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Subtle Steric Differences Impact the Structural and Conducting Properties of Radical Gold Bis(dithiolene) Complexes

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Abstract

Among single component molecular conductors, neutral radical gold dithiolene complexes $[(R\text{-thiazdt})_2\text{Au}]^{\bullet}$ derived from the *N*-alkyl-1,3-thiazoline-2-thione-4,5-dithiolate (R-thiazdt) ligand provide an extensive series of conducting, non-dimerized, $\frac{1}{2}$ -filled band systems. Analogs of the known R = isopropyl derivative were investigated here with R = NMe₂, cyclopropyl (cPr) and n-propyl (nPr), aiming at rationalizing the different solid state structures adopted by these compounds despite very closely related substituents on the heterocyclic nitrogen atom. An original crisscross organization within dimerized chains is observed with R = NMe₂, differing however from the analogous iPr derivative by a 180° rotation of the heterocyclic nitrogen substituent. On the other hand, the cyclopropyl and n-propyl substituents lead to robust, uniform, non-dimerized chains with a strongly 1D electronic structure and a formal $\frac{1}{2}$ -filled electronic structure. The semiconducting behaviour of these two radical complexes is characteristic of a Mott insulator, whose sensitivity to external pressure has been evaluated up to 2.5 GPa.

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Single component molecular conductors have been defined by opposition to classical charge transfer salts as conducting materials involving only one single molecular entity. Two main classes of compounds respond today to this definition: (i) neutral complexes of electroactive tetrathiafulvalene dithiolene ligands successfully isolated as metallic compounds by Kobayashi *et al.*^[1] and (ii), fully organic neutral radical species such as thiazyl radicals (dithiadiazolyl and dithiazolyl),^[2,3,4] spiro-bis(phenalenyl)-boron^[5] and tetrathiafulvalene (TTF) based radicals.^[6,7] Neutral dithiolene complexes lacking the tetrathiafulvalene backbone were also reported for their metallic conductivity, such as trimetallic molybdenum dithiolene cluster complexes,^[8] or more recently [Ni(dmit)₂] under pressure.^[9] Neutral gold dithiolene complexes offer another alternative.^[10] They are indeed obtained as radical species from the 1e⁻ oxidation of the d⁸ anionic Au^{III} [Au(dithiolene)₂]⁻ complexes. Strong intermolecular interactions in the solid state between such delocalized radical species are however not a sufficient condition to observe a sizeable or even metallic conductivity because these radical species most often dimerize to form a 2e⁻ -eventually delocalized- bond.^[10] Even the regular stacking of planar π -type radicals leading to a 1/2-filled conduction band is threatened by the so-called Peierls transition which opens a gap at the Fermi level through a dimerization of the stacks.

We recently prepared a series of neutral radical gold dithiolene complexes, [Au(R-thiazdt)₂][•] (Chart 1a) based on the *N*-ethyl-1,3-thiazoline-2-thione-4,5-dithiolate (Et-thiazdt) ligand with one radical per site,^[11] which are not affected by the usual propensity of radical species to dimerize in the solid state, and which behave as Mott insulators. Several modifications of this prototypical [Au(Et-thiazdt)₂][•] complex have been reported, aiming at a deeper understanding of the structure-properties relationship within this original single-component-conductor family. For instance, the use of diselenolene ligands, rather than dithiolene ones (Chart 1b), enhances the orbital overlaps and increases the room temperature conductivity by two orders of magnitude.^[12] Contrariwise, the replacement of the exocyclic sulfur atom by an oxygen one leads to decreased intra-layer interactions and smaller conductivities (Chart 1b). Such structural modifications do not entail substantial changes in the solid state packing but essentially the strength of the intermolecular interactions through so-called anisotropic chemical pressure effects.^[12] The nature of the substituent R on the nitrogen atom in these *N*-substituted 1,3-thiazoline-2-thione-4,5-dithiolate ligands provides another tool to modify these series in a deeper manner. For instance, introduction of hydroxyethyl substituents revealed solid state structures

with added OH...S hydrogen bonds (Chart 1c).^[13] Also, the higher steric constraints brought by bulkier isopropyl groups (Chart 1c) led to an original crisscross overlap of the radical species within uniform stacks, with a very specific electronic structure, highly resistant to physical pressure effects.^[14] It confirms that steric constraints considerably affect the organization of the molecules in the solid state and also the associated physical properties which result from strong intermolecular interactions such as the conducting one.

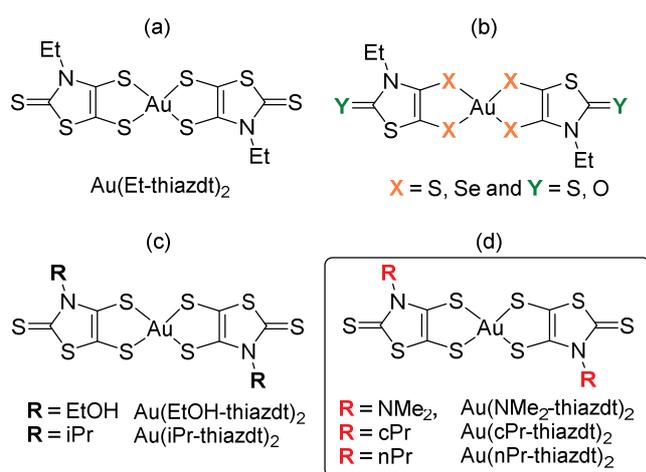


Chart 1

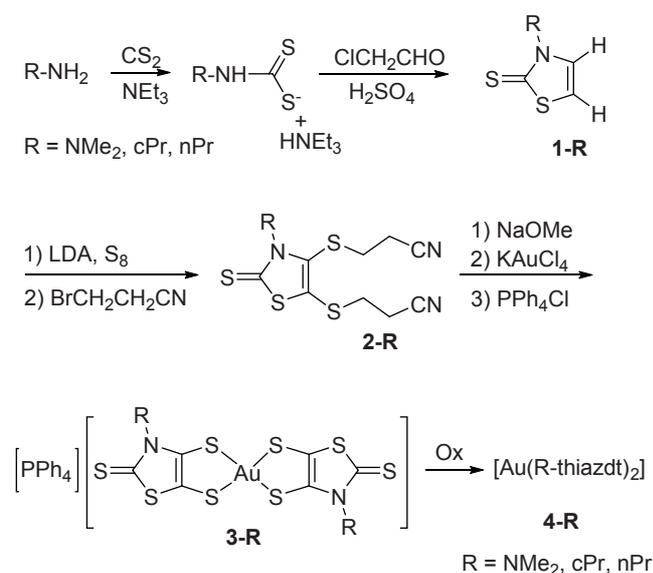
In this paper, we want to unravel the striking evolutions of structural and electronic properties we observed within an extended series of single component conductors based to these N-isopropyl gold dithiolene complexes, by replacing the isopropyl substituent by either a dimethylamino (NMe_2), a cyclopropyl (cPr) or a n-propyl substituent (nPr). While the dimethylamino group is expected to exhibit steric constraints similar to that of the isopropyl one, the somewhat less bulky but less flexible cyclopropyl group provides another variation, which should also contrast with the linear n-propyl derivative, as highlighted below.

Results and discussion

Syntheses

An efficient strategy for the preparation of 1,3-thiazoline-2-thiones consists in the use of primary amine as starting compound.^[15] This chemical route allows the formation of various N-substituted heterocycles depending on the primary amine used in the first step,^[16] as here N,N-dimethylhydrazine, cyclopropylamine and n-

propylamine. The preparation of the anionic gold dithiolene complexes **3-NMe₂**, **3-cPr** and **3-nPr** follows a described procedure, outlined in Scheme 1.^[11] As a first step, the N-alkyl-1,3-thiazoline-2-thione **1** was formed by reacting the dithiocarbamate salt, obtained by adding carbon disulfide and triethylamine to a solution of the primary amine, with chloroacetaldehyde followed by cyclisation and dehydration in the presence of sulfuric acid. Bis-metallation of **1-R** (R = NMe₂, cPr, nPr) with LDA followed by the addition of S₈ and bromopropionitrile led to the formation of the protected dithiolate ligands **2-R** (Scheme 1).



Scheme 1. Synthetic route toward the neutral radical gold complexes **4-R** (R = NMe₂, cPr, nPr)

The dithiolene ligands were generated by adding sodium methanolate to **2-R** and subsequent addition of KAuCl₄ and PPh₄Cl to the medium led to monoanionic Au^{III} complexes [PPh₄][Au(R-thiazdt)₂] **3-R** (R = NMe₂, cPr, nPr) which were recrystallized in CH₃CN. According to this procedure, crystals amenable for X-ray diffraction were obtained for two monoanionic complexes **3-NMe₂** and **3-nPr** as tetraphenylphosphonium salts. The monoanionic complex **3-cPr** was also prepared with the Et₄N⁺ counter ion, [NEt₄][Au(cPr-thiazdt)₂], allowing us to obtain crystals suitable for X-ray diffraction analyses. The description of the molecular structures of these three monoanionic species is detailed below. Due to the dissymmetrical character of the dithiolate ligand, these monoanionic complexes **3-R** are susceptible to exist in two configurations, the *trans* and *cis* ones. ¹H NMR spectra exhibit at room

temperature one set of signals indicating a fast exchange in solution between the *cis* and *trans* isomers. In order to generate the neutral radical complexes, we focused on the electrocrystallization approach as it generally allows the formation of good quality crystals at the anode.^[17] Upon application of a constant current intensity of 0.3-0.5 μA to a solution of $[\text{PPh}_4][\text{Au}(\text{R-thiazdt})_2]$ **3-R** (R = NMe₂, cPr, nPr) in CH₃CN containing NBu₄PF₆ as supporting electrolyte, crystals of the three neutral species **4-R** were obtained on the electrode.

Electrochemical properties

The redox properties of the three monoanionic complexes $[\text{PPh}_4][\text{Au}(\text{R-thiazdt})_2]$ **3-R** were investigated by cyclic voltammetry carried out in CH₂Cl₂ using NBu₄PF₆ as supporting electrolyte. The redox potentials are gathered in Table 1 together with those of analogous complexes for comparison purposes and a representative CV is given in Figure 1 for **3-nPr**. For all these complexes, the shapes of the CV are very similar. They exhibit two reversible oxidation waves assigned to the successive formation of the neutral and the monocationic complex and one irreversible reduction wave corresponding to the reduction to the dianionic species (Figure 1). Whatever the nature of R, the redox potentials are very close to those found for analogous complexes with ethyl and isopropyl substituents $[\text{Au}(\text{R-thiazdt})_2]$ R = Et, iPr. Note the sharp potential responses on the reduction processes from cation-to-neutral and neutral-to-anion that reveal adsorption phenomena for all these complexes.

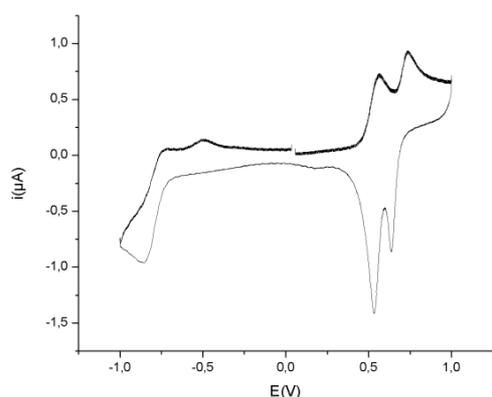


Figure 1 Cyclic voltammogram of $[\text{Au}(\text{nPr-thiazdt})_2]^{1-}$ **3-nPr** in CH₂Cl₂ with NBu₄PF₆ 0.1 M as electrolyte, E in V vs SCE, Pt electrode, scan rate 100 $\text{mV}\cdot\text{s}^{-1}$.

Table 1. Redox potentials of $[\text{Au}(\text{R-thiazdt})_2]^{-1}$ **3-R** complexes (E in V vs. SCE) in CH_2Cl_2 , NBu_4PF_6 , Pt, scan rate $100 \text{ mV}\cdot\text{s}^{-1}$.

R	E_{red}^*	$E_{\text{pa}1}/E_{\text{pc}1}^{-}$ 1/0	$E_{\text{pa}2}/E_{\text{pc}2}^{0/+1}$	Ref.
NMe ₂	-0.96	0.51/0.43**	0.66/0.57	this work
cPr	-0.95	0.52/0.46**	0.65/-	this work
nPr	-0.85	0.56/0.53**	0.73/0.64	this work
iPr	-0.95	0.56/0.50**	0.69/0.59	14
Et	-0.90	0.55/0.49	0.70/0.61	11

* Irreversible process. ** Adsorption

Molecular structures

The monoanionic gold dithiolene complex **3-NMe₂** crystallizes as Ph_4P^+ salt in the monoclinic system, space group $\text{P}2_1/\text{n}$. Two types of crystals were obtained for the monoanionic **3-nPr** complex, the n-propyl derivative was isolated either as acetonitrile solvate, formulated as $(\text{Ph}_4\text{P})(\mathbf{3-nPr})\cdot(\text{CH}_3\text{CN})_2$ or as the pure compound $(\text{Ph}_4\text{P})(\mathbf{3-nPr})$, both crystallizing in the monoclinic system, space group $\text{C}2/\text{c}$ while the Et_4N^+ salt of **3-cP** crystallizes in the triclinic system, space group $\text{P}-1$. The molecular structures of the complexes are presented in Figure 2. All complexes, with such dissymmetrically substituted ligands, adopt a *trans* configuration with a square planar geometry around the gold atom. The only exception is $(\text{Ph}_4\text{P})(\mathbf{3-nPr})$ which adopts also a *cis* configuration (Figure 2d). All the metallacycles are slightly distorted along the $\text{S}\cdots\text{S}$ axis with angles from 2 to 5° (Table 2). Selected bond lengths are collected in Table 2 and are similar in the four structures.

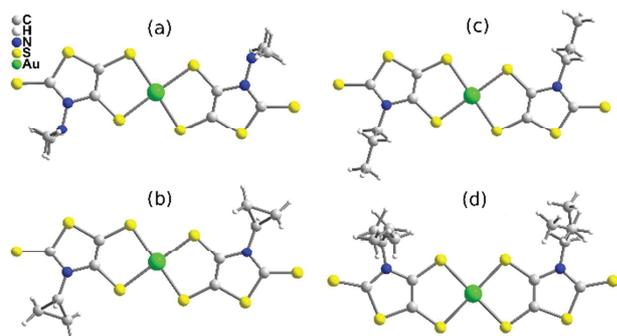
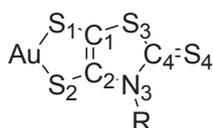


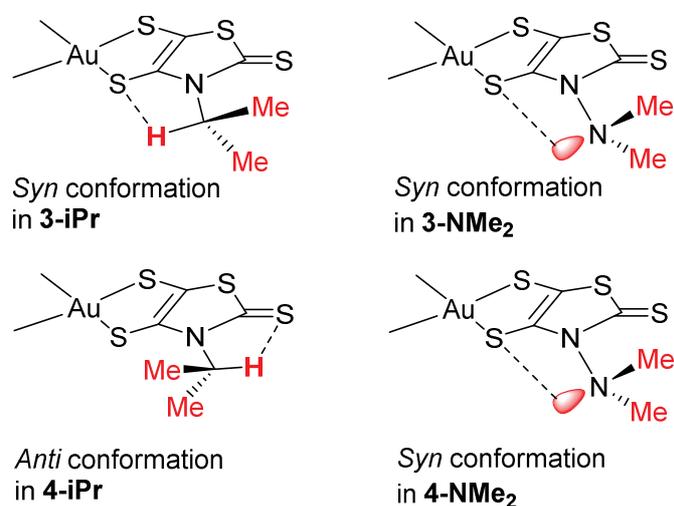
Figure 2. Detail of the monoanionic complexes **3-R** showing the orientation of the nitrogen substituents. (a) R = NMe₂; (b) R = cyclopropyl (cPr); (c) R = nPr in the solvate (Ph₄P)(**3-nPr**)•(CH₃CN)₂; (d) R = nPr in (Ph₄P)(**3-nPr**).

Table 2. Selected bond lengths (Å) torsion angle $\theta_{S_1...S_2}$ (°) within monoanionic and neutral gold dithiolene complexes *trans*-**3-R** and **4-R** (R = NMe₂, cPr, nPr).



	3-NMe₂	4-NMe₂	3-cPr	4-cPr	3-nPr	4-nPr
Charge	monoanion	neutral	monoanion	neutral	monoanion	neutral
Au–S1	2.3310(9) 2.3355(10)	2.3218(10)	2.3242(11)	2.3167(8)	2.3189(9)	2.3182(14)
Au–S2	2.3263(11) 2.3201(10)	2.3159(11)	2.3208(16)	2.3206(36)	2.3239(7)	2.3206(16)
S1–C1	1.740(4) 1.739(4)	1.714(5)	1.732(6)	1.714(14)	1.739(3)	1.710(6)
S2–C2	1.749(4) 1.738(4)	1.714(3)	1.752(4)	1.725(14)	1.747(4)	1.735(6)
C1–C2	1.330(5) 1.344(5)	1.365(6)	1.347(8)	1.344(19)	1.331(5)	1.369(9)
C1–S3	1.746(4) 1.758(4)	1.748(4)	1.748(4)	1.750(14)	1.748(4)	1.737(6)
C2–N3	1.410(5) 1.404(5)	1.384(6)	1.397(7)	1.414(17)	1.408(4)	1.388(8)
N3–C4	1.352(5) 1.358(5)	1.374(6)	1.366(5)	1.361(17)	1.357(5)	1.366(8)
S3–C4	1.747(4) 1.760(4)	1.752(6)	1.742(7)	1.725(14)	1.733(4)	1.748(7)
C4–S4	1.672(4) 1.656(4)	1.648(5)	1.659(5)	1.671(14)	1.667(4)	1.646(6)
$\theta_{S_1...S_2}$	4.81(8)	4.96(16)	2.63(2)	1.06(30)	3.42(7)	2.14(12)

Despite having very similar size, the orientation of the R substituent of the heterocyclic nitrogen atom with regard to the thiazoline ring strongly differs from one complex to the other. The *syn* disposition of the methyl groups in **3-NMe₂** (Scheme 2), with one methyl above and one methyl below the plane of the thiazoline-2-thione core and the lone pair oriented towards the metallacycle, recalls the *syn* conformation observed for the methyl groups in the isopropyl (iPr) analogous complex **3-iPr**.^[14] Besides, the cyclopropyl groups in **3-cPr** are located above and below the thiazoline plane (Figure 2b) and the n-propyl substituents are strongly disordered in the *cis* **3-nPr** complex (Figure 2d).



Scheme 2.

All the neutral complexes **4-R** (R = NMe₂, cPr, nPr) crystallize as the *trans*-isomers and they all exhibit a planarization of the metallacycles upon oxidation leading to quasi planar skeletons. Geometrical modifications between the monoanionic and neutral species are those generally observed in gold bis(dithiolene) complexes such as, within the metallacycles, a shortening of the C–S bonds and an elongation of the C=C bond (Table 2). This phenomenon is due to the non-innocent character of the dithiolene ligand. Within the thiazoline-2-thione ring, the modification of the bond lengths is less pronounced. Focusing now on the alkyl group, we note that its orientation with respect to the thiazoline ring is the same as the one observed for the monoanionic species for all the three complexes. For instance, the *syn* configuration of the NMe₂ (Scheme 2) maintains the orientation of the lone pair

towards the metallacycle, a possible consequence of a stabilizing interaction between the lone pair of the nitrogen atom and the closest sulfur atom of the metallacycle. This orientation contrasts with the *anti* conformation observed for the methyl groups in the isopropyl (iPr) analogous complex **4-iPr** (Scheme 2).^[14]

Structural and electronic properties of the radical complexes

Transport measurements were carried out on crystals of the neutral radical species **4-R** along the long axis of the crystals which correspond in all cases to the stacking axis of the complexes **4-R**. Conductivity values obtained at room temperature are collected in Table 3 together with those of some analogous complexes with R = Et, iPr for comparison. For the three novel complexes, the temperature dependence of the resistivity at ambient pressure shows a semiconducting behavior with an activation energy in the range of 0.07–0.18 eV, as deduced from the fit of the data with a law of the type $\rho = \rho_0 \exp(E_a/kT)$ (Figures S2 & S3). Temperature dependence of the magnetic susceptibility was measured for both **4-nPr** and **4-cPr**, revealing a very weak, temperature independent response.

Table 3. Conductivity values at room temperature and ambient pressure for the radical complexes **4-R** together with energy activation (E_a).

4-R	σ_{RT} (S.cm ⁻¹)	E_a (eV)	Ref
R = NMe ₂	0.045	0.18	this work
R = cPr	0.037	0.14	this work
R = nPr	0.36	0.07	this work
R = iPr	5	0.06	14
R = Et	0.33	0.12	11
R = EtOH	0.05-0.07	0.14	13

We will see below that the large differences of conductivity and activation energies are intimately correlated to the details of the solid state arrangement adopted by these complexes, which furthermore strongly differ from what was observed earlier in **4-Et** or **4-iPr**. The highly conducting **4-Et** species adopts indeed a non-dimerized, uniform chain structure with strong inter-chain interactions. It even becomes metallic under pressure.^[11] On the other hand, the larger steric constraints of the iPr groups, which adopt an *anti* conformation (See Scheme 2) led to an

original crisscross stacking of the radicals and, despite a uniform stack, the apparition of a small direct gap in the band structure due to an avoided band crossing.^[14]

If we consider now the NMe₂-substituted complex **4-NMe₂**, it crystallizes in the orthorhombic system, space group I222 (n° 23), with the gold complex on a two-fold axis. The neutral radical molecules do form stacks along the *a* axis, with a crisscross orientation comparable to that in **4-iPr** (Figure 3). However, at variance with **4-iPr** which exhibits an apparently similar structure, the stacks are not uniform any more, as shown for example from the Au•••Au distances at 3.4660(6) Å and 3.5395(6) Å, indicating that the complexes are actually dimerized along the stacking axis.

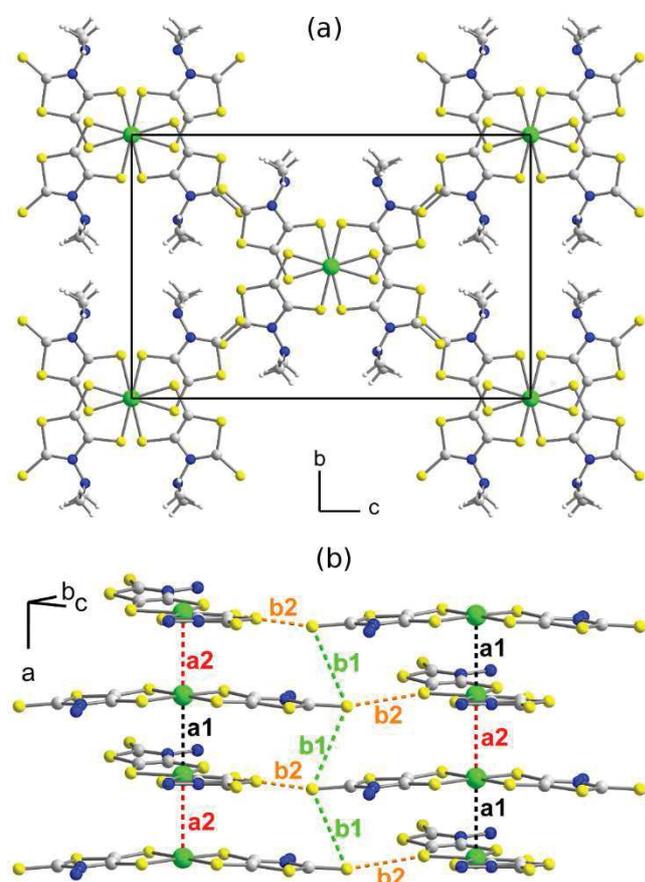


Figure 3. (a) Projection view along *a* of the unit cell of **4-NMe₂**. (b) Details of the overlap interactions a1, a2, b1 and b2. The dotted lines are only a guide to the eyes to visualize interacting molecules.

The calculated overlap energies β associated with these two intra-stack interactions, *a*₁ and *a*₂ (Figure 3b) amount to $\beta_{a_1} = 0.8$ and $\beta_{a_2} = -0.357$ eV

respectively. Two other interactions b_1 and b_2 are found between the stacks, associated with short $S\cdots S$ inter-columnar contacts, with associated overlap energies of $\beta_{b1} = 0.213$ eV and $\beta_{b2} = 0.036$ eV. The consequences of this dimerization on the electronic structure are clearly seen in the calculated band structures of **4-NMe₂** and **4-iPr** shown in Figure 4. With four different molecular orientations in the unit cell, each molecular energy level gives rise to four bands. As a consequence of the stack dimerization in **4-NMe₂**, a large gap is observed within the four SOMO bands marked with an asterisk (*), with the two lower bands filled and the two upper bands empty. On the other hand, in **4-iPr**, an avoided crossing at Fermi level generates a much smaller gap. The direct consequence is a room temperature conductivity in **4-NMe₂** decreased by two orders of magnitude and an activation energy multiplied by 3 (Figure S2). The origin of these striking differences between the two complexes despite the very similar shape and volume of the iPr and NMe₂ substituents can be traced back to the details of their relative orientation. We have already noted the *anti* conformation of the iPr group in **4-iPr** at variance with the *syn* orientation of the NMe₂ group in **4-NMe₂** (Scheme 2). As shown in Figure 5, it brings a small but notable difference in the dihedral angle between complexes along the chains, which then can contribute to the observed differences in the stacking patterns, and the resulting electronic structures.

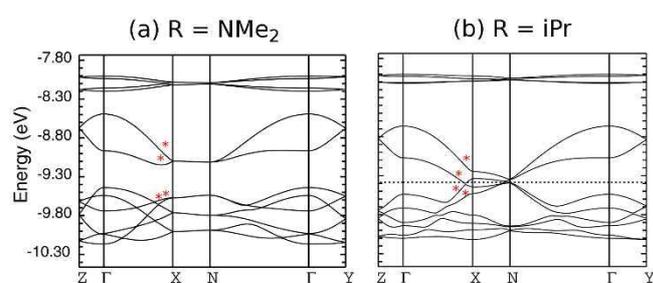


Figure 4 Calculated band structure of the gold complexes a) **4-NMe₂** and b) **4-iPr** with the HOMO-1(4 bands), SOMO (4 bands marked with red asterisks) and LUMO (4 bands) where $\Gamma = (0, 0, 0)$, $X = (\frac{1}{2}, 0, 0)$, $Y = (0, \frac{1}{2}, 0)$, $Z = (0, 0, \frac{1}{2})$, and $N = (\frac{1}{2}, 0, \frac{1}{2})$ of the Brillouin zone of the triclinic lattice. The dashed line in **4-iPr** refers to the Fermi level assuming a metallic filling of the levels.

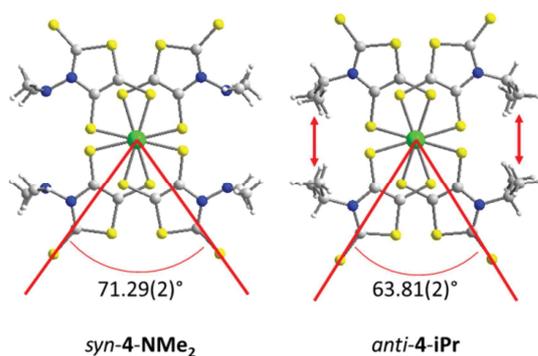


Figure 5. Comparison of overlap patterns with the chains in **4-iPr** and **4-NMe₂**.

The crystal structures of the n-propyl and cyclopropyl derivatives (**4-nPr**, **4-cPr**) are closely related and will be described together in the following. Albeit both compounds are semiconductors, **4-nPr** exhibits a higher conductivity and smaller activation energy than **4-cPr** (Table 2). The analysis and comparison of both structures can provide a rationale for these differences. As shown in Figure 6, **4-nPr** crystallizes in the triclinic system, space group $P\bar{1}$, and **4-cPr** in the monoclinic system, space group $P2_1/c$. In both structures, the radical gold complexes are located on inversion centers, and stack into uniform chains running along a axis. Much weaker inter-stack interactions further develop in both structures along b axis, generating (a,b) conducting layers, isolated from each other along c axis. A detail of the interactions within each layer is given in Figure 7. The associated $\beta_{\text{HOMO-HOMO}}$ overlap interactions amount to $\beta_a = -0.4067$ eV, $\beta_{b1} = -0.0078$ eV and $\beta_{b2} = 0.0383$ eV in **4-nPr**, and to $\beta_a = -0.1975$ eV, $\beta_{b1} = 0.0314$ eV and $\beta_{b2} = 0.0080$ eV in **4-cPr**. The resulting band structure calculations shown in Figure 8 reflect this structural organization. With only one molecular orientation per unit cell, it is composed of one band per molecular energy level. The β_a value twice as large in **4-cPr** than in **4-nPr** gives rise to a much larger band dispersion along the stacking $\Gamma \rightarrow X$ direction in the former complex, which also explains its higher conductivity and smaller activation energy (see above, Figure S3).

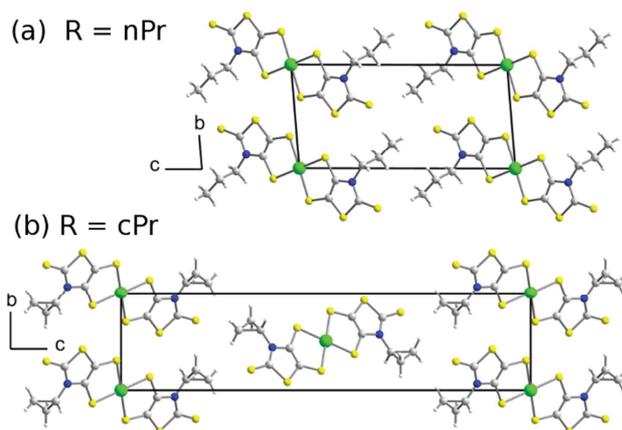


Figure 6. Projection views along a axis of the unit cells of (a) **4-nPr** and (b) **4-cPr**.

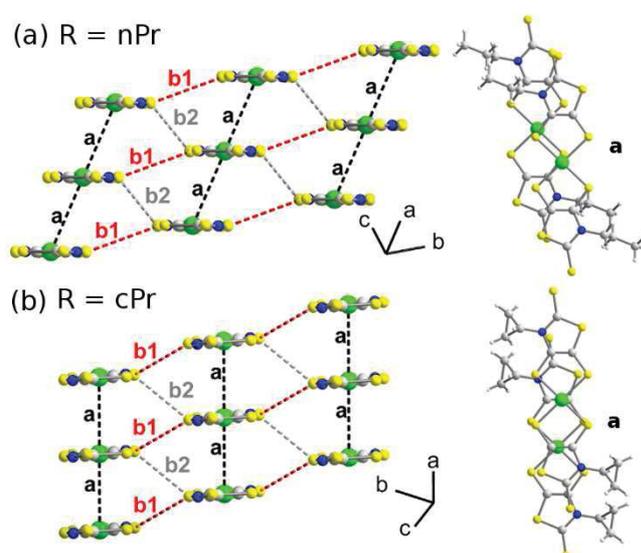


Figure 7. The overlap interactions a , b_1 and b_2 in the layers (left), and detail of the intra-stack overlap a (right) in (a) **4-nPr** and (b) **4-cPr**. The dotted lines are only a guide to the eyes to visualize interacting molecules.

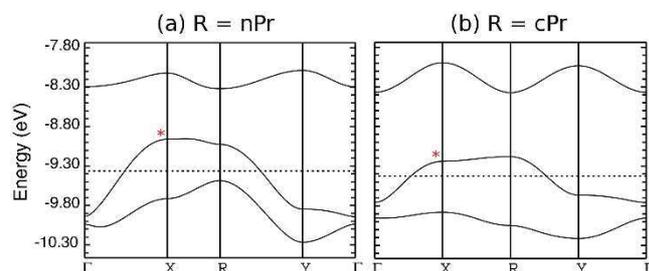


Figure 8. Band structures of the (a) **4-nPr** and (b) **4-cPr** gold complexes. The SOMO bands are marked with a red asterisk. $\Gamma = (0, 0, 0)$, $X = (\frac{1}{2}, 0, 0)$, $Y = (0, \frac{1}{2}, 0)$, and $R = (\frac{1}{2}, -\frac{1}{2}, 0)$ of the Brillouin zone of the monoclinic lattice. The dashed line refers to the Fermi level assuming a metallic filling of the levels.

The stronger overlap interaction within the stacks in **4-cPr**, together with weak inter-stack interactions give rise in both compounds to a strongly one-dimensional band structure, as evidenced from calculated open Fermi surfaces (Figure S4) for both compounds. It follows that the observed semiconducting properties cannot be attributed to an energy gap as described above in **4-NMe₂** or **4-iPr**, but is rather the consequence of strong electron localization. We are faced here with two rare examples of so-called molecular Mott insulators, an electronic ground state where the radical species are neither strongly delocalized in a 1/2-filled conduction band, nor paired two-by-two, eventually in a dimerized chain (Peierls transition). Such an electronic state was already observed in **4-Et** and opens very interesting perspectives. Indeed, this electronic structure is particularly sensitive to external stimuli such as pressure or electric field pulses. In **4-Et**, application of external pressure leads indeed to the stabilization of a metallic state (above 1.3 GPa),^[11,12] while electric field pulses were used to generate a remnant and reversible highly conducting state, of interest toward their use in Resistive Random Access Memories (RRAM).^[18] Along these lines, we therefore investigated the pressure dependence of the most conducting Mott insulator system, i.e. **4-nPr**, up to 2.5 GPa. The RT conductivity increases by two orders of magnitude from 0.36 S cm⁻¹ at ambient pressure, up to 60 S cm⁻¹ at 2.5 GPa, following an exponential law characteristic of semiconductors up to 1.8 GPa (Figure S5). At higher pressure, the increase of the conductivity is slowed down but also perturbed by the freezing of the pressure transmitting medium around 2.1 GPa. Although the conductivity at 2.5 GPa is similar to **4-iPr** value, it is not clear if the conducting regime is reached for **4-nPr** as observed in **4-iPr**. The temperature dependence of the resistivity at different pressures (Figure 9) shows that **4-nPr** keeps its semi-conducting behavior, at least up to 2.5 GPa. The activation energy deduced from the Arrhenius fit of the data in the temperature range 80-150 K decreases continuously from 73 meV at ambient pressure down to 28 meV at 2.5 GPa but does not vanishes (Figure S6 and S7), at variance with the behavior reported for **4-iPr**.^[14] Again the difference between the two complexes finds its origin in their structural organization of the two complexes. The crisscross overlap stacking in **4-iPr** and the numerous inter-stack interactions in the perpendicular plane allowed indeed for a three-dimensional overlap pattern while, as already mentioned, the calculated Fermi surface for **4-nPr** (Figure S4), with

its strong one-dimensional nature does not favor the metallic state to the same extent.

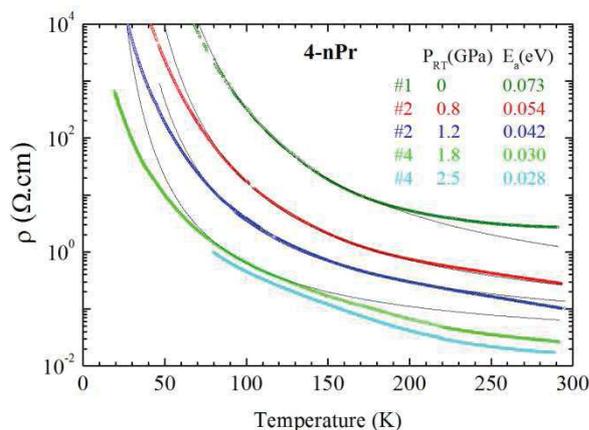


Figure 9. Temperature dependence of the resistivity of **4-nPr** at different pressures. The continuous lines are the Arrhenius fit to the data giving the activation energy.

Conclusion

In the search for novel single-component conductors, we have shown here how the use of sterically demanding, but closely related substituents such as $R = NMe_2$, cyclopropyl and *n*-propyl on the heterocyclic nitrogen atom of the *N*-alkyl-1,3-thiazoline-2-thione-4,5-dithiolate (*R*-thiazdt) ligand give rise, in the corresponding radical gold dithiolene complexes $[(R\text{-thiazdt})_2]^*$, to varied solid state structures. The relative orientation of the methyl groups in the isopropyl and NMe_2 groups vs. the thiazoline heterocycle is the only difference between both complexes but completely modifies their structural and electronic properties, despite an apparently comparable original crisscross stacking. On the other hand, the $R = \text{cyclopropyl}$ and $R = n\text{-propyl}$ radical gold complexes avoid this dimerization and organize into uniform chains with strong one-dimensional character and semiconducting behaviour. The reluctance of these radical complexes toward usual dimerization makes them attractive Mott insulator systems and the evolutions of their conductivity under pressure (up to 2.5 GPa) demonstrate that other stimuli such as electric field pulses could induce a transition to a metallic state. Further investigations along these lines are currently underway, as well as other chemical modifications, with even smaller ($R = \text{Me}$) or larger ($R = \text{tBu}$) substituents.

Experimental section

General. NMR spectra were recorded at room temperature using CDCl_3 unless otherwise noted. Chemical shifts are reported in ppm and ^1H NMR spectra were referenced to residual CHCl_3 (7.26 ppm), ^{13}C NMR spectra were referenced to CHCl_3 (77.2 ppm). Mass spectra were recorded with a Bruker MaXis 4G for **3-nPr** and **4-cPr**, with a Thermo Fisher Q-Exactive for **3-cPr**, **3-NMe₂** and **4-nPr**, and with Waters Q-ToF 2 instrument for all the other derivatives by the Centre Régional de Mesures Physiques de l'Ouest, Rennes. CVs were carried out on a 10^{-3} M solution of complex in CH_2Cl_2 with NBu_4PF_6 0.1 M with a Pt working electrode. Potentials were measured *versus* Saturated Calomel Electrode (SCE). Chemicals and materials from commercial sources were used without further purification.

N-alkyl-1,3-thiazoline-2-thiones (1): To a solution of primary amine (50 mmol, propylamine: 4.1 mL, cyclopropylamine: 3.5 mL, N,N-dimethylhydrazine: 3.8 mL) and triethylamine (7.8 mL, 56.0 mmol) in 250 mL of diethylether, CS_2 (3.4 mL, 56.0 mmol) was slowly added. The reaction mixture was stirred for 30 minutes at 0°C . The white solid was filtered, washed with diethyl ether and dried under vacuum. The dithiocarbamate salts were obtained as white powders and used in the next step without further purification.

Triethylammonium propyldithiocarbamate: white powder. Yield: 9.6 g (81%); mp = 136°C . ^1H NMR (D_2O , 300 MHz) δ (ppm) : 0.79 (t, 3H, CH_3 , $^3J_{\text{HH}} = 7.2$ Hz), 1.15 (t, 9H, CH_3 , $^3J_{\text{HH}} = 7.4$ Hz), 1.51 (sext, 2H, CH_2 , $^3J_{\text{HH}} = 7.2$ Hz), 3.06 (q, 6H, CH_2 , $^3J_{\text{HH}} = 7.4$ Hz), 3.34 (t, 2H, CH_2 , $^3J_{\text{HH}} = 7.2$ Hz). ^{13}C NMR (D_2O , 75 MHz) δ : 8.3 (CH_3), 10.7 (CH_3), 21.3 (CH_2), 46.6 (N- CH_2), 49.7 (N- CH_2^-), 209.9 (C=S).

Triethylammonium cyclopropyldithiocarbamate: Yield: 9.7 g (83%); mp = 118°C . ^1H NMR (D_2O , 300 MHz) δ (ppm) : 0.56 (m, 2H, CH_2), 0.69 (m, 2H, CH_2), 1.17 (t, 9H, CH_3 , $^3J_{\text{HH}} = 7.3$ Hz), 2.83 (m, 1H, CH), 3.09 (q, 6H, CH_2 , $^3J_{\text{HH}} = 7.3$ Hz). ^{13}C NMR (D_2O , 75 MHz) δ (ppm) : 6.2 (CH_3), 8.2 (CH_2), 29.9 (CH), 46.6 (CH_2), 212.9 (C=S).

Triethylammonium 2,2-dimethylhydrazine-1-carbodithioate: Yield: 4.1 g (35%); mp = 144°C . ^1H NMR (D_2O , 300 MHz) δ (ppm) : 1.16 (t, 9H, CH_3 , $^3J_{\text{HH}} = 7.3$ Hz), 3.08 (q, 6H, CH_2 , $^3J_{\text{HH}} = 7.3$ Hz), 3.14 (s, 6H, CH_3).

To a solution of the triethylammonium dithiocarbamate salts prepared above (54 mmol, 12.8 g for R = nPr, 12.6 g for R = cPr and 12.8 g for R = NMe₂) in acetonitrile (100 mL) chloroacetaldehyde (7.2 mL, 56.7 mmol, 50% in water) was slowly added. The pale yellow solution was stirred at room temperature overnight except for R = NMe₂ which was refluxed overnight. The volume was reduced to about 10 mL by rotary evaporation and the brown residue was slowly added to 25 mL of concentrated sulfuric acid cooled at 0° C. The reaction mixture was stirred during 20 minutes, hydrolyzed with ice water and extracted with dichloromethane. The organic phase was washed with water, dried over MgSO₄ and the solvent was removed under reduced pressure to give thiazoline-2-thione **1** which was purified by column chromatography using dichloromethane as eluent for **1-nPr** and **1-cPr** and using dichloromethane /diethyl ether (9.9/0.1) as eluent for **1-NMe₂**.

1-nPr, orange oil; 76 % yield (6.5 g); R_f = 0.55 (SiO₂, CH₂Cl₂); ¹H NMR (300 MHz) δ 0.92 (t, 3H, CH₃, ³J_{HH} = 7.4 Hz), 1.80 (sext, 2H, CH₂, ³J_{HH} = 7.4 Hz), 4.07 (t, 2H, CH₂, ³J_{HH} = 7.4 Hz), 6.56 (d, 1H, =CH, ³J_{HH} = 4.6 Hz), 7.00 (d, 1H, =CH, ³J_{HH} = 4.6 Hz). ¹³C NMR (75 MHz) δ 10.8 (CH₃), 21.5 (CH₂), 51.2 (CH₂), 110.8 (=C), 131.3 (=C), 186.9 (C=S). HRMS (ASAP) calcd. for C₆H₁₀NS₂⁺: 160.02492, found: 160.0249.

1-cPr, brown oil; 72 % yield (6.1 g); R_f = 0.58 (SiO₂, CH₂Cl₂); ¹H NMR (300 MHz) δ 0.97 (m, 2H, CH₂), 1.18 (m, 2H, CH₂), 3.42 (m, 1H, CH), 6.51 (d, 1H, =CH, ³J_{HH} = 4.7 Hz), 6.93 (d, 1H, =CH, ³J_{HH} = 4.7 Hz); ¹³C NMR (CDCl₃, 75 MHz) δ (ppm) : 7.6 (CH₂), 32.3 (CH), 110.6 (=C), 131.1 (=C), 188.9 (C=S). HRMS (ASAP) calcd for C₆H₈NS₂⁺: 158.00982, found: 158.0098; Anal. Calcd. for C₆H₇NS₂: C, 45.83 ; H, 4.49 ; N, 8.91. Found : C, 45.96 ; H, 4.35 ; N, 8.84.

1-NMe₂, orange solid; 30% yield (2.6 g); mp = 102°C. R_f = 0.53 (SiO₂, CH₂Cl₂/Et₂O (9.9-0.1)). ¹H NMR (CDCl₃, 300 MHz) δ (ppm) : 3.02 (s, 6H, CH₃), 6.53 (d, 1H, =CH, ³J_{HH} = 4.9 Hz), 7.16 (d, 1H, =CH, ³J_{HH} = 4.9 Hz). ¹³C NMR (CDCl₃, 75 MHz) δ (ppm) : 44.2 (CH₃), 109.9 (=C), 128.4 (=C), 185.0 (C=S). HRMS (ASAP) calcd for C₅H₈N₂S₂Na⁺: 183.00266, found: 183.0028. Anal. Calcd for C₅H₈N₂S₂: C, 37.47, H, 5.03, N, 17.48. Found : C, 37.12, H, 5.06, N, 17.38 .

4,5-bis(cyanoethylthio)-1,3-thiazoline-2-thiones 2: Under inert atmosphere, at -10 °C, freshly prepared LDA (12.0 mmol) was added to a solution of 1,3-thiazoline-2-

thione **1** (8.0 mmol 1.27 g for **1-nPr**, 1.26 g for **1-cPr**, 1.28 g for **1-NMe₂**) in 40 mL of anhydrous THF. After stirring for 30 minutes at -10°C, sulfur S₈ (0.38 g, 12.0 mmol) was added and the solution was stirred for an additional 30 minutes. A solution of 16.0 mmol of LDA was added to the medium. The reaction mixture was stirred for 3 hours, at -10°C, and sulfur was added (0.51 g, 16.0 mmol) followed 30 min later by the addition of 3-bromopropionitrile (6.6 mL, 80.0 mmol). The temperature was allowed to rise to room temperature and the reaction was stirred overnight. The solvent was removed under reduced pressure and the residue was dissolved in dichloromethane. The organic phase was washed with water and then dried over MgSO₄. The 4,5-biscyanoethylthio-1,3-thiazoline-2-thione **2** was purified by column chromatography using dichloromethane as eluent for **2-nPr** and **2-cPr** and using dichloromethane /diethyl ether (9.9/0.1) as eluent for **2-NMe₂**.

2-nPr, brown powder; 46 % yield (1.21 g); mp = 103 °C; R_f = 0.75 (SiO₂, CH₂Cl₂). ¹H NMR (300 MHz) δ 1.02 (t, 3H, CH₃, ³J_{HH} = 7.5 Hz), 1.78 (sext, 2H, CH₂, ³J_{HH} = 7.5 Hz), 2.75 (t, 4H, CH₂, ³J_{HH} = 7.0 Hz), 3.10 (t, 2H, CH₂, ³J_{HH} = 7.0 Hz), 3.15 (t, 2H, CH₂, ³J_{HH} = 7.0 Hz), 4.35 (m, 2H, CH₂). ¹³C NMR (75 MHz) δ 10.7 (CH₃), 18.3 (CH₂), 20.9 (CH₂), 31.3 (S-CH₂), 31.9 (CH₂), 50.7 (CH₂), 117.0 (CN), 125.1 (=C), 135.6 (=C), 186.9 (C=S). HRMS (ASAP) calcd for C₁₂H₁₆N₃S₄⁺: 330.02216, found: 330.0220; Anal. calcd for C₁₂H₁₅N₃S₄: C, 43.74, H, 4.59, N, 12.75. Found: C, 43.98, H, 4.74, N, 13.03.

2-cPr, brown powder; 75 % yield (1.96 g); mp = 113°C; R_f = 0.57 (SiO₂, CH₂Cl₂); ¹H NMR (300 MHz) δ 1.21 (m, 2H, CH₂), 1.36 (m, 2H, CH₂), 2.72 (t, 2H, CH₂, ³J_{HH} = 6.8 Hz), 2.74 (t, 2H, CH₂, ³J_{HH} = 6.8 Hz), 2.94 (m, 1H, CH), 3.08 (t, 2H, CH₂, ³J_{HH} = 6.8 Hz), 3.19 (t, 2H, CH₂, ³J_{HH} = 6.9 Hz); ¹³C NMR (75 MHz) δ 10.6 (CH₂), 18.6 (CH₂), 31.4 (CH₂), 31.9 (CH), 117.1 (CN), 126.3 (=C), 137.5 (=C), 189.9 (C=S); HRMS (ASAP) calcd for C₁₂H₁₄N₃S₄⁺: 328.00706, found: 328.0070; Anal. calcd for C₁₂H₁₃N₃S₄: C, 44.01, H, 4.00, N, 12.83. Found: C, 43.80, H, 3.80, N, 12.59.

2-NMe₂, orange-brown oil; 49 % yield (1.30 g); R_f = 0.53 (SiO₂, CH₂Cl₂/Et₂O, 9.9/0.1); ¹H NMR (300 MHz) δ 2.74 (t, 4H, CH₂, ³J_{HH} = 6.9 Hz), 3.06 (t, 2H, CH₂, ³J_{HH} = 6.9 Hz), 3.18 (t, 2H, CH₂, ³J_{HH} = 6.9 Hz), 3.25 (s, 6H, CH₃); ¹³C NMR (CDCl₃, 75 MHz) δ (ppm) : 18.4 (CH₂), 30.4 (CH₂), 31.1 (CH₂), 42.9 (CH₃), 117.3 (CN), 121.3 (=C), 138.1 (=C), 186.2 (C=S); HRMS (ESI) calcd for C₁₁H₁₄N₄S₄Na⁺: 352.9999, found:

353.0003; Anal. calcd for C₁₁H₁₄N₄S₄: C, 39.97 ; H, 4.27 ; N, 16.95 ; S, 38.81. Found: C, 40.05 ; H, 4.37 ; N, 16.95 ; S, 38.60.

Anionic gold dithiolene complexes 3-R: Under inert atmosphere and at room temperature, to a dry two-neck flask containing 1,3-thiazoline-2-thione **2-R** (0.28 g, 0.85 mmol), 5 mL of 1M solution of sodium methalonnate in MeOH was added. After complete dissolution of the compound, potassium tetrachloroaurate (III) hydrate (0.19 g, 0.5 mmol) dissolved in 7 mL of dry methanol was added to the reaction mixture followed, 5 h later, by the addition of tetraphenylphosphonium chloride (0.19 g, 0.51 mmol) in 7 mL of anhydrous methanol. The reaction was stirred overnight at room temperature under argon atmosphere. The dark green suspension was filtered under vacuum. The resulting solid was washed with MeOH and recrystallized from acetonitrile.

3-nPr: dark green crystals; 68 % yield (0.28 g); mp = 199 °C; ¹H NMR (300 MHz) δ 0.95 (t, 6H, CH₃, ³J_{HH} = 7.5 Hz), 1.75 (m, 4H, CH₂), 3.98 (m, 4H, CH₂), 7.61 (m, 8H, CH_{Ar}), 7.73 (m, 8H, CH_{Ar}), 7.89 (m, 4H, CH_{Ar}); ¹³C NMR (CDCl₃, 75 MHz) δ 11.0 (CH₃), 20.6 (CH₂), 48.9 (CH₂), 112.2 (C=C), 116.5 (C_{Ar}), 117.7 (C_{Ar}), 130.5 (C_{Ar}), 132.1 (C=C), 134.1 (C_{Ar}), 135.6 (C_{Ar}), 191.2 (C=S); UV-vis-NIR (CH₂Cl₂) λ (nm), (ε M⁻¹.cm⁻¹): 364 (5196), 294 (32095), 260 (45449), 234 (79940); HRMS (ESI) calcd for C₆₀H₅₄N₂P₂S₈Au⁺: 1317.11881, found: 1317.1192; Anal. calcd for C₃₆H₃₄AuN₂PS₈: C, 44.16, H, 3.50, N, 2.86, S, 26.20. Found: C, 44.00, H, 3.50, N, 2.70, S, 26.64.

3-cPr: dark green crystals; 52 % yield (0.22 g); mp = 257 °C; ¹H NMR (300 MHz) δ 1.14 (m, 2H, CH₂), 1.23 (m, 2H, CH₂), 2.82 (m, 1H, CH), 7.59 (m, 8H, CH_{Ar}), 7.76 (m, 8H, CH_{Ar}), 7.88 (m, 4H, CH_{Ar}). ¹³C NMR (75 MHz) δ 8.7 (CH₂), 29.5 (N-CH), 112.4 (C=C), 116.5 (C_{Ar}), 117.7 (C_{Ar}), 130.5 (C_{Ar}), 134.1 (C_{Ar}), 134.5 (C=C), 135.5 (C_{Ar}), 194.2 (C=S); UV-vis-NIR (CH₂Cl₂) λ (nm), (ε M⁻¹.cm⁻¹): 370 (31377), 396 (25628), 262 (35868), 232 (59101); HRMS (ESI) calcd for C₁₂H₁₀N₂S₈Au: 634.82808, found: 634.8281; Anal. calcd for C₃₆H₃₄AuN₂PS₈: C, 44.34, H, 3.10, N, 2.87. Found: C, 43.89, H, 2.94, N, 2.74 .

3-NMe₂: dark red crystals; 70 % yield (0.29 g); mp = 238 °C. ¹H NMR (CDCl₃, 300 MHz) δ (ppm) : 3.18 (s, 12H, CH₃), 7.61 (m, 8H, CH_{Ar}), 7.76 (m, 8H, CH_{Ar}), 7.89 (m, 4H, CH_{Ar}). ¹³C NMR (CDCl₃, 75 MHz) δ (ppm) : 42.7 (CH₃), 111.0 (C=C), 117.4 (C_{Ar}),

118.6 (C_{Ar}), 131.3 (C_{Ar}), 134.9 (C_{Ar}), 135.2 (C=C), 136.6 (C_{Ar}), 194.7 (C=S). UV-vis-NIR (CH₂Cl₂) λ (nm), (ϵ M⁻¹.cm⁻¹): 378 (20479), 314 (23592), 232 (49401); HRMS (ESI) calcd for C₁₂H₁₀N₄S₈Au⁻: 640.84988, found: 640.8501. Anal. calcd for C₃₆H₃₄AuN₂PS₈: C, 41.62 ; H, 3.29 ; N, 5.71. Found: C, 41.46 ; H, 3.13 ; N, 5.57.

Neutral gold dithiolene complexes 4-R: In a two compartment cell equipped with Pt electrodes (diameter 1mm, length 2 cm) TBAPF₆ (620 mg, 1.6 mmol) dissolved in 16 mL of CH₃CN or a mixture of CH₃CN/CH₂Cl₂ was introduced in both compartments as supporting electrolyte. The monoanionic gold complex (8 mg) was introduced in the anodic compartment. The current intensity was adjusted from 0.3 to 0.5 μ A, and the reaction was left during 5 days to 5 weeks. Crystals of the radical neutral complex were collected on the anode.

4-nPr: dark green needles. HRMS (ESI) calcd for C₁₂H₁₄N₂S₈Au⁻: 638.85938, found: 638.8596. Anal. calcd for C₁₂H₁₄AuN₂S₈: C, 22.53, H, 2.21, N, 4.38. Found: C, 22.06, H, 1.92, N, 4.33.

4-cPr: dark green needles. mp = 265 °C; HRMS (ESI) calcd for C₁₂H₁₀N₂S₈Au⁻: 634.82808, found: 634.8281.

4-NMe₂: black needles; mp = 208 °C; Anal. calcd for C₁₀H₁₂N₄S₈Au: C, 18.72, H, 1.88, N, 8.73, S, 39.97. Found: C, 18.19, H, 1.81, N, 8.21, S, 40.42.

X-ray crystal structures.

Single-crystal diffraction data were collected on APEXII, Bruker-AXS diffractometer, Mo-K α radiation (λ = 0.71073 Å) for all compounds. The structures were solved by direct methods using the *SIR97* program^[19], and then refined with full-matrix least-square methods based on *F*² (*SHELXL-97*)^[20] with the aid of the *WINGX*^[21] program. All non-hydrogen atoms were refined with anisotropic atomic displacement parameters. H atoms were finally included in their calculated positions. Details of the data collections and final refinements are given in Table 4. In the crystal structures of (Et₄N)(**3-cPr**), **4-cPr** and **4-nPr**, relatively large residual densities (2-5 e Å⁻³) observed in the last Fourier difference maps are located systematically in the vicinity of the gold atom. CCDC 1559432-1559438 contain the supplementary

crystallographic data for this paper. These data are provided free of charge by the Cambridge Crystallographic Data Centre

Table 4 Crystallographic data

	(Ph ₄ P)(3-NMe ₂)	(Ph ₄ P)(3-nPr) (CH ₃ CN) ₂	(Ph ₄ P)(3-nPr)	(Et ₄ N)(3-cPr)
Formulae	C ₃₄ H ₃₂ AuN ₄ PS ₈	C ₄₀ H ₄₀ AuN ₄ PS ₈	C ₃₆ H ₃₄ AuN ₂ PS ₈	C ₂₀ H ₃₀ AuN ₃ S ₈
FW (g.mol ⁻¹)	981.05	1061.17	979.07	765.91
System	monoclinic	monoclinic	monoclinic	triclinic
Space group	P2 ₁ /n	C2/c	C2/c	P-1
a (Å)	10.1748(3)	27.2174(4)	45.0689(7)	7.3215(4)
b (Å)	16.7965(4)	7.44430(10)	7.44970(10)	10.2503(5)
c (Å)	22.3624(5)	24.8551(4)	30.6610(5)	10.4945(6)
α (deg)	90.00	90.00	90.00	97.741(2)
β (deg)	98.1500(10)	116.0530(10)	131.1580(10)	106.169(2)
γ (deg)	90.00	90.00	90.00	110.746(2)
V (Å ³)	3783.16(17)	4524.28(12)	7750.6(2)	683.10(6)
T (K)	150(2)	293(2)	293(2)	150(2)
Z	4	4	8	1
D _{calc} (g.cm ⁻³)	1.722	1.558	1.678	1.862
μ (mm ⁻¹)	4.404	3.689	4.298	6.011
Total refls	30779	35141	40037	13319
Abs corr	multi-scan	multi-scan	multi-scan	multi-scan
T _{min} , T _{max}	0.331, 0.616	0.429, 0.691	0.902, 0.958	0.563, 0.786
Uniq refls	8667	5195	8881	3144
R _{int}	0.0421	0.0463	0.0627	0.0482
Uniq refls (I > 2σ(I))	7629	3950	6006	3061
R ₁	0.0338	0.0276	0.0444	0.0373
wR ₂ (all data)	0.0760	0.0661	0.0575	0.0921
GOF	1.040	1.113	1.075	1.035
Res. density (e ⁻ .Å ⁻³)	2.337, -1.266	0.962, -0.684	0.848, -0.53	5.796, -0.944
CCDC n°	1559432	1559433	1559434	1559435

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Table 4 Crystallographic data (continued)

	4-NMe₂	4-cPr	4-nPr
Formulae	C ₁₀ H ₁₂ AuN ₄ S ₈	C ₁₂ H ₁₀ AuN ₂ S ₈	C ₁₂ H ₁₄ AuN ₂ S ₈
FW (g.mol ⁻¹)	641.68	635.67	639.70
System	orthorhombic	monoclinic	triclinic
Space group	I222	P2 ₁ /c	P-1
a (Å)	7.0055(8)	4.7474(5)	4.4422(4)
b (Å)	13.7316(14)	6.6311(5)	7.3666(7)
c (Å)	20.649(2)	27.607(3)	14.6283(13)
α (deg)	90.00	90.00	85.148(3)
β (deg)	90.00	90.459(3)	88.746(3)
γ (deg)	90.00	90.00	74.489(3)
V (Å ³)	1986.3(4)	869.05(15)	459.61(7)
T (K)	150(2)	150(2)	150(2)
Z	4	2	1
D _{calc} (g.cm ⁻³)	2.146	2.429	2.311
μ (mm ⁻¹)	8.248	9.422	8.908
Total refls	10473	5451	7203
Abs corr	multi-scan	multi-scan	multi-scan
T _{min} , T _{max}	0.538, 0.814	0.623, 0.828	0.660, 0.837
Uniq refls	2283	1951	2044
R _{int}	0.0370	0.0620	0.0371
Uniq refls (I > 2σ(I))	2029	1479	2015
R ₁	0.0190	0.0709	0.0380
wR ₂ (all data)	0.0398	0.1486	0.0960
GOF	1.106	1.153	1.062
Res. density (e ⁻ .Å ⁻³)	1.043, -0.681	2.511, -2.438	5.591, -1.542
CCDC n°	1559436	1559437	1559438

Transport measurements.

The resistivity measurements were carried out along the long axis of the needle shaped single crystals which correspond to the stacking axis of the **4-R** complexes. Gold pads were evaporated on the crystals in order to improve the quality of the contacts and gold wires were glued with silver paste on those contacts. For **4-NMe₂** and **4-nPr**, a standard four points technique was used with a low frequency lock-in detection (applied current $I_{ac} = 0.1-1\mu\text{A}$) for measured resistances below 10 kΩ and dc measurement for higher resistances (applied current $I_{dc} = 0.1-1\mu\text{A}$). For **4-cPr**, resistivity was deduced from a two points measurement because of the small size of the crystals. The $\rho(T)$ curve was checked by two different techniques: a dc

measurement with an applied current $I_{dc} = 0.1\mu\text{A}$ and a current measurement with an applied voltage of 1V using a dc voltage source/picoammeter Keithley 6487.

Resistivity measurements were also performed under high hydrostatic pressure on very thin needles of **4-nPr** (section around $50\mu\text{m}^2$). A NiCrAl clamped cell was used up to 2.5 GPa with silicon oil (Daphne 7373) as the pressure transmitting medium. The pressure at room temperature was extracted from the resistance of a manganin gauge in the pressure cell and it is this value which is indicated in the figures. However, the loss of pressure during cooling is estimated to 0.2 GPa, slightly decreasing with pressure. The highest pressure $\rho(T)$ curve was measured in 3 points after one contact have been lost during the pressurization. Low temperatures have been provided by a home-made cryostat built around a 4K pulse tube.

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Conflict of interest

The authors declare no conflict of interest.

Keywords: S ligands • Gold • Stacking interactions • Conducting materials • Single component conductors

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Graphical Abstract

