuCl_2(Et_2dtc)]$.

Although we had spectroscopic evidence for the formation of a mixture in the "[AuCl₂(Me₂dtc)]" system, it is noteworthy that the reported ¹H NMR data for ionic [Au(Me₂dtc)₂]Cl is identical to that for neutral [AuCl₂(Me₂dtc)], ¹⁵ and it also overlaps with the HDO signal in DMSO-d₆. Given the importance of both [AuCl₂(Me₂dtc)] and [AuBr₂(Me₂dtc)] as potential metallopharmaceuticals, we revisited the "[AuX2(Me2dtc)]" (X = Cl, Br) system using AuBr₃ rather than $K[AuCl_4]$ as starting material. Two distinct peaks in the ¹H NMR spectrum of the "[AuBr₂(Me₂dtc)]" mixture synthesized by precipitation from water, as well as two CH₃ peaks in the ¹³C{¹H} NMR spectrum (which were correlated by HMBC spectra) were observed, con-



¹H NMR spectra Fig. 6 Portions of the (acetone-d₆) [Au(Et₂dtc)₂][BF₄] (blue, top); [AuCl₂(Et₂dtc)] (green, middle) and the assynthesized mixture which precipitated from water (red, bottom) with the coloured circles indicating which peaks correspond to which component of the mixture.

firming that the Me₂dtc system also forms a mixture (see ESI, Fig. S21 and 22; also Fig. S6 for X-ray structures†). Here we note that the "[AuX₂(Me₂dtc)]" system is not particularly soluble and we were unable to see the 13C signal for the dithiocarbamate carbon by 13C{1H} NMR spectroscopy. However, HSOC spectra on the "[AuCl₂(Me₂dtc)]" system exhibited two distinct correlations between a broad peak for the CH₃ protons (overlapping *neutral* and *cationic*) and the separate dithiocarbamate carbons which could not be seen by ¹³C{¹H} NMR spectroscopy (see ESI, Fig. S49†).

To explain these findings, we monitored the synthesis of "[AuCl₂(Et₂dtc)]" by ¹H NMR spectroscopy. A solution of Na (Et₂dtc) in DMSO-d₆ was added portionwise to a solution of K [AuCl₄] in DMSO-d₆ at RT, and a series of ¹H NMR spectra were obtained (Fig. 7). The spectra clearly show that as soon as Na(Et2dtc) is added a mixture of neutral [AuCl2(Et2dtc)] and cationic [Au(Et2dtc)2] is formed, even though K[AuCl4] is in excess, and that the majority product is the cationic species $[Au(Et_2dtc)_2]^+$ rather than the desired neutral $[AuCl_2(Et_2dtc)]$. Further additions of Na(Et2dtc) do not significantly affect the ratio of products formed, and after complete addition of $Na(Et_2dtc)$ a product ratio of 7:5 cationic: neutral was obtained.

A critical reinterpretation of existing literature data shows that there are indications, even in the older literature, that the synthesis of "[AuX2(dtc)]" always forms a mixture rather than a pure compound. For example, van der Kerk and co-workers identified that the oxidation product of $[Au(R_2dtc)]_n$ with elemental I_2 (R = Me, Et, ⁿPr, ⁿBu) led to the formation of a putative Au(II) species $[AuI(R_2dtc)]^{10}$. In reality this was likely to be a mixture of [AuI₂(R₂dtc)] and [Au(R₂dtc)₂][AuI₂]. Oxidation of $[Au(R_2dtc)]_n$ with an excess of the other halogens led to the isolation of yellow solid material, assigned as "[AuX₂(R₂dtc)]". Based on our observations (above), yellow compounds indicated the presence of cationic [Au(dtc)₂]⁺ hence we repeated this reaction using $[Au(Et_2dtc)]_n$ and an excess of Br₂ in CH₂Cl₂. After complete addition of Br₂, the

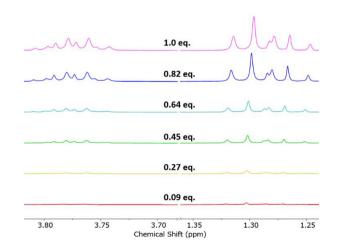


Fig. 7 Stacked plot of ¹H NMR spectra (DMSO-d₆) following the portionwise addition of Na(Et2dtc) to Na[AuCl4]. The majority product is $[Au(Et_2dtc)_2]^+$ and the minority product is $[AuCl_2(Et_2dtc)]$.

¹H NMR spectrum of the product mixture clearly showed two distinct sets of peaks, one of which was matched to $[Au(Et_2dtc)_2][BF_4]$ (*vide supra*). We were able to grow two distinct crystal morphologies from the mixture: orange rods corresponding to $[AuBr_2(Et_2dtc)]$ and yellow plates of $[Au(Et_2dtc)_2][AuBr_2]$ (Fig. 8, also see ESI† for further discussion). These structures provide further confirmation that both the neutral and cationic products are always present when " $[AuX_2(dtc)]$ " is synthesized.

Other authors have isolated both neutral [AuX₂(dtc)] and cationic [Au(dtc)₂]⁺ components from the same reaction mixture. Radanović *et al.* obtained [AuX₂(Me₂dtc)] and [Au(Me₂dtc)₂]X (X = Cl, Br) from the oxidative addition of tetramethylthiuram disulfide to [AuX₃(N-methylimidazole)].³² Loseva *et al.* structurally characterized a co-crystal containing two molecules of [AuCl₂(i Pr₂dtc)] and three [Au(i Pr₂dtc)₂]⁺ cations, with one [AuCl₂]⁻ anion and two [ClO₄]⁻ anions to balance the charge.³³ Finally, during the course of this work,

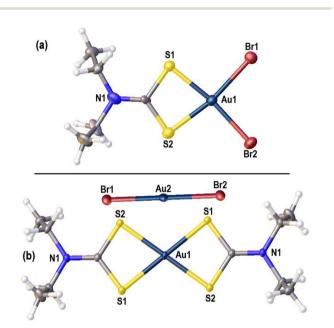


Fig. 8 (a) Solid-state molecular structure of [AuBr₂(Et₂dtc)] showing the Au1-centred molecule. Range of selected bond lengths (Å): Au–S 2.288(6)–2.329(6); Au–Br 2.440(3)–2.451(3); (b) solid-state molecular structure of [Au(Et₂dtc)₂][AuBr₂]. Selected bond lengths (Å): Au1–S1 2.3261(6); Au1–S2 2.3216(5).

Angeloski *et al.* reported that the addition of auric acid to a solution of $Na(^iPr_2dtc)$ in a 1:1 ratio formed " $[Au(^iPr_2dtc)_2][AuCl_4]$ ". Their 1H NMR spectra clearly show a mixture of the cationic species and $[AuCl_2(^iPr_2dtc)]$ was formed, with the cationic species as the majority product. ¹⁷

We were able to conclusively demonstrate that the "[AuX₂(dtc)]" system always exists as a mixture by reacting pure [Au(Et₂dtc)₂][BF₄] (synthesized as above) with one equivalent of K[AuCl₄], forming [Au(Et₂dtc)₂][AuCl₄] *in situ* (Scheme 4). The reaction was monitored by 1H NMR spectroscopy: after 1 day small shoulders corresponding to the neutral species had appeared and, over time, the peaks corresponding to the cationic species dropped in intensity and peaks corresponding to the neutral species grew (Fig. 9). This indicates that the [AuX₂(dtc)]/[Au(dtc)₂]⁺ system – with Au-containing anions that can interconvert – always exists as a mixture rather than as a single pure species, irrespective of how it is synthesized.

It is clear that – in solution – the series of supposedly "pure" compounds " $[AuX_2(dtc)]$ " actually exist as a mixture containing $[AuX_2(dtc)]$ and $[Au(dtc)_2]^+$ (with the charge balanced by a variety of anions), presumably a result of the dissociation of labile dithiocarbamate ligands. A critical reappraisal of existing literature data, coupled to repeated syntheses of selected published procedures, is strongly indicative that the series of compounds " $[AuX_2(dtc)]$ " have always been synthesized as mixtures rather than as pure individual species, despite many claims to the contrary. Indeed, some reported literature syntheses of "pure" $[AuX_2(dtc)]$ actually generate a mixture where the $[Au(dtc)_2]^+$ cation is the major component!

Identification of this mixture is not straightforward, even with routine access to high-resolution NMR spectroscopy, as the signals for neutral and cationic species often overlap. However, 2D NMR spectroscopy (notably HSQC) allows for the observation of distinct correlations between the alkyl/aryl substituents and the dithiocarbamate carbons for both neutral and cationic compounds, and this can be used as a diagnostic indicator for the presence of both species in a sample even if the dithiocarbamate carbon cannot be observed using 1D $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy. UV-vis spectroscopy can also be used as a diagnostic tool since the UV-vis spectrum of the cationic species has a characteristic secondary maximum absorption ($\lambda_{\text{max}} \sim 287 \text{ nm}$) which is much less pronounced in the neutral complex (see ESI, Fig. S66–S68†). However, it cannot be used to accurately quantify the components of the mixture

$$\mathsf{Et_2N} = \underbrace{\left\{ \begin{array}{c} \mathsf{S} / / / / \\ \mathsf{S} \end{array} \right\}}^{\left[\mathsf{BF_4}\right] \ominus} \underbrace{\left\{ \begin{array}{c} \mathsf{Na[AuCl_4]} \\ \mathsf{-Na[BF_4]} \end{array} \right\}}_{-\mathsf{Na[BF_4]}} \mathsf{Et_2N} = \underbrace{\left\{ \begin{array}{c} \mathsf{S} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{Cl} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl} / / / / \\ \mathsf{S} \end{array} \right\}}_{-\mathsf{Na[Au]}} \underbrace{\left\{ \begin{array}{c} \mathsf{Cl}$$

Scheme 4 Regeneration of the [AuCl₂(Et₂dtc)]/[Au(Et₂dtc)₂]⁺ mixture by metathesizing stable [Au(Et₂dtc)₂][BF₄] with Na[AuCl₄].

complex, and ultimately isolating this compound as a single species.

Our as-synthesized mixture of "[AuCl₂(Et₂dtc)]" (with a product ratio of 5 + 3 entionic pourts)) was discolved in both

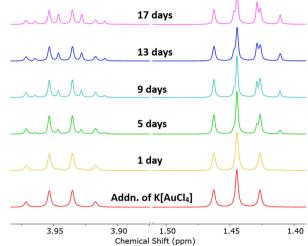


Fig. 9 Stacked plot of ¹H NMR spectra in acetone-d₆ following the addition of K[AuCl₄] to [Au(Et₂dtc)₂][BF₄] (red, bottom) at 4 day intervals.

since the mixture contains multiple anions ($[AuCl_4]^-$, $[AuCl_2]^-$, Cl^-) in unknown concentrations.

Here, we also note that elemental analysis (EA) is not an appropriate bulk characterisation method for these compounds since the neutral and cationic species with $[AuX_4]^-$ anion have identical empirical formulae, thus they cannot be distinguished by EA. We demonstrated this by obtaining 'acceptable' EA data (*i.e.* $\pm 0.4\%$) for "[AuCl₂(dtc)]" using iPr_2 dtc and iBu_2 dtc ligands, despite submitting a mixture of $[AuCl_2(dtc)]$ and $[Au(dtc)_2][AuCl_4]$ for analysis. A recent patent from Nardon *et al.* also acknowledged this fact: "However, the syntheses therein proposed [of $[AuX_2(dtc)]]$ do not permit to obtain the compounds in pure form, but as a mixture of two different complexes having the same empirical formula. This result was not evident from the elemental analysis". 34

Selective formation of [AuCl₂(dtc)]

In 2005, Ronconi et al. observed a minor impurity in their ¹H NMR spectrum of "[AuX₂(dtc)]" (dtc = MSDT or ESDT), ¹³ which was assigned as the dimeric species $[AuBr_2(\mu^2-dtc)]_2$ based on the observation of a linear "AuBr2" moiety in infra-red spectral data.14 Given our findings above, it is clear that the minor impurity is actually $[Au(dtc)_2]^+$, most likely with an $[AuBr_2]^$ anion which would account for the linear "AuBr2" moiety observed by IR spectroscopy. Interestingly, this supposed "dimer" slowly converted to the "monomer" upon leaving a solution of the mixture in acetone-d₆ to stand for 24 hours. Complete conversion did not occur, but a strong solvent dependence on the interconversion between majority and minority products was noted. Certain solvents (CDCl₃, CD₂Cl₂, acetone-d₆) enabled conversion to the majority product but other solvents (DMSO-d₆, CD₃OD) showed no obvious change after 24 hours. 14 This observation prompted us to investigate the "[AuCl₂(Et₂dtc)]" system in more detail with the aim of converting the mixture towards the desired [AuCl₂(Et₂dtc)]

Our as-synthesized mixture of "[AuCl₂(Et₂dtc)]" (with a product ratio of 5:3 cationic:neutral) was dissolved in both acetone-d₆ and DMSO-d₆, and the behaviour of the system was monitored by ¹H NMR spectroscopy at room temperature (Fig. 10). After 24 hours a shift in the ratio of the two components had taken place, although complete conversion to [AuCl₂(Et₂dtc)] had not occurred. Long-term monitoring by ¹H NMR spectroscopy indicated that the system stopped converting after 39 days in acetone-d₆ (final product ratio 1:2.6 cationic:neutral) but for DMSO-d₆ the process was faster, reaching the end point in 19 days (final ratio 1:7.3 cationic:neutral). We were unable to completely convert the system to [AuCl₂(Et₂dtc)] in either solvent, although it was noted that heating the mixture did increase the rate of conversion.

The precipitation of orange crystals was observed during the isomerisation process in acetone-d₆ (and also CD₂Cl₂) but not in DMSO-d₆. Unit cell parameters of all the crystals we measured matched the cell parameters of [AuCl₂(Et₂dtc)] and no evidence for any other compound was observed. This indicated the neutral species [AuCl2(Et2dtc)] was less soluble in acetone-d₆ than the cationic [Au(Et₂dtc)₂]⁺, potentially enabling the synthesis of pure [AuCl₂(Et₂dtc)]. Thus, addition of an acetone solution of Na(Et2dtc) to an acetone solution of Na[AuCl₄] at 30 mM concentration led to the formation of a dark vellow precipitate. Although the amount of precipitate was less than that obtained from the reaction conducted in water, it was clear from the ¹H NMR spectrum of the product that a mixture of [AuCl₂(Et₂dtc)] and [Au(Et₂dtc)₂]⁺ was still present. Gratifyingly, the ratio was now 9:4 in favour of the neutral complex.

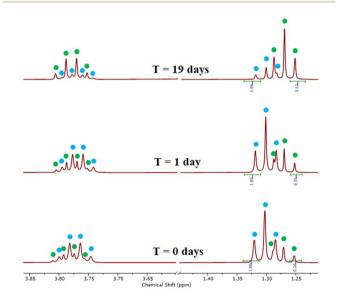


Fig. 10 Comparison of 1H NMR spectra (in DMSO-d₆) following conversion of the "[AuCl₂(Et₂dtc)]" mixture, synthesized from water. Blue dots correspond to peaks from the cationic [Au(Et₂dtc)₂]⁺ species and green dots correspond to peaks from neutral [AuCl₂(Et₂dtc)].

The synthesis of "[AuCl₂(Et₂dtc)]" reported by Cordón et al. was also conducted in acetone, but notably no precipitate was observed when Na(Et2dtc) was added to in situ-generated "AuCl₃". Repeating their procedure did not result in the formation of any precipitate, but ¹H NMR spectroscopy showed the product we obtained was clearly a mixture of [AuCl₂(Et₂dtc)] and [Au(Et₂dtc)₂]⁺, contrary to the claim of a pure product by Cordón et al. Close examination of the ¹H NMR spectra supplied in their ESI (Fig. S3 and S4 from ref. 8) does appear to show small shoulders on the signals associated with the ethyl groups, likely corresponding to minor amounts of [Au(Et₂dtc)₂]⁺ which indicates that they did not obtain pure [AuCl₂(Et₂dtc)] either.

We speculated that the lack of precipitate was likely due to a co-solvent (CCl₄) which was able to solubilise all the reaction products, rather than 3 mL of residual acetone. We adapted this synthesis to replace the toxic acetone/CCl4 mix with DMSO, since both [AuCl₂(Et₂dtc)] and [Au(Et₂dtc)₂]⁺ exhibited high solubility in DMSO-d₆, as well as exhibiting a high neutral: cationic ratio. Thus, upon addition of Na(Et2dtc) to K [AuCl₄] in DMSO, no precipitate was observed. To precipitate the product a significant amount of water was added, but this resulted in a reduced isolated yield (41%). The ¹H NMR spectrum of the precipitated product showed that a mixture was still present, albeit the ratio of [AuCl₂(Et₂dtc)]: [Au(Et₂dtc)]⁺ was now 9:1, higher than our previous syntheses conducted in water and acetone.

Since the strategy of keeping everything dissolved did result in a higher amount of neutral product, albeit a reduced isolated yield with DMSO as solvent, we returned to acetone but at a tenfold-lower concentration of reactants. During the course of this work, Angeloski et al. reported that refluxing a mixture of [AuCl₂(ⁱPr₂dtc)] and [Au(ⁱPr₂dtc)₂]⁺ in acetone rapidly converted the mixture to the point where [AuCl₂(ⁱPr₂dtc)] was the dominant component.¹⁷ Thus, slow addition of Na(Et2dtc) to K[AuCl4] at 3 mM concentration did not generate any precipitate. Refluxing this reaction for 16 hours resulted in a mixture of [AuCl₂(Et₂dtc)] and $[Au(Et_2dtc)_2]^+$ with the vast majority (~17:1) being the neutral species (Fig. 11). Slow evaporation of the solvent led to the precipitation of orange crystals of pure [AuCl₂(Et₂dtc)]: these are indefinitely stable in the solid state, but upon dissolution the isomerisation resumes and the mixture reforms (see ESI, Fig. S65†).

The synthesis detailed above, conducted in dilute refluxing acetone, represents the best approach to generating pure solid [AuX₂(dtc)]. We have used this method to demonstrate that pure, crystalline material of all [AuX2(dtc)] mentioned in this paper can be isolated and 'clean' NMR spectra can be obtained if the pure crystalline solids are dissolved in a deuterated solvent and data obtained immediately (see ESI† for spectra). This may explain the observation of Pettenuzzo et al. who claim to have isolated a single species of "[AuBr₂(dtc)]" (where dtc is derived from the ethyl ester of 4-carboxypyrrolidine) by transmetallation from [Zn(dtc)₂] using AuBr₃.35

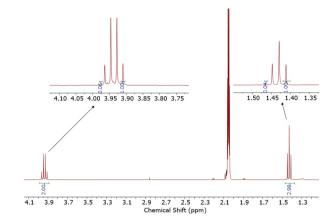


Fig. 11 ¹H NMR spectrum (in acetone-d₆) of the product obtained from the synthesis of "[AuCl2(Et2dtc)]" in dilute refluxing acetone. The majority product is [AuCl₂(Et₂dtc)].

Although the nature of the dtc ligand does have an effect on the ratio of neutral: cationic species observed after heating the reaction mixture (i.e. poorly soluble dtc ligands such as Me₂dtc and pyrr-dtc require longer heating to convert the mixture; see ESI† for details), in all cases we saw a majority of the desired neutral compound from which pure material can be fractionally crystallized. However, it is important to note that if the crystallisation process is too quick (e.g. the vessel has too many nucleation points such as scratches on the glass, or the compound is not especially soluble in acetone) the yellow cationic complex co-crystallises with the orange neutral species. In particular, we observed this for the "[AuBr₂(Me₂dtc)]" system, where fractional crystallisation of a post-refluxed mixture yielded crystals of both neutral [AuBr₂(Me₂dtc)] and ionic [Au $(Me_2dtc)_2$ [AuBr₂] (see ESI, Fig. S6†).

Finally, we note that gold dithiocarbamate complexes in both the +1 and +3 oxidation state are compounds of interest as potent new metallopharmaceuticals. Whilst the speciation of Au(1) dithiocarbamates in solution is not in question,³⁶ we have demonstrated that the same cannot be said about "[AuX2(dtc)]". The use of "[AuX2(dtc)]" as a structural mimic of cis-platin, using various dtc ligands, has been thoroughly investigated and neutral "[AuX2(dtc)]" compounds with simple ligands such as Me2dtc and Et2dtc have been shown to be active against cultured human cancer cell lines. 3,12,26,37,38 Conversely, a recent publication from Fregona and co-workers observed conversion from the neutral "[AuBr₂(dtc)]" to cationic [Au(dtc)₂]⁺ (where dtc is derived from proline alkyl esters) in a 1:9 DMSO/H₂O mix,³⁹ and it was asserted that the cationic species [Au(dtc)₂]⁺ were more cytotoxic than the neutral species owing to "decomposition" of the neutral species supposedly generating the cationic species in situ. 15 Clearly the presence of both components in a product mixture has led to confusing and contradictory cytotoxic data. More work is needed to deconvolute the effects of inadvertently studying the cytotoxicity of a mixture containing both compounds where one, or even both, components of the mixture may be biologically active.

Derivatives of $[AuX_2(dtc)]$ where $X = Me, ^9 C_6F_5, ^8$ mesityl, 40 thiolate, 5 etc. appear to be significantly more stable and do not seem to exist as solution-phase mixtures. Presumably this is due to the reduced stability of the corresponding $[AuX_4]^-$ and $[AuX_2]^-$ anions when X is not a halide. Although we have demonstrated that it is possible to isolate pure $[AuX_2(dtc)]$ (X = halide) in the solid phase by fractional crystallisation, despite many literature reports to the contrary it is not possible to generate pure $[AuX_2(dtc)]$ in solution since mixtures of $[AuX_2(dtc)]$ and $[Au(dtc)_2]^+$ are always obtained.

Conclusions

A detailed investigation into the synthesis of the supposedly pure series of compounds "[AuX₂(dtc)]" (X = Cl, Br; dtc = dithiocarbamate) has been conducted. Contrary to the prevailing understanding of this series of compounds, we discovered that they actually exist as a mixture of multiple species in solution: the desired [AuX₂(dtc)] alongside the [Au(dtc)₂]⁺ cation with a variety of different anions such as [AuX₄]⁻, [AuX₂]⁻ and X⁻ to balance the charge. In some cases, reported literature syntheses of "pure" neutral [AuX₂(dtc)] actually result in the formation of a mixture where cationic [Au(dtc)₂]⁺ is the majority product.

It is possible to vary the synthetic conditions to convert the mixture such that neutral [AuX₂(dtc)] is the majority product but it is not possible to synthesise it as a single species in solution. Fractional crystallisation does yield pure [AuX₂(dtc)] as a crystalline solid. However, it is important to note that [Au(dtc)₂]⁺ is stable as a single species in solution so long as the counteranion is not a haloaurate species. Anion metathesis to [AuCl₄]⁻ subsequently results in the mixture reforming. Many of these compounds exhibit CH····Au interactions in the solid state which are strongly indicative of hydrogen bonding to both Au(i) and Au(iii), and will be of interest to researchers studying these still somewhat controversial interactions.

The supposedly pure "[AuX₂(dtc)]" (X = Cl, Br, I) have been employed in a variety of different applications, but most notably as highly promising metallodrugs against cultured human cancer cell lines. In view of our new insights into the synthesis and speciation of this series of compounds in solution, great care needs to be taken with any future studies into this mixture. A critical reappraisal of the literature has already highlighted uncertainties surrounding the precise nature of the biologically active species and future studies may need to focus on analogues which do not isomerise in solution in order to generate reliable cytotoxic data.

Experimental

The aqueous synthesis of [AuX₂(dtc)]/[Au(dtc)₂]⁺ mixtures

A slight modification of the method published by Nilakantan $et\ al.$ was used. 16

General procedure. An aqueous solution of Na(dtc) or Li(dtc) (1.0 eq., 0.22 M) was added dropwise to an aqueous solution of

 $M[AuCl_4]$ (M = Na, K; 1.0 eq., 0.22 M) with vigorous stirring (AuBr₃ was used for the analogous bromide compounds). A yellow-orange precipitate formed immediately upon addition of the dithiocarbamate salt. After the addition was complete the solution was stirred for 1 h, then isolated by vacuum filtration, washed with distilled water (10 mL) and Et₂O (5 mL). On smaller scales the solid product was dried *in vacuo*, but on larger scales the product was dissolved in CH_2Cl_2 , dried with MgSO₄, filtered and then reisolated by removing all volatiles *in vacuo*.

In most cases, two distinct products were observed in NMR spectra. These have simply been assigned as 'major' and 'minor' components. The analytically pure samples synthesized by a different method have been definitively assigned.

"[AuCl₂(Me₂dtc)]" mixture. Na(Me₂dtc)·2(H₂O) (118 mg, 0.66 mmol, 1.0 eq.) was dissolved in water (5 mL) and added dropwise to a vigorously stirred solution of K[AuCl₄] (250 mg, 0.66 mmol, 1.0 eq.) in water (3 mL) over 15 minutes. The solution was then stirred for 1 h. A yellow precipitate was isolated by vacuum filtration, washed with distilled water $(3 \times 5 \text{ mL})$ followed by Et₂O (2 × 5 mL) and dried in vacuo. Yield: 215 mg of a vellow solid. ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.38 (s, overlapping with H₂O peak, 12H, CH₃, major product), 3.35 (s, 6H, CH₃, minor product). ¹H NMR (400.1 MHz, CD₃CN): δ 3.33 (s, 12H, CH₃, major product), 3.31 (s, 6H, CH₃, minor product). ¹H NMR (400.1 MHz, (CD₃)₂CO): δ 3.55 (s, 12H, CH₃, major product), 3.55 (s, 6H, CH₃, minor product). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): 40.29 (s, CH₃), 40.00 (s, overlapping with solvent peak, CH₃). No peak was observed for the quaternary carbon centre.

"[AuCl₂(Et₂dtc)]" mixture. Na(Et₂dtc)·3(H₂O) (119 mg, 0.53 mmol, 1.0 eq.) was dissolved in water (3 mL) and added dropwise to a vigorously stirred solution of K[AuCl₄] (200 mg, 0.53 mmol, 1.0 eq.) in water (2 mL) over 15 minutes. The suspension was then stirred for 1 h. A yellow precipitate was isolated by vacuum filtration, washed with water (3 × 5 mL) and Et_2O (2 × 5 mL) then dried in vacuo. Yield: 152 mg of a yellow solid. ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.79 (overlapping q, $^{3}J_{HH}$ = 7.2 Hz, 4H, CH₂, minor product), 3.78 (overlapping q, ${}^{3}J_{HH}$ = 7.2 Hz, 8H, CH₂, major product), 1.3 (t, ${}^{3}J_{HH}$ = 7.2 Hz, 12H, CH₃, major product), 1.27 (t, ${}^{3}J_{HH}$ = 7.2 Hz, 6H, CH₃, minor product). ¹H NMR (400.1 MHz, CDCl₃): δ 3.81 (overlapping q, ${}^{3}J_{HH}$ = 7.3 Hz, 8H, CH₂, major product), 3.76 (overlapping q, ${}^{3}J_{HH}$ = 7.3 Hz, 4H, CH₂, minor product), 1.46 (t, ${}^{3}J_{HH}$ = 7.3 Hz, 12H, CH₃, major product), 1.43 (t, ${}^{3}J_{HH}$ = 7.3 Hz, 6H, CH₃, minor product). ¹H NMR (400.1 MHz, (CD₃)₂CO): δ 3.94 (overlapping q, ${}^{3}J_{HH}$ = 7.3 Hz, 8H, CH₂, major product), 3.93 (overlapping q, ${}^{3}J_{HH}$ = 7.3 Hz, 4H, CH₂, minor product), 1.44 (t, $^{3}J_{HH}$ = 7.3 Hz, 12H, CH₃, major product), 1.43 (t, $^{3}J_{HH}$ = 7.3 Hz, 6H, CH₃, minor product).

An alternative synthesis for "[AuCl₂(Et₂dtc)]" was also attempted whereby all reagents were added together in EtOH/ water (1:1 volume), then heated to reflux for 1 h at 110 °C. ¹⁶ The solution was then cooled and the solid isolated by filtration. This was washed with distilled water and Et₂O, then dried *in vacuo*. ¹H NMR spectroscopy indicated the presence of the same product mixture.

"[AuCl₂(iPr₂dtc)]" mixture. Na(iPr₂dtc) (120 mg, 0.60 mmol, 1.0 eq.) was dissolved in water (5 mL) and added dropwise to a vigorously stirred solution of K[AuCl₄] (250 mg, 0.60 mmol, 1.0 eq.) in water (2 mL) over 15 minutes. The solution was then stirred for 1 h. A yellow precipitate was isolated by vacuum filtration, washed with water (3 × 5 mL) followed by Et₂O (2 × 5 mL) and dried in vacuo. Yield 235 mg of a yellow solid. X-ray quality crystals were obtained through vapour diffusion of pentane into a concentrated CH2Cl2 solution. ¹H NMR (400.1 MHz, DMSO-d₆): δ 4.64 (overlapping sept, $^{3}J_{HH}$ = 6.8 Hz, 4H, CH, major product) 4.55 (overlapping sept, ${}^{3}J_{HH}$ = 6.7 Hz, 2H, CH, minor product), 1.48 (d, ${}^{3}J_{HH}$ = 6.8 Hz, 24H, CH₃, major product), 1.44 (d, ${}^{3}J_{HH}$ = 6.8 Hz, 12H, CH₃, minor product). ${}^{13}C{}^{1}H$ NMR (100.6 MHz, DMSO-d₆): δ 193.06 (s, quaternary), 185.69 (s, quaternary), 55.05 (s, CH), 54.72 (s, CH), 19.60 (s, CH₃). Elemental analysis: calc. for C₇H₁₄AuCl₂NS₂ (444.16): C 18.93%, H 3.18%, N 3.15%. Found: C 18.68%, H 2.93%, N 3.30%.

"[AuCl₂(¹Bu₂dtc)]" mixture. K(¹Bu₂dtc) (161 mg, 0.66 mmol, 1.0 eq.) was dissolved in water (3 mL) and added dropwise to an aqueous solution (2 mL) of K[AuCl₄] (250 mg, 0.66 mmol, 1.0 eq.) whilst vigorously stirring. A yellow precipitate formed, and the suspension was further stirred for 1 hour. The solid was filtered, washed with water (3 \times 5 mL) and Et₂O (2 \times 5 mL) then dried in vacuo. Yield: 171 mg of a yellow solid. X-ray quality crystals were obtained through vapour diffusion of pentane into a concentrated CH₂Cl₂ solution. ¹H NMR (400.1 MHz, CDCl₃): δ 3.56 (d, ${}^{3}J_{HH}$ = 7.6 Hz, 4H, CH₂, minor), 3.48 (d, ${}^{3}J_{HH}$ = 7.6 Hz, 8H, CH₂, major), 2.31–2.18 (overlapping m, 3H, CH, major and minor), 1.03 (overlapping d, ${}^{3}J_{HH} = 6.7$ Hz, 12H, CH₃, major), 1.02 (overlapping d, ${}^{3}J_{HH}$ = 6.7 Hz, 24H, CH₃, minor). ¹³C{¹H} NMR (100.6 MHz, CDCl₃): δ 194.79 (s, quaternary), 58.52 (s, CH₂), 27.26 (s, CH), 20.17 (s, CH₃). Elemental analysis: calc. for C₉H₁₈AuCl₂NS₂ (472.21): C 22.89%, H 3.84%, N 2.97%. Found: C 23.06%, H 3.95%, N 2.96%.

"[AuCl2(pyrr-dtc)]" mixture. Na(pyrr-dtc) (56 mg, 0.33 mmol, 1.0 eq.) was dissolved in water (6 mL) and added dropwise to an aqueous solution (4 mL) of K[AuCl₄] (125 mg, 0.33 mmol, 1.0 eq.) with vigorous stirring. A yellow precipitate formed, and the suspension was further stirred for 1 hour. The solid was then isolated by filtration, washed with water (3 × 5 mL) and Et_2O (2 × 5 mL) and dried in vacuo. Yield: 123 mg of a yellow solid. The ¹H NMR spectrum for this product agrees with the chemical shift assignments in previously reported spectra,²⁶ although the authors describe singlet multiplicity in both peaks instead of the expected complex multiplets observed here. Crystals were obtained through vapour diffusion of pentane into a concentrated CH₂Cl₂ solution. ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.85-3.80 (m, 4H, NCH₂), 2.08-2.04 (m, 4H, CH₂). ${}^{13}C{}^{1}H$ NMR (100.6 MHz, DMSO-d₆): δ 51.11 (s, NCH₂), 51.04 (s, NCH₂), 23.55 (s, CH₂), 23.33 (s, CH₂). No peak was observed for the quaternary carbon centre.

"[AuCl₂(pip-dtc)]" mixture. Na(pip-dtc) (121 mg, 0.66 mmol, 1.0 eq.) was dissolved in water (3 mL) and added dropwise to an aqueous solution (2 mL) of K[AuCl₄] (250 mg, 0.66 mmol,

1.0 eq.) whilst vigorously stirring. A yellow precipitate formed, and the suspension was further stirred for 1 hour. The solid was filtered, washed with water (3 × 5 mL) and Et₂O (2 × 5 mL) then dried *in vacuo*. Yield: 232 mg of a yellow solid. The ¹H NMR spectrum for this product agrees with the chemical shift assignments in previously reported spectra, ²⁵ but the multiplets obscure the signals corresponding to the expected minor byproduct. Crystals were obtained through vapour diffusion of pentane into a concentrated CH₂Cl₂ solution. ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.85–3.82 (m, 4H, NCH₂), 1.74–1.72 (m, 6H, CH₂). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 191.68 (s, quaternary), 50.63 (s, minor, NCH₂), 50.32 (s, major, NCH₂), 25.14 (s, major, 3,5-CH₂), 24.93 (s, minor, 3,5-CH₂), 23.41(s, 4-CH₂).

"[AuCl₂(p-tolyl₂dtc)]" mixture. Li(p-tolyl₂dtc) (185 mg, 0.66 mmol, 1.0 eq.) was dissolved in water (12 mL) and added dropwise to an aqueous solution (8 mL) of K[AuCl₄] (250 mg, 0.66 mmol, 1.0 eq.) whilst vigorously stirring. A brown precipitate formed and the suspension was further stirred for 1 h. The solid was isolated by filtration, washed with water $(3 \times 5 \text{ mL})$ and Et₂O $(2 \times 5 \text{ mL})$ and dried in vacuo. The solid was then redissolved in a minimum amount of CH2Cl2 and purified by flash chromatography through a silica plug (eluent: CH₂Cl₂) to remove the brown contaminant. Yield: 177 mg of an orange solid. X-ray quality crystals were obtained through vapour diffusion of pentane into a concentrated CH2Cl2 solution. ¹H NMR (400.1 MHz, DMSO-d₆): δ 7.64 (m, 4H, $C_{\Delta r}$ -H, minor product), 7.62 (m, 8H, CAr-H, major product), 7.45-7.43 (m, 4H, C_{Ar}-H, minor product), 7.42-7.40 (m, 8H, C_{Ar}-H, major product), 2.37 (s, 6H, CH₃, minor product), 2.34 (s, 12H, CH₃, major product). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 141.44 (s, quaternary, major), 140.88 (s, quaternary, minor), 136.43 (s, quaternary, minor), 135.42 (s, quaternary, major), 131.22 (s, CH, major), 131.01 (s, CH, minor), 126.57 (s, CH, minor), 126.32 (s, CH, major), 20.83 (s, CH₃, major), 20.78 (s, CH_3 , minor).

"[AuBr₂(Me₂dtc)]" mixture. Na(Me₂dtc)·2(H₂O) (66 mg, 0.46 mmol, 1.0 eq.) was dissolved in water (3 mL) and added dropwise to an aqueous solution (3 mL) of AuBr₃ (200 mg, 0.46 mmol, 1.0 eq.) whilst vigorously stirring. An orange precipitate formed and the suspension was further stirred for 1 h. The solid was isolated by filtration, washed with water (5 × 1 mL) and Et₂O (5 × 1 mL) and dried *in vacuo*. Yield: 128 mg of an orange solid. ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.38 (s, 6H, major product), 3.35 (s, 6H, minor product). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 194.27 (s, quaternary, major), 188.69 (s, quaternary, minor), 40.73 (s, CH₃, major), 40.09 (s, CH₃, minor).

"[AuBr₂(Et₂dtc)]" mixture. [Au(Et₂dtc)]_n (172 mg, 0.5 mmol, 1.0 eq.) was suspended in CH₂Cl₂ (25 mL). A solution of Br₂ (80 mg, 0.5 mmol, 1.0 eq.) in CH₂Cl₂ (10 mL) was added dropwise over 15 minutes with vigorous stirring. The solid dissolved, resulting in a dark orange solution. X-ray quality crystals were obtained by allowing this solution to slowly evaporate to dryness. Yield: 217 mg of orange crystals. ¹H NMR (400.1 MHz, CDCl₃): δ 3.79 (q, ${}^{3}J_{\rm HH}$ = 7.3 Hz, 8H, CH₂, major

product), 3.70 (q, ${}^{3}J_{\text{HH}}$ = 7.3 Hz, 4H, CH₂, minor product), 1.44 (overlapping t, ${}^{3}J_{\text{HH}}$ = 7.3 Hz, 12H, CH₃, major product), 1.41 (overlapping t, ${}^{3}J_{\text{HH}}$ = 7.3 Hz, 6H, minor product). ${}^{13}\text{C}\{^{1}\text{H}\}$ NMR (100.6 MHz, CDCl₃): δ 46.86 (s, CH₂, major product), 45.63 (s, CH₂, minor product), 12.61 (s, CH₃, major product), 12.41 (s, CH₃, minor product).

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Note: "[AuBr₂(Et₂dtc)]" was also synthesized using the general procedure to provide an authentic sample for comparison of NMR spectra: Na(Et₂dtc)·3(H₂O) (119 mg, 0.67 mmol, 1.0 eq.) was dissolved in water (3 mL) and added dropwise to a vigorously stirred solution of AuBr₃ (291 mg, 0.67 mmol, 1.0 eq.) in water (3 mL) over 15 minutes. The suspension was then stirred for 1 h. An orange precipitate was isolated by vacuum filtration, washed with water (3 × 5 mL) and Et₂O (2 × 5 mL) then dried *in vacuo*. Yield: 230 mg of an orange solid.

"[AuBr₂(p-tolyl₂dtc)]" mixture. Li(p-tolyl₂dtc) (185 mg, 0.66 mmol, 1.0 eq.) was dissolved in water (12 mL) and added dropwise to an aqueous solution (8 mL) of AuBr₃ (291 mg, 0.66 mmol, 1.0 eq.) whilst vigorously stirring. A brown precipitate formed and the suspension was further stirred for 1 h. The solid was isolated by filtration, washed with water (3 \times 5 mL) and Et₂O (2 × 5 mL) and dried in vacuo. The solid was dissolved in CH2Cl2 and filtered through a plug of silica to remove the brown contaminant. Yield: 295 mg of an orange solid. X-ray quality crystals were obtained through slow evaporation of a concentrated CH₂Cl₂ solution. ¹H NMR (400.1 MHz, CDCl₃): δ 7.42 (d, ${}^3I_{HH}$ = 8.4 Hz, 4H, C_{Ar}-H, minor product), 7.34 (d, ${}^{3}J_{HH}$ = 8.4 Hz, 8H, C_{Ar}-H, major product), 7.31-7.27 (overlapping m, 12H, CAr-H, major and minor product), 2.41 (s, 6H, CH₃, major product), 2.37 (s, 12H, CH₃, minor product). ${}^{13}C{}^{1}H$ NMR (100.6 MHz, CDCl₃): δ 141.66 (s, quaternary, major), 141.04 (s, quaternary, minor), 136.37 (s, quaternary, minor), 135.17 (s, quaternary, major), 131.35 (s, CH, major), 131.11 (s, CH, minor), 126.40 (s, CH, minor), 125.99 (s, CH, major), 21.34 (s, CH₃, major), 21.32 (s, CH₃, minor).

The acetone synthesis of $[AuX_2(dtc)]/[Au(dtc)_2]^+$ mixtures (X = Cl, Br) and the isolation of pure $[AuX_2(dtc)]$

General procedure. "Au X_3 " (1.0 eq.) was dissolved in acetone to give a clear yellow solution. A solution of M(dtc) (1.0 eq.) in acetone was added dropwise to the "Au X_3 " solution over a period of 2 hours. The resulting slightly cloudy yellow solution was refluxed for 24 hours, then cooled to RT and filtered, followed by removal of all volatiles to yield an orange solid.

Orange crystals of analytically pure $[AuX_2(dtc)]$ were obtained by redissolving the crude orange solid in a minimum volume of acetone, then allowing the solvent to slowly evaporate. Before complete solvent loss occurred, the remaining liquid was decanted away from the orange crystals. These were then washed with water and Et_2O , and dried *in vacuo*. **Note:** the vessel used for slow evaporation needs to be clean and free from excessive nucleation points (e.g. scratches) otherwise yellow crystals of $[Au(dtc)_2]^+$ co-crystallize with the desired orange solid.

The analytical data below are based on the orange crystalline material obtained by slow evaporation of an acetone solution. Unless stated otherwise, the percentage yields given are those of pure orange crystalline material based off a crystallization run starting from 50 mg of crude orange solid.

[AuCl₂(Me₂dtc)]. Following the general procedure, K[AuCl₄] (173 mg, 0.46 mmol) was dissolved in acetone (125 mL) and Na(Me₂dtc)·2H₂O (66 mg, 0.46 mmol) was dissolved in acetone (125 mL). Yield: 114 mg of crude solid. 24 mg of the crude product was redissolved in acetone and recrystallized as described above, affording 8 mg of orange crystals (35%).

¹H NMR (400.1 MHz, DMSO-d₆): δ 3.38 (s, 6H, CH₃).

¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 40.02 (s, CH₃).

[AuCl₂(Et₂dtc)]. Following the general procedure, K[AuCl₄] (251 mg, 0.66 mmol) was dissolved in acetone (45 mL) and Na(Et₂dtc)·3H₂O (149 mg, 0.66 mmol) was dissolved in acetone (55 mL). Yield: 151 mg of crude solid. The entire crude product was redissolved in acetone and recrystallized as described above, affording 84 mg of orange crystals (55%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.78 (q, ${}^{3}J_{\text{HH}}$ = 7.2 Hz, 4H, CH₂), 1.27 (t, ${}^{3}J_{\text{HH}}$ = 7.2 Hz, 6H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 186.96 (s, quaternary), 46.52 (s, CH₂), 11.99 (s, CH₃).

[AuCl₂(ⁱPr₂dtc)]. Following the general procedure, K[AuCl₄] (200 mg, 0.53 mmol) was dissolved in acetone (45 mL) and K(ⁱPr₂dtc) (114 mg, 0.53 mmol) was dissolved in acetone (55 mL). Yield: 228 mg of crude solid. 50 mg was recrystallized from acetone, affording 25 mg of orange crystals (50%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 4.55 (septet, J = 6.7 Hz, 2H, CH), 1.44 (d, J = 6.7 Hz, 12H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 185.20 (s, quaternary), 54.57 (s, CH), 19.12 (s, CH₃).

[AuCl₂(ⁱBu₂dtc)]. Following the general procedure, K[AuCl₄] (200 mg, 0.53 mmol) was dissolved in acetone (45 mL) and K(ⁱBu₂dtc) (120 mg, 0.53 mmol) was dissolved in acetone (55 mL). Yield: 226 mg of crude solid. 50 mg was recrystallized from acetone, affording 29 mg of orange crystals (57%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.60 (d, J = 7.6 Hz, 4H, CH₂), 2.30–2.20 (m, 2H, CH), 0.92 (d, J = 6.6 Hz, 12H, CH₃). ¹³C(¹H) NMR (100.6 MHz, DMSO-d₆): δ 189.51 (s, quaternary), 58.00 (s, CH₂), 26.35 (s, CH), 19.44 (s, CH₃).

[AuCl₂(pyrr-dtc)]. Following the general procedure, K[AuCl₄] (200 mg, 0.53 mmol) was dissolved in acetone (45 mL) and Na (pyrr-dtc) (90 mg, 0.53 mmol) was dissolved in acetone (55 mL). Yield: 166 mg of crude solid. 23 mg was recrystallized from acetone, affording 9 mg of orange crystals (39%). 1 H NMR (400.1 MHz, DMSO-d₆): δ 3.83–3.80 (m, 4H, NCH₂), 2.07–2.03 (m, 4H, CH₂). 13 C{ 1 H} NMR (100.6 MHz, DMSO-d₆): δ 51.07 (s, NCH₂), 23.36 (s, CH₂). No peak was observed for the quaternary carbon centre.

[AuCl₂(pip-dtc)]. Following the general procedure, K[AuCl₄] (200 mg, 0.53 mmol) was dissolved in acetone (45 mL) and Na (pip-dtc) (97 mg, 0.53 mmol) was dissolved in acetone (55 mL). Yield: 204 mg of crude solid. 53 mg was recrystallized from acetone, affording 47 mg of orange crystals (89%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.83–3.80 (m, 4H, NCH₂), 1.72–1.70 (m, 6H, 3,4,5-CH₂). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 184.66 (s, quaternary) 50.66 (s, NCH₂), 24.95 (s, NCH₂CH₂), 23.47 (s, CH₂CH₂CH₂).

[AuCl₂(*p*-tolyl₂dtc)]. Following the general procedure, K [AuCl₄] (200 mg, 0.53 mmol) was dissolved in acetone (45 mL) and Li(*p*-tolyl₂dtc) (148 mg, 0.53 mmol) was dissolved in acetone (55 mL). The crude product from this synthesis was contaminated with a brown residue, which was removed by dissolving the crude product in CH₂Cl₂ and passing through a silica plug before removing all solvents. Yield: 162 mg of crude solid. 50 mg was recrystallized from acetone, affording 29 mg of orange crystals (59%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 7.63 (m, 4H, C_{Ar}-H), 7.45–7.43 (m, 4H, C_{Ar}-H), 2.37 (s, 6H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 196.54 (s, quaternary), 141.42 (s, quaternary), 135.40 (s, quaternary), 131.21 (s, CH), 126.31 (s, CH), 20.82 (s, CH₃).

[AuBr₂(Me₂dtc)]. Following the general procedure, AuBr₃ (217 mg, 0.50 mmol) was dissolved in acetone (125 mL) and Na(Me₂dtc)·2H₂O (71 mg, 0.50 mmol) was dissolved in acetone (125 mL). Yield: 62 mg of crude solid. 20 mg of the crude product was redissolved in acetone and recrystallized as described above, affording 12 mg of orange crystals (60%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.35 (s, 6H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 39.59 (s, CH₃).

[AuBr₂(Et₂dtc)]. Following the general procedure, AuBr₃ (158 mg, 0.36 mmol) was dissolved in acetone (45 mL) and Na (Et₂dtc)·3H₂O (82 mg, 0.36 mmol) was dissolved in acetone (55 mL). Yield: 166 mg of crude product. 50 mg was recrystallized from acetone, affording 31 mg of orange crystals (62%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.75 (q, ³ $J_{\rm HH}$ = 7.2 Hz, 4H, CH₂), 1.27 (t, ³ $J_{\rm HH}$ = 7.2 Hz, 6H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 188.38 (s, quaternary), 46.04 (s, CH₂), 11.98 (s, CH₃).

[AuBr₂(p-tolyl₂dtc)]. Following the general procedure, AuBr₃ (150 mg, 0.34 mmol) was dissolved in acetone (45 mL) and Li (p-tolyl₂dtc) (96 mg, 0.34 mmol) was dissolved in acetone (55 mL). The crude product from this synthesis was contaminated with a brown residue, which was removed by dissolving the crude product in CH₂Cl₂ and passing through a silica plug before removing all solvents. Yield: 124 mg of crude solid. 53 mg was recrystallized from acetone, affording 36 mg of orange crystals (68%). ¹H NMR (400.1 MHz, DMSO-d₆): δ 7.64 (m, 4H, C_{Ar}–H), 7.45–7.43 (m, 4H, C_{Ar}–H), 2.36 (s, 6H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 197.87 (s, quaternary), 141.39 (s, quaternary), 135.17 (s, quaternary), 131.21 (s, CH), 126.43 (s, CH), 20.83 (s, CH₃).

[Au(Et₂dtc)₂][BF₄]. K[AuCl₄] (250 mg, 0.66 mmol, 1.0 eq.) was dissolved in acetone (100 mL). Na[BF₄] (72.6 mg, 0.66 mmol, 1.0 eq.) was dissolved in acetone (20 mL) and added dropwise to the K[AuCl₄] solution. A solution of Na(Et₂dtc)·3H₂O (298 mg, 1.32 mmol, 2.0 eq.) in acetone (120 mL) was then added dropwise over a period of 2 hours. After complete addition, the slightly cloudy yellow mixture was allowed to stir at RT for 4 hours, before the mixture was filtered to remove the white precipitate. The filtrate was concentrated under reduced pressure to a volume of ~20 mL, followed by addition of 120 mL of hexane to the solution causing precipitation of an orange solid. The supernatant liquid was removed, and the product was dried under vacuum for 2 days. Yield: 248 mg

(64.6%) of an orange solid. Crystals were obtained by slow evaporation of a concentrated acetone solution. ¹H NMR (400.1 MHz, (CD₃)₂CO): δ 3.94 (q, ³ $J_{\rm HH}$ = 7.2 Hz, 8H, CH₂), 1.44 (t, ³ $J_{\rm HH}$ = 7.2 Hz, 12H, CH₃). ¹³C{¹H} NMR (100.6 MHz, (CD₃)₂CO): δ 195.53 (s, quaternary), 47.83 (s, CH₂), 12.48 (s, CH₃). ¹H NMR (400.1 MHz, DMSO-d₆): δ 3.77 (q, ³ $J_{\rm HH}$ = 7.2 Hz, 8H, CH₂), 1.30 (t, ³ $J_{\rm HH}$ = 7.2 Hz, 12H, CH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-d₆): δ 193.88 (s, quaternary), 46.56 (s, CH₂), 12.10 (s, CH₃). Elemental analysis: calc. for C₁₀H₂₀AuBF₄N₂S₄ (580.27): C 20.70%, H 3.47%, N 4.83%. Found: C 20.79%, H 2.99%, N 4.58%.

Data availability

The data supporting this article have been included as part of the ESI.†

 $\label{eq:continuous} \begin{array}{llll} & \text{Crystallographic data: CCDC reference numbers} & 2374277 \\ [\text{AuCl}_2(^{\mathrm{i}}\text{Pr}_2\text{dtc})], & 2374278 & [\text{AuCl}_2(^{\mathrm{i}}\text{Bu}_2\text{dtc})], & 2374279 \\ [\text{Au}(^{\mathrm{i}}\text{Bu}_2\text{dtc})_2][\text{AuCl}_2], & 2374280 & [\text{AuCl}_2(p\text{-tolyl}_2\text{dtc})], & 2374281 \\ [\text{Au}(p\text{-tolyl}_2\text{dtc})][\text{AuCl}_2], & 2374282 & [\text{AuBr}_2(p\text{-tolyl}_2\text{dtc})], & 2374283 \\ [\text{Au}(p\text{-tolyl}_2\text{dtc})][\text{AuBr}_2], & 2374284 & [\text{AuBr}_2(\text{Et}_2\text{dtc})], & 2374285 \\ [\text{Au}(\text{Et}_2\text{dtc})_2][\text{AuBr}_2], & 2374286 & [\text{Au}(\text{Et}_2\text{dtc})_2][\text{BF}_4], & 2374287 \\ [\text{AuBr}_2(\text{Me}_2\text{dtc})], & 2374288 & [\text{Au}(\text{Me}_2\text{dtc})_2][\text{AuBr}_2] & \text{contain crystallographic data in CIF format.} \\ \end{array}$

Conflicts of interest

There are no conflicts to declare.

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