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Inhibitors and Antiproliferative Agents**

Journal:	<i>Dalton Transactions</i>
Manuscript ID:	DT-ART-10-2014-003105.R1
Article Type:	Paper
Date Submitted by the Author:	03-Nov-2014
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## Gold(I) Thiotetrazolates as Thioredoxin Reductase Inhibitors and Antiproliferative Agents

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### Abstract

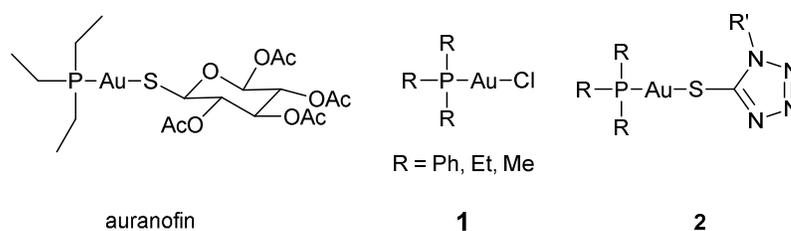
Gold(I) complexes with phosphane and thiotetrazolate ligands were prepared and investigated as new type of bioactive gold metallodrugs. The complexes triggered very efficient inhibition of the enzyme thioredoxin reductase (TrxR), which is an important molecular target for gold species. Strong cytotoxic effects were observed in MDA-MB-231 breast adenocarcinoma and HT-29 colon carcinoma cells and the complexes also caused strong effects in vincristine resistant Nalm-6 leukemia cells. Cellular uptake studies showed elevated cellular gold levels for complexes containing a triphenylphosphane ligand, whereas trifurylphosphane analogues accumulated at significantly lower cellular concentrations.

**Keywords:** cellular uptake, gold complexes, tetrazoles, thioredoxin reductase

## Introduction

The therapeutic use of gold and its complexes dates back thousands of years and it was also a hot topic in alchemy during medieval ages and the Renaissance.<sup>1</sup> However, elaboration of gold based antiproliferative agents is a comparably young and intensively developing area of medicinal and pharmaceutical chemistry.<sup>2-5</sup> The precise mechanisms of antitumor activity of gold compounds are still not completely understood. As a result of a series of mechanistic studies it was suggested that binding of gold to the selenoenzyme thioredoxin reductase (TrxR) leads to its inhibition, which results in the alteration of mitochondrial functions, increased formation of reactive oxygen species and finally causes cell death *via* apoptosis.<sup>6,7</sup> Both gold(I) and gold(III) derivatives were found to be efficient inhibitors of thioredoxin reductase<sup>6</sup>, yet gold(III) based complexes are nowadays considered as prodrugs that are reduced before binding to give gold(I) species.<sup>8</sup>

The choice of ligands appears to be one of the most challenging tasks in the design of novel gold based antitumor agents. Owing to the “soft acidic” nature of gold(I), its coordination compounds are highly prone to rapid metabolic transformations *via* interaction with numerous sulfur containing proteins and peptides *in vivo*.<sup>2</sup> Therefore, stability is a crucial point in the design of gold(I) species, and usually typical “soft bases” such as phosphorus- and sulphur-containing nucleophiles are utilized as ligands to enhance the stability of the resulting gold(I) complexes under physiological conditions. Gold(I) phosphane complexes are the most frequently investigated group of gold based compounds.<sup>3,9-15</sup> Many of them, including auranofin (Figure 1) and chlorido gold(I) phosphane (**1**, Figure 1) complexes, were shown to be effective antiproliferative agents and potent TrxR inhibitors.<sup>6,7,16</sup>



**Figure 1.** Examples of gold(I) phosphane complexes

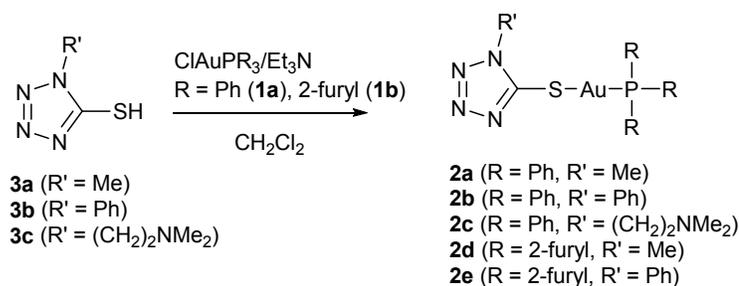
Here we report on the synthesis and biological evaluation of a new series of tetrazole-containing gold(I) phosphane complexes (**2**, Figure 1). Being rather new ligands in bioinorganic chemistry, tetrazole derivatives have been gaining more attention in recent years due to the development of more efficient synthesis methods.<sup>17</sup> Moreover, the tetrazole ring is a well known bioisoster of the carboxylic group and several tetrazole-containing platinum complexes have earlier been reported as having a promising antiproliferative profile and low toxicity *in vivo*.<sup>18-20</sup>

Gold(I) complexes with polynitrogen-containing azoles have been relatively little studied so far. Among them, mixed-ligand phosphane gold(I) 5-thiotetrazolates **2** are one of the most investigated groups.<sup>21</sup> The obvious structural analogy between this group of gold(I) complexes and auranofin makes them interesting subjects for antitumor drug design. Despite this fact, there have been up to date no data reported in the literature on their antiproliferative or other biological effects. In order to elucidate any possible structure-activity relationships within this group of gold(I) complexes, a panel of (phosphane)gold(I) 1-R-5-thiotetrazolates have been synthesized and characterized by means of spectroscopic and structural methods. Their antiproliferative activity against two human cancer cell lines as well as the inhibitory effects toward TrxR and glutathione reductase (GR) have been evaluated, whereupon cellular uptake studies were performed using high resolution atomic absorption spectroscopy (HR AAS). Additionally, antiproliferative activity of the best candidates was additionally investigated against resistant leukemia cell lines.

## Results and discussion

### Synthesis and characterization

Tetrazole-containing gold(I) complexes **2a-e** bearing triphenylphosphane or trifurylphosphane ligands were synthesized starting from corresponding 1-substituted 5-thiotetrazoles **3a-c** and chloridogold(I) phosphanes ClAuPR<sub>3</sub> [R = Ph (**1a**), 2-furyl (**1b**)] in the presence of triethylamine according to Scheme 1. Synthesis of compounds **2a** and **2b** under similar conditions was reported elsewhere earlier.<sup>22, 23</sup> Complexes **2c-e** were obtained for the first time.



**Scheme 1.** Synthesis of thiotetrazole-containing gold(I) phosphane complexes **2a-e**

Composition and structure of the products were confirmed by elemental analysis, ESI mass spectrometry, and NMR spectroscopy (<sup>1</sup>H and <sup>13</sup>C). Complexes were additionally characterized by means of IR spectroscopy and thermal analysis (TG/DSC). The molecular and crystal structure of **2e** was additionally confirmed by single crystal X-ray diffraction. In general the

spectral and structural characteristics of complexes **2a** and **2b** were found to be consistent with previously published results.<sup>23,24</sup>

ESI(+) mass spectra of **2a-e** showed pseudomolecular ions  $[M+H]^+$  along with the respective  $[Au(PR_3)_2]^+$  species. Assignments of signals observed in  $^1H$  and  $^{13}C$  NMR spectra of all complexes were made based on previously published data<sup>23, 25</sup> and are in agreement with the suggested structures (see Scheme 1). In particular,  $^1H$  NMR spectra of **2a-e** lack signals of NH protons that are observed in the spectra of uncoordinated tetrazolylthiones **3a-c** at approx. 14 ppm. Coordination also leads to significant upfield shifts (approx. 5-6 ppm) of the C(5) signals, which are observed at approx. 158 ppm in the  $^{13}C$  NMR spectra of **2a-e** (163-164 ppm in the spectra of **3a-c**). Other resonances remain mainly unchanged when compared to the spectra of the starting materials. Due to  $^{13}C$ - $^{31}P$  coupling, phenyl and furyl carbons of the phosphane ligands appear as doublet resonances.

When compared with the spectra of the free ligands **3a-c**, IR spectra of **2a-e** show the same pattern of changes including disappearance of the intensive bands at 1510, 1350-1360, 1040-1050, and 790-800  $cm^{-1}$  characteristic for the stretching and deformation vibrations of C=S and N-C=S fragments.<sup>22, 23</sup> New partially splitted bands at 690-710  $cm^{-1}$  assigned to the stretching vibrations of C-S bond<sup>23</sup> were observed. Altogether, these changes support the formation of a covalent Au-S bond between the triarylphosphane gold(I) moiety and the thiolate function of the tetrazole-containing ligands.

According to the results of DSC/TG, all the complexes were found to be thermally stable up to 180-200 °C. Exothermal decomposition of **2a-c** and **2e** at higher temperatures is preceded by their melting at 180-195 °C, while **2d** melts at noticeably lower temperature (approx. 120 °C).

### Crystal and molecular structure of complex **2e**

Complex **2e** crystallizes in the monoclinic space group  $C2/c$ , with eight formula units in the unit cell. Crystal data and structure refinement details for this compound are summarized in Table 1. Similarly to the previously reported complexes **2a** and **2b**,<sup>24</sup> complex **2e** presents a molecular complex showing bidentate linear coordination environment of the gold atoms (Figure 2a).

In complex **2e**, the gold atom is in a standard linear coordination, being attached to the phosphorus atom with an Au1-P1 distance of 2.2475(6) Å and to the sulfur atom with an Au1-S1 distance of 2.3051(7) Å. The coordination angle P1-Au1-S1 takes on a value of 170.93(3)°. The geometries of the coordination units are very close in three compounds **2a**, **2b** and **2e** (see Table 2).

In the complex **2e**, the oxygen atoms of the furane rings are oriented away from the gold atom. The angles Au1-P1-C12 [116.21(8)°] and Au1-P1-C22 [114.05(9)°] are larger than the

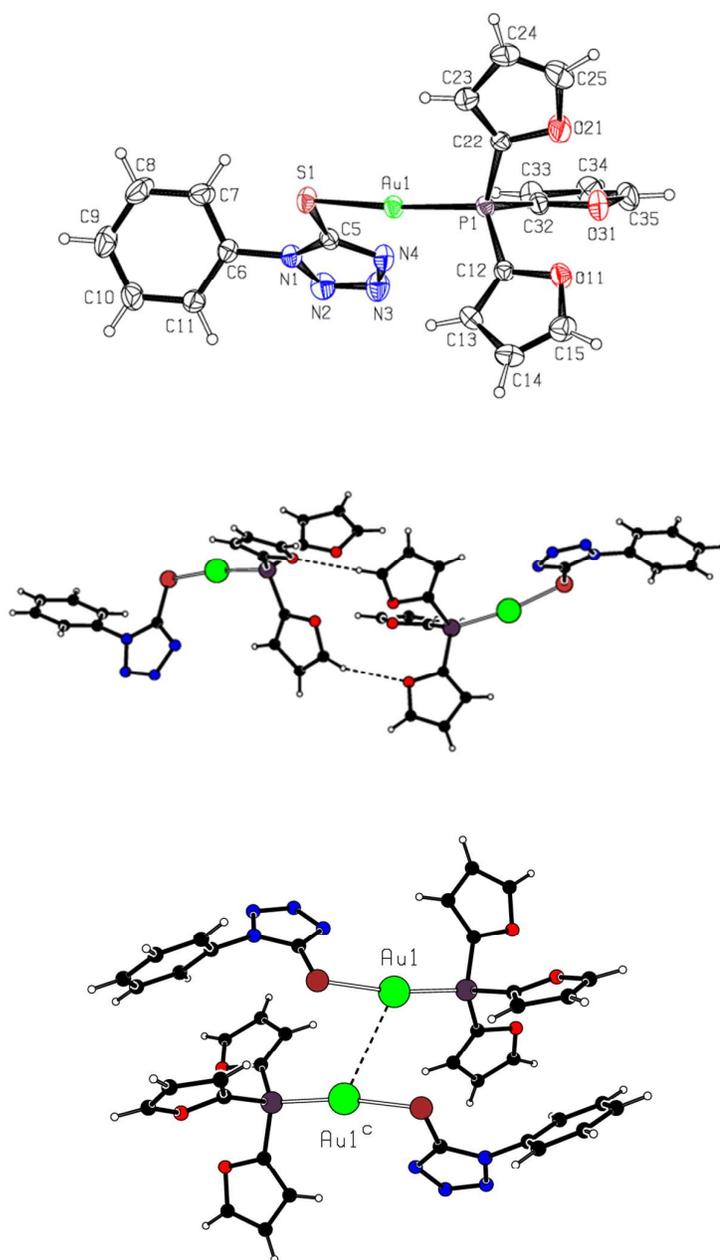
tetrahedral standard, whereas the angle Au1–P1–C32 [107.76(9)°] is close to that but somewhat smaller. The phosphane C–P–C valence angles, lying in a range of 103.83(12)–107.55(12)°, are smaller than the tetrahedral one. These geometric features are very close to those in complexes **2a** and **2b**. The fragment AuPR<sub>3</sub> (R = 2-furyl) displays a distorted not propeller-like structure due to the large variation in dihedral angles Au–P–C–C [–5.8 (3), 8.1 (3) and 16.6 (4)°].

As for the tetrazole ring geometry, it is typical of 1*H*-tetrazoles in the three complexes **2a**, **2b** and **2e** (see Table 2), with the shortest bond N2–N3 and remaining bond distances lying in rather narrow ranges. In the crystal structure of complex **2e**, there are non-classic intermolecular hydrogen bonds C35–H35⋯O11<sup>b</sup> between two neighboring complex molecules [symmetry transformation: (*b*)  $-x, y, -z + \frac{1}{2}$ ; hydrogen bond geometry: H35⋯O11<sup>b</sup> 2.55 Å, C35⋯O11<sup>b</sup> 3.424(3) Å, C35–H35⋯O11<sup>b</sup> 157°]. These bonds form dimeric entities comprising *R*<sup>2</sup><sub>2</sub>(14) hydrogen-bonded rings (Figure 2*b*).

The complex molecules are bonded into dimeric units through very weak aurophilic bonding,<sup>26</sup> with separation of the gold atoms Au1...Au1<sup>c</sup> of 3.3892(5) Å (symmetry transformation as in Fig. 2*c*). These units are connected *via* the mentioned above hydrogen bonds to give polymeric chains running along the *a* axis.

**Table 1.** Main crystal data and structure refinement details for complexes **2e**.<sup>a</sup>

Formula	C <sub>19</sub> H <sub>14</sub> AuN <sub>4</sub> O <sub>3</sub> PS
Formula weight	606.34
T/K	296(2)
Crystal system	Monoclinic
Space group	<i>C</i> 2/ <i>c</i>
Crystal size /mm	0.35×0.34×0.12
<i>a</i> /Å	13.79810(10)
<i>b</i> /Å	13.08130(10)
<i>c</i> /Å	22.1996(3)
β/°	91.0696(5)
<i>V</i> /Å <sup>3</sup>	4006.27(7)
<i>Z</i>	8
<i>D</i> <sub>c</sub> /g cm <sup>-3</sup>	2.011
μ/mm <sup>-1</sup>	7.557
Reflections	5371
Restraints	0
Parameters	262
<i>R</i> 1/ <i>wR</i> 2[ <i>I</i> >2σ( <i>I</i> )]	0.0210/0.0484
<i>R</i> 1/ <i>wR</i> 2 [all data]	0.0283/0.0505
Goodness-of-fit	1.004



**Figure 2.** top: complex **2e**, with the atom numbering scheme and displacement ellipsoids drawn at the 30% probability level; middle: hydrogen-bonded dimer in the crystal structure of **2e**; bottom: dimeric unit formed by aurophilic bonding in the crystal structure of **2e** [symmetry transformation: (c)  $1 - x, y, \frac{1}{2} - z$ ].

**Table 2.** Some geometric features ( $\text{\AA}$ ,  $^\circ$ ) of complexes **2a**, **2b** and **2e**.

Compound	<b>2a</b> <sup>a</sup>	<b>2b</b> <sup>a</sup>	<b>2e</b>
Au1–P1	2.261(1)	2.265(2)	2.2475(6)
Au1–S1	2.325(1)	2.304(2)	2.3051(7)
P1–Au1–S1	170.11(3)	172.68(5)	170.93(3)
C5–S1–Au1	97.5(1)	105.3(2)	99.28(9)
S1–C5	1.740(4)	1.733(7)	1.726(3)
N1–C5	1.342(5)	1.346(9)	1.350(3)
N1–N2	1.354(5)	1.378(8)	1.361(3)
N2–N3	1.289(5)	1.295(9)	1.285(4)
N3–N4	1.354(5)	1.366(8)	1.359(4)
N4–C5	1.331(5)	1.346(8)	1.326(3)

<sup>a</sup> from ref. <sup>24</sup>**Table 3.** Antiproliferative activity and inhibition activity against TrxR and GR of gold(I) phosphane complexes **1a-b** and **2a-e**, n.d.: not determined.

Compound	IC <sub>50</sub> , $\mu\text{M}$		Selectivity GR/TrxR (x-fold)	IC <sub>50</sub> , $\mu\text{M}$	
	TrxR	GR		MDA-MB-231	HT-29
<b>1a</b>	0.256 $\pm$ 0.002 <sup>27</sup>	4.2 $\pm$ 0.7 <sup>27</sup>	16	9.5 $\pm$ 0.7	4.2 $\pm$ 0.9 <sup>10</sup>
<b>1b</b>	0.207 $\pm$ 0.021	n.d.		9.9 $\pm$ 0.7	10.0 $\pm$ 1.8
<b>2a</b>	0.037 $\pm$ 0.010	0.97 $\pm$ 0.25	26	7.9 $\pm$ 1.9	12.0 $\pm$ 2.7
<b>2b</b>	0.046 $\pm$ 0.012	0.55 $\pm$ 0.04	12	7.8 $\pm$ 2.0	10.2 $\pm$ 1.5
<b>2c</b>	0.165 $\pm$ 0.065	0.66 $\pm$ 0.19	4	8.9 $\pm$ 1.1	8.6 $\pm$ 0.7
<b>2d</b>	0.033 $\pm$ 0.005	0.43 $\pm$ 0.05	13	8.5 $\pm$ 1.5	13.3 $\pm$ 2.9
<b>2e</b>	0.053 $\pm$ 0.001	0.51 $\pm$ 0.07	10	9.3 $\pm$ 0.2	11.5 $\pm$ 1.7
<b>3a</b>	n.d.	n.d.	-	> 100	> 100
<b>3b</b>	n.d.	n.d.	-	> 100	> 100
<b>3c</b>	n.d.	n.d.	-	> 100	> 100

**TrxR and GR inhibition activity**

As mentioned above, inhibition of selenoenzyme thioredoxin reductase (TrxR) is considered to be an important mechanism of bioactivity of gold(I) species.<sup>3, 6, 7</sup> Therefore, the potential of tetrazole-based gold(I) complexes to inhibit TrxR was studied on isolated rat liver TrxR using the DTNB (dithiobisnitrobenzoic acid) reduction assay. As far as selenocysteine is considered to be the relevant binding site for gold complexes, another structurally and functionally similar disulfide reductase, i.e. glutathione reductase (GR, from yeast), was included into the test as a reference. Instead of selenocysteine, GR contains a cysteine residue in its active site and thus can be used to check the specificity of TrxR inhibition by the tested compounds.<sup>28, 29</sup>

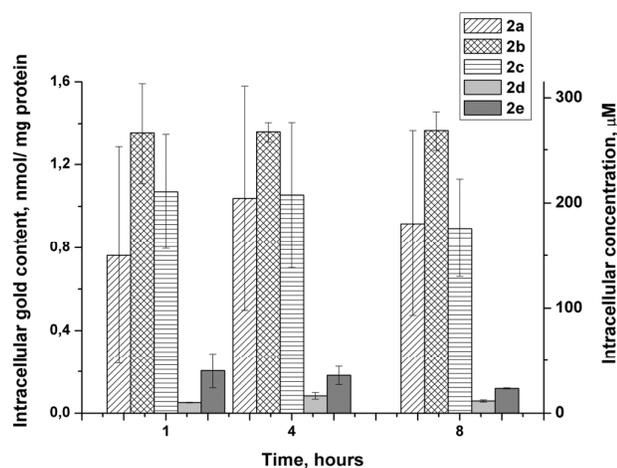
According to the results given in Table 3, tetrazole-containing gold(I) complexes turned out to be very effective inhibitors of TrxR with  $IC_{50}$  values in the low nanomolar range. With the exception of **2c**, all complexes showed  $IC_{50}$  values in the range 30–50 nM and at least 10-fold selectivity for inhibition of TrxR over GR. In all cases **2a-e** were more effective enzyme inhibitors than the respective chlorido complexes **1a** and **1b**.

### Antiproliferative activity

Antiproliferative activity of thiotetrazole-containing gold(I) complexes **2a-e** against two human cancer cell lines (MDA-MB-231 and HT-29) was assessed using the crystal violet assay. Chlorido(triphenylphosphane)gold(I) (**1a**) and chlorido[tri(2-furyl)phosphane]gold(I) (**1b**) were included into the test as positive controls. According to the results (Table 3), the tetrazole-containing complexes **2a-e** demonstrate considerable antiproliferative activity against both human cancer cell lines with  $IC_{50}$  values found in the micromolar or low micromolar range (approx. 8.0-13.0  $\mu$ M). These values are comparable to those of the positive controls and the platinum anticancer drug cisplatin ( $IC_{50}$  values of 7.0  $\mu$ M in HT-29 cells<sup>30</sup> and 4.0  $\mu$ M in MDA-MB-231 cells<sup>31</sup>). The metal-free thiotetrazole derivatives **3a-3c** were investigated as negative controls and showed no inhibition of tumor cell proliferation.

### Cellular uptake

In order to evaluate the extent of cellular uptake, we measured the gold content in HT-29 cells exposed to 10.0  $\mu$ M (approx. equal to the  $IC_{50}$  calculated for this cell line) solutions of the tested compounds for a period of 1, 4 and 8 h by a method based on high resolution continuum source atomic absorption spectroscopy (HR-CS AAS)<sup>32, 33</sup>. Results were corrected for the respective protein contents of the samples and presented in Figure 3 as nmol gold per milligram cellular protein. Besides, based on certain biophysical parameters of HT-29 cells<sup>32</sup>, the nmol Au/mg protein values obtained were used to estimate the respective intracellular molar concentrations.



**Figure 3.** Cellular gold content of HT-29 cells exposed to 10  $\mu\text{M}$  of gold(I) complexes **2a-2e**

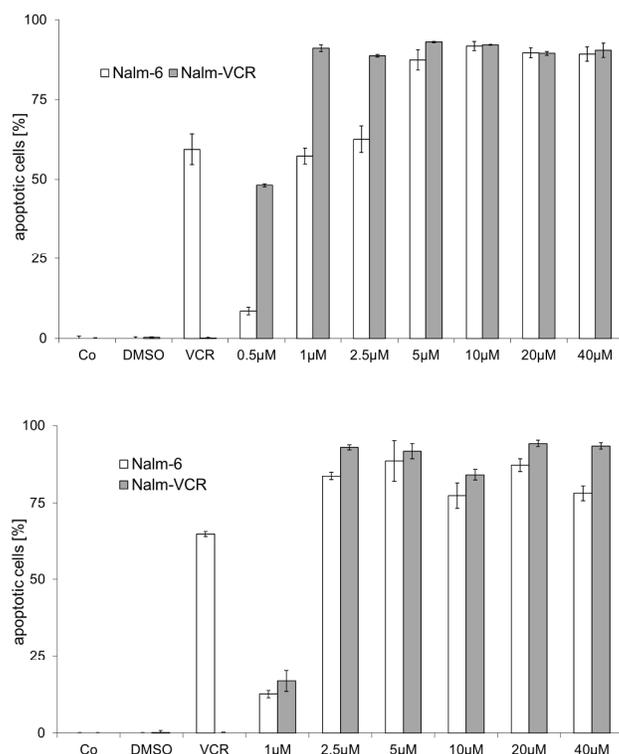
As shown in Figure 4, the highest values of intracellular accumulation were observed for triphenylphosphine gold(I) complexes **2a-2c**. The content of gold in HT-29 cells reached by these complexes after 1 h of exposure correlated to intracellular concentrations of 150 (**2a**), 265 (**2b**) and 210 (**2c**)  $\mu\text{M}$ , respectively. Taking into account the exposure concentration of 10.0  $\mu\text{M}$ , one can conclude that for these complexes at least 15-fold accumulation can be achieved already within the first hour of the experiment. This is markedly higher than the 8-fold accumulation reported for the chlorido derivative **1a** after 6 h of exposure.<sup>10</sup> Drastically lower levels of cellular uptake were found for [tri(2-furyl)phosphane]gold(I) complexes **2d** and **2e**, which can be explained by their lower lipophilicity. Intracellular concentrations of 10 and 40  $\mu\text{M}$  observed for **2d** and **2e**, respectively, after 1 h of exposure are 7–15 times lower in comparison with their triphenylphosphane-based analogs **2a** and **2b**.

#### Antiproliferative activity in a resistant cell line

Resistance to therapeutic agents is a major problem in current tumor chemotherapy. Hence, the development of drugs that can overcome drug resistance in tumor tissues is of major interest. Here we used wild type and vincristine resistant NALM-6 cells, which overexpress the drug efflux pump p-glycoprotein,<sup>33</sup> to evaluate the possible overcoming of drug resistance by the gold complexes under study. Complexes **2a** and **2d** were selected for these experiments as examples with high (**2a**) and low (**2d**) cellular uptake but comparable toxicity and TrxR inhibitory potential (see above). Apoptosis induction was assessed as DNA fragmentation in the cells.

In fact, both **2a** and **2d** triggered strong effects in wild type and vincristine resistant NALM-6 cells already in low micromolar concentrations (> 50% DNA fragmentation at 1.0  $\mu\text{M}$  with **2a**

and 2.5  $\mu\text{M}$  with **2d**), which clearly demonstrated that both complexes could overcome p-glycoprotein related multidrug resistance.



**Figure 4.** Effects of **2a** (top) and **2d** (bottom) on DNA fragmentation in wild type and vincristine resistant Nalm-6 cells after 72 h of exposure; Co: control, DMSO: cells treated with same amount of DMSO, VCR: vincristine (20 nM)

## Conclusions

A series of different gold(I) phosphane complexes with thiotetrazolato ligands were prepared and biologically evaluated. The crystal structure of complex **2e** showed the expected linear coordination geometry around the gold center. Interestingly, dimers were formed by intermolecular hydrogen bonding between the furane moieties.

Complexes **2a-e** turned out to be very effective inhibitors of the enzyme TrxR. Comparison with the respective chloride analogues **1a** and **1b** indicated that the thiotetrazolato ligand was important for this effect. The lowest activity in the series **2a-e** was observed for **2c**, which contains an aliphatic side chain with a tertiary nitrogen at the thiotetrazolato ligand. Notably, strong TrxR inhibition with thiophenolate containing gold(I) carbene complexes was also observed in a previous study.<sup>34</sup> However, the enhanced activity against TrxR did not translate into elevated cytotoxicity since complexes **2a-e** were of comparable activity with **1a** and **1b** in cell proliferation assays.

Cellular uptake studies showed a very high accumulation of the triphenylphosphane containing complexes **2a-c**, whereas the triphenylfurane analogues **2d** and **2e** were found at significantly lower levels in the cells. It can be speculated that possible dimer formation by hydrogen bonding of the trifurylphosphane moiety might contribute to this effect besides differences in the lipophilicity. It should be mentioned that we observed higher cellular accumulation of triphenylphosphane gold complexes compared to trialkylphosphane derivatives in a number of recent studies<sup>10, 11, 35</sup> and that lipophilicity is generally considered as an important parameter for cellular uptake of gold phosphane species.<sup>36</sup>

Importantly, selected compounds **2a** and **2d** showed decent activity in p-glycoprotein overexpressing vincristine-resistant Nalm-6 cells, which indicates that this type of gold compounds might be useful for the design of resistance overcoming drugs.

In summary, gold complexes with thiotetrazolate represent a new promising class of gold species, which trigger important biological effects such as TrxR inhibition and cytotoxicity.

## Experimental

### Materials and methods

All reagents and solvents were used as received from Sigma, Aldrich, or Acros Organics. <sup>1</sup>H and <sup>13</sup>C NMR spectra were recorded on a Bruker DRX-400 AS NMR System. ESI(+) mass spectra were recorded on a Bruker micrOTOF instrument using dichloromethane as a solvent. The purity of the target compounds (>95%) was confirmed by elemental analysis. For all compounds undergoing biological evaluation, the experimental values differed less than 0.5% from the calculated ones.

IR spectra of powder samples were registered on a Nicolet Thermo Avatar 330 spectrometer using Smart Diffuse Reflection accessory in the range of 4000–400 cm<sup>-1</sup>. The TG and DSC curves were obtained with a NETZSCH STA 429 thermoanalyzer in a dynamic nitrogen atmosphere (heating rate of 10 K·min<sup>-1</sup>, aluminum oxide, mass 5–6 mg and temperature range from room temperature up to 600 °C).

### Synthesis of gold(I) complexes **2a-e**

Compounds **2a-c** were obtained using the following general procedure. 0.5 mmol (248 mg) of chlorido(triphenylphosphane)gold(I) and 0.5 mmol of corresponding 1-R'-5-thiotetrazole **3a-c** were dissolved in 20 ml of CH<sub>2</sub>Cl<sub>2</sub>. Triethylamine (0.1 mL) was added dropwise to the prepared solution, and the mixture was stirred overnight at room temperature. Then the mixture was evaporated to dryness, and the residue was dissolved in hot THF. After cooling to the room

temperature, needle-like crystals of triethylammonium chloride formed and were separated by filtration. The filtrate was evaporated again, and the solid residue was recrystallized from CH<sub>2</sub>Cl<sub>2</sub>/hexane (**2a** and **2b**) or acetonitrile (**2c**) to obtain the colorless product.

**(1-Methyltetrazol-5-ylthiolato)(triphenylphosphane)gold(I) (2a)**

Yield (from CH<sub>2</sub>Cl<sub>2</sub>/hexane): 235 mg (82%). *Anal.* Calc. for C<sub>20</sub>H<sub>18</sub>N<sub>4</sub>PSAu: C, 41.82; H, 3.16; N, 9.75; S, 5.58. Found: C, 42.31; H, 3.30; N, 9.97; S, 5.32%. MS (ESI<sup>+</sup>, 100 V, CH<sub>2</sub>Cl<sub>2</sub>, m/z): 575.07 [M+H]<sup>+</sup>; 721.15 [Au(PPh<sub>3</sub>)<sub>2</sub>]<sup>+</sup>. DSC/TG: 189–190 °C (mp); 282 °C (T<sub>max</sub>, exothermal dec.). <sup>1</sup>H NMR [400 MHz, CDCl<sub>3</sub>, δ (ppm)]: 7.62–7.47 (m, 15H, C<sub>6</sub>H<sub>5</sub>); 3.96 (s, 3H, CH<sub>3</sub>). <sup>13</sup>C NMR [100 MHz, CDCl<sub>3</sub>, δ (ppm), *J* (Hz)]: 158.0 (s, C(5)<sup>tz</sup>); 134.3 (d, <sup>2</sup>*J*(C,P) = 13.8, *ortho*-C from PPh<sub>3</sub>); 131.9 (d, <sup>4</sup>*J*(C,P) = 2.5, *para*-C from PPh<sub>3</sub>); 129.3 (d, <sup>3</sup>*J*(C,P) = 11.7, *meta*-C from PPh<sub>3</sub>); 128.9 (d, <sup>1</sup>*J*(C,P) = 59.0, *ipso*-C from PPh<sub>3</sub>); 34.0 (s, CH<sub>3</sub><sup>tz</sup>). FT-IR (ν, cm<sup>-1</sup>): 3054 (s), ν(CH<sub>aryl</sub>); 2942 (m), ν(CH<sub>alkyl</sub>); 1478 (s), ν(CC<sub>aryl</sub>); 1436 (vs), 1377 (s), δ(CH<sub>aryl</sub>)+δ(CH<sub>alkyl</sub>); 1332 (m), 1313 (m), 1264 (s), 1222 (m), 1162 (s), ν(N=N<sub>tz</sub>)+ν(N-N<sub>tz</sub>); 1101 (s), 1077 (m), 1026 (m), 996 (m), 969 (m), δ(CN<sub>4</sub>)+δ(C<sub>6</sub>H<sub>5</sub>); 749 (vs), ν(C–P); 693 (vs), ν(C–S).

**(1-Phenyltetrazol-5-ylthiolato)(triphenylphosphane)gold(I) (2b)**

Yield (from CH<sub>2</sub>Cl<sub>2</sub>/hexane): 290 mg (91%). *Anal.* Calc. for C<sub>25</sub>H<sub>20</sub>N<sub>4</sub>PSAu: C, 47.18; H, 3.17; N, 8.80; S, 5.04. Found: C, 47.32; H, 3.22; N, 9.08; S, 4.88%. MS (ESI<sup>+</sup>, 100 V, CH<sub>2</sub>Cl<sub>2</sub>, m/z): 637.09 [M+H]<sup>+</sup>; 721.15 [Au(PPh<sub>3</sub>)<sub>2</sub>]<sup>+</sup>. DSC/TG: 195–196 °C (mp); 213 °C (T<sub>max</sub>, exothermal dec.). <sup>1</sup>H NMR [400 MHz, CDCl<sub>3</sub>, δ (ppm)]: 7.76–7.74 (m, 2H, CH from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>); 7.59–7.40 (m, 18H, CH from C<sub>6</sub>H<sub>5</sub><sup>tz</sup> and PPh<sub>3</sub>); 3.96 (s, 3H, CH<sub>3</sub>). <sup>13</sup>C NMR [100 MHz, CDCl<sub>3</sub>, δ (ppm), *J* (Hz)]: 158.1 (s, C(5)<sup>tz</sup>); 135.2 (s, *ipso*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>); 134.3 (d, <sup>2</sup>*J*(C,P) = 13.8, *ortho*-C from PPh<sub>3</sub>); 131.9 (s, *meta*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>), 131.9 (d, <sup>4</sup>*J*(C,P) = 2.3, *para*-C from PPh<sub>3</sub>); 129.3 (d, <sup>3</sup>*J*(C,P) = 11.7, *meta*-C from PPh<sub>3</sub>); 129.1 (s, *para*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>), 128.9 (d, <sup>1</sup>*J*(C,P) = 59.0, *ipso*-C from PPh<sub>3</sub>); 124.6 (s, *ortho*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>). FT-IR (ν, cm<sup>-1</sup>): 3074 (w), 3051 (w), 3032 (w), ν(CH<sub>aryl</sub>); 1595 (m), 1500 (s), 1481 (m), 1435 (vs), 1418 (m), 1402 (m), 1377 (s), ν(CC<sub>aryl</sub>)+δ(CH<sub>aryl</sub>); 1330 (w), 1311 (w), 1271 (s), 1229 (s), 1182 (m), 1159 (m), ν(N=N<sub>tz</sub>)+ν(N-N<sub>tz</sub>); 1103 (vs), 1079 (m), 1028 (m), 998 (m), 980 (w), δ(CN<sub>4</sub>)+δ(C<sub>6</sub>H<sub>5</sub>); 758 (vs), 748 (vs), ν(C–P); 713 (s), 696 (vs) ν(C–S).

**[1-(2-(N-dimethylamino)ethyl)tetrazol-5-ylthiolato](triphenylphosphane)gold(I) (2c)**

Yield (from acetonitrile): 265 mg (84%). *Anal.* Calc. for C<sub>23</sub>H<sub>25</sub>N<sub>5</sub>PSAu: C, 43.75; H, 3.99; N, 11.09; S, 5.08. Found: C, 43.72; H, 3.44; N, 11.07; S, 5.01%. MS (ESI<sup>+</sup>, 100 V, CH<sub>2</sub>Cl<sub>2</sub>, m/z): 632.13 [M+H]<sup>+</sup>; 721.15 [Au(PPh<sub>3</sub>)<sub>2</sub>]<sup>+</sup>. DSC/TG: 195–196 °C (mp); 256 °C (T<sub>max</sub>, exothermal

dec.).  $^1\text{H}$  NMR [400 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm),  $J$  (Hz)]: 7.62–7.47 (m, 15H,  $\text{C}_6\text{H}_5$  from  $\text{PPh}_3$ ); 4.41 (t,  $^3J(\text{H,H}) = 7.2$ , 2H,  $\alpha\text{-CH}_2$ ); 2.85 (t,  $^3J(\text{H,H}) = 7.2$ , 2H,  $\beta\text{-CH}_2$ ); 2.32 (s, 6H,  $\text{N}(\text{CH}_3)_2$ ).  $^{13}\text{C}$  NMR [100 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm),  $J$  (Hz)]: 157.6 (s,  $\text{C}(5)^{\text{tz}}$ ); 134.3 (d,  $^2J(\text{C,P}) = 13.8$ , *ortho*-C from  $\text{PPh}_3$ ); 131.9 (d,  $^4J(\text{C,P}) = 2.5$ , *para*-C from  $\text{PPh}_3$ ); 129.3 (d,  $^3J(\text{C,P}) = 11.7$ , *meta*-C from  $\text{PPh}_3$ ); 128.9 (d,  $^1J(\text{C,P}) = 59.0$ , *ipso*-C from  $\text{PPh}_3$ ); 57.5 (s,  $\alpha\text{-CH}_2$ ); 45.5 (s,  $\text{N}(\text{CH}_3)_2$ ); 45.4 (s,  $\beta\text{-CH}_2$ ). FT-IR ( $\nu$ ,  $\text{cm}^{-1}$ ): 3067 (m), 3049 (m),  $\nu(\text{CH}_{\text{aryl}})$ ; 2965 (s), 2940 (s), 2856 (m), 2825 (s), 2791 (m),  $\nu(\text{CH}_{\text{alkyl}})$ ; 1585 (m), 1480 (s),  $\nu(\text{CC}_{\text{aryl}})$ ; 1437 (vs), 1380 (s),  $\delta(\text{CH}_{\text{aryl}})+\delta(\text{CH}_{\text{alkyl}})$ ; 1301 (s), 1260 (m), 1250 (m), 1179 (s), 1160 (m),  $\nu(\text{N}=\text{N}_{\text{tz}})+\nu(\text{N}-\text{N}_{\text{tz}})$ ; 1100 (s), 1066 (m), 1023 (m), 997 (m), 982 (m), 920 (m),  $\delta(\text{CN}_4)+\delta(\text{C}_6\text{H}_5)$ ; 777 (m),  $\nu(\text{C}-\text{N})$ ; 750 (vs),  $\nu(\text{C}-\text{P})$ ; 710 (s), 690 (vs)  $\nu(\text{C}-\text{S})$ .

Complexes **2d** and **2e** were obtained using the same procedure starting from chlorido[tri(2-furyl)phosphane]gold(I) and corresponding 1-R-5-thiotetrazole (R = Me (**3a**), Ph (**3b**)).

#### (1-Methyltetrazol-5-ylthiolato)[tri(2-furyl)phosphane]gold(I) (**2d**)

Yield (from THF/hexane): 220 mg (81%). *Anal.* Calc. for  $\text{C}_{14}\text{H}_{12}\text{N}_4\text{O}_3\text{PSAu}$ : C, 30.89; H, 2.22; N, 10.29. Found: C, 30.97; H, 2.38; N, 10.71%. MS ( $\text{ESI}^+$ , 100 V,  $\text{CH}_2\text{Cl}_2$ ,  $m/z$ ): 545.01  $[\text{M}+\text{H}]^+$ ; 661.03  $[\text{Au}(2\text{-furyl})_2]^+$ . DSC/TG: 120–122 °C (mp); 232 °C ( $T_{\text{max}}$ , exothermal dec.).  $^1\text{H}$  NMR [400 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm)]: 7.80 (m, 3H,  $\text{CH}^{\text{furyl}}$ ); 7.34 (m, 3H,  $\text{CH}^{\text{furyl}}$ ); 6.57 (m, 3H,  $\text{CH}^{\text{furyl}}$ ); 3.97 (s, 3H,  $\text{CH}_3$ ).  $^{13}\text{C}$  NMR [100 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm),  $J$  (Hz)]: 157.8 (s,  $\text{C}(5)^{\text{tz}}$ ); 150.0 (d,  $^3J(\text{C,P}) = 6.4$ , C(5) from furyl); 141.3 (d,  $^1J(\text{C,P}) = 93.8$ , C(2) from furyl); 125.8 (d,  $^2J(\text{C,P}) = 27.1$ , C(3) from furyl); 111.7 (d,  $^3J(\text{C,P}) = 9.8$ , C(4) from furyl); 34.1 (s,  $\text{CH}_3$ ). FT-IR ( $\nu$ ,  $\text{cm}^{-1}$ ): 3115 (m), 2981 (w),  $\nu(\text{CH}_{\text{aryl}})$ ; 2951 (w),  $\nu(\text{CH}_{\text{alkyl}})$ ; 1546 (m), 1465 (m), 1450 (m),  $\nu(\text{CC}_{\text{aryl}})$ ; 1381 (m), 1366 (m),  $\delta(\text{CH}_{\text{aryl}})+\delta(\text{CH}_{\text{alkyl}})$ ; 1271 (m), 1215 (m),  $\nu(\text{N}=\text{N}_{\text{tz}})+\nu(\text{N}-\text{N}_{\text{tz}})$ ; 1170 (m),  $\nu(\text{C}-\text{O})$ ; 1130 (m), 1110 (s), 910 (m), 881 (m),  $\delta(\text{CN}_4)+\delta(\text{furyl})$ ; 760 (s),  $\nu(\text{C}-\text{P})$ ; 705 (m)  $\nu(\text{C}-\text{S})$ .

#### (1-Phenyltetrazol-5-ylthiolato)[tri(2-furyl)phosphane]gold(I) (**2e**)

Yield (from  $\text{CH}_2\text{Cl}_2$ /hexane): 270 mg (89%). *Anal.* Calc. for  $\text{C}_{19}\text{H}_{14}\text{N}_4\text{O}_3\text{PSAu}$ : C, 37.64; H, 2.33; N, 9.24. Found: C, 37.43; H, 2.27; N, 9.12%. MS ( $\text{ESI}^+$ , 100 V,  $\text{CH}_2\text{Cl}_2$ ,  $m/z$ ): 607.03  $[\text{M}+\text{H}]^+$ ; 661.03  $[\text{Au}(2\text{-furyl})_2]^+$ . DSC/TG: 179–180 °C (mp); 214 °C ( $T_{\text{max}}$ , exothermal dec.).  $^1\text{H}$  NMR [400 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm)]: 7.80–7.79 (m, 3H,  $\text{CH}^{\text{furyl}}$ ); 7.78–7.76 (m, 2H,  $\text{C}_6\text{H}_5^{\text{tz}}$ ); 7.54–7.51 (m, 2H,  $\text{C}_6\text{H}_5^{\text{tz}}$ ); 7.48–7.45 (m, 1H,  $\text{C}_6\text{H}_5^{\text{tz}}$ ); 7.34–7.32 (m, 3H,  $\text{CH}^{\text{furyl}}$ ); 6.57–6.56 (m, 3H,  $\text{CH}^{\text{furyl}}$ ).  $^{13}\text{C}$  NMR [100 MHz,  $\text{CDCl}_3$ ,  $\delta$  (ppm),  $J$  (Hz)]: 157.8 ( $\text{C}(5)^{\text{tz}}$ ); 150.0 (d,  $^3J(\text{C,P}) = 6.3$ , C(5) from furyl); 141.4 (d,  $^1J(\text{C,P}) = 92.5$ , C(2) from furyl); 135.2 (s, *ipso*-C from  $\text{C}_6\text{H}_5^{\text{tz}}$ ); 129.3

(s, *meta*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>); 129.2 (s, *para*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>); 125.7 (d, <sup>2</sup>J(C,P) = 27.1, C(3) from furyl); 124.5 (s, *ortho*-C from C<sub>6</sub>H<sub>5</sub><sup>tz</sup>); 111.7 (d, <sup>3</sup>J(C,P) = 9.8, C(4) from furyl). FT-IR (ν, cm<sup>-1</sup>): 3158 (m), 3122 (m), ν(CH<sub>aryl</sub>); 1597 (m), 1550 (m), 1496 (s), ν(CC<sub>aryl</sub>); 1459 (m), 1397 (m), 1369 (s), δ(CH<sub>aryl</sub>); 1313 (m), 1266 (m), 1217 (m), ν(NN<sub>tz</sub>)+ν(N-N<sub>tz</sub>); 1170 (s), (C-O); 1130 (s), 1100 (m), 1080 (m), 1040 (m), 1010 (vs), 909 (m), 881 (m), 830 (m), δ(CN<sub>4</sub>)+δ(furyl); 752 (vs), ν(C-P); 692 (s) ν(C-S).

### Crystallographic studies

Single crystal X-ray data of complex **2e** were collected at room temperature on a SMART APEX II diffractometer using graphite monochromated Mo-K $\alpha$  radiation ( $\lambda = 0.71073 \text{ \AA}$ ). The reflection data were corrected on absorption. The structures were solved by direct methods with the program SIR2004<sup>37</sup> and refined on  $F^2$  by full-matrix least squares technique with SHELXL-2013.<sup>38</sup> All non-hydrogen atoms were refined anisotropically. The hydrogen atoms were placed in calculated positions and refined using a “riding” model, with  $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$ . Molecular graphics was performed with the program PLATON.<sup>39</sup> Crystallographic data of complex **2e** (excluding structure factors) have been deposited with the Cambridge Crystallographic Data Centre, CCDC, 12 Union Road, Cambridge CB21EZ, UK. Copies of the data can be obtained free of charge on quoting the depository number CCDC-1027451. (Fax: +44-1223-336-033; E-Mail: deposit@ccdc.cam.ac.uk, <http://www.ccdc.cam.ac.uk>)

### TrxR/GR inhibition assay

To determine the inhibition of TrxR and GR an established microplate reader based assay was performed with minor modifications.<sup>12</sup> For this purpose commercially available rat liver TrxR and baker yeast GR (both from Sigma–Aldrich) were used and diluted with distilled water to achieve concentrations of 0.4 U/mL (TrxR) and 20.0 U/mL (GR). The compounds were freshly dissolved as stock solutions in DMF. To each 25  $\mu\text{L}$  aliquots of the enzyme solution each 25  $\mu\text{L}$  of potassium phosphate buffer pH 7.0 containing the compounds in graded concentrations or vehicle (DMF) without compounds (control probe) were added and the resulting solutions (final concentration of DMF: max. 0.5% V/V) were incubated with moderate shaking for 75 min at 37  $^{\circ}\text{C}$  in a 96 well plate. To each well 225  $\mu\text{L}$  of reaction mixture (1000  $\mu\text{L}$  reaction mixture consisted of 500  $\mu\text{L}$  potassium phosphate buffer pH 7.0, 80  $\mu\text{L}$  100 mM EDTA solution pH 7.5, 20  $\mu\text{L}$  bovine serum albumin solution 0.05%, 100  $\mu\text{L}$  of 20 mM NADPH solution and 300  $\mu\text{L}$  of distilled water) were added and the reaction was started by addition of 25  $\mu\text{L}$  of an 20 mM ethanolic dithio-bis-2-nitrobenzoic acid (DTNB) solution. After proper mixing, the formation of 5-thio-2-nitrobenzoic acid (5-TNB) was monitored with a microplate reader (Perkin Elmer

Victor X4) at 405 nm in 35 s intervals for 350 s. For each tested compound the non interference with the assay components was confirmed by a negative control experiment using an enzyme free solution. The  $IC_{50}$  values were calculated as the concentration of compound decreasing the enzymatic activity of the untreated control by 50% and are given as the means and error of 2–3 independent experiments.

### **Cell culture and Cytotoxicity Assay in MDA-MB-231 and HT-29 Cells**

The antiproliferative effects in MDA-MB-231 and HT-29 cells after 72 h (HT-29) or 96 h (MDA-MB-231) exposure to the gold complexes were evaluated using crystal violet assay.

In short: HT-29 human colon carcinoma cells and MDA-MB-231 breast cancer cells were maintained in cell culture medium (minimum essential eagle medium supplemented with 2.2 g  $NaHCO_3$ , 110 mg  $L^{-1}$  sodium pyruvate and 50 mg  $L^{-1}$  gentamicin sulfate adjusted to pH 7.4) containing 10% (V/V) fetal calf serum at 37 °C/5%  $CO_2$  and passaged once a week according to standard procedures. For the experiments, the compounds were prepared freshly as stock solutions in DMF and diluted with the cell culture medium to the final assay concentrations (0.1% V/V DMF). The assay procedure is described in more detail in our recent papers.<sup>10, 40</sup> The  $IC_{50}$  value was described as that concentration reducing proliferation of untreated control cells by 50%.

### **Cellular uptake studies**

#### **Sample preparation for cellular uptake studies**

The cellular uptake was measured according to a previously described procedure.<sup>32</sup> In short: cells were grown until at least 70% confluency in 75  $cm^2$  cell culture flasks. Stock solutions of the gold complexes in DMF were freshly prepared and diluted with cell culture medium to the desired concentration (final DMF concentration: 0.1% V/V, final gold complex concentration: 10  $\mu M$ ). The cell culture medium of the cell culture flasks was replaced with 4.0 mL of the cell culture medium solutions containing the compounds and the flasks were incubated at 37 °C/5%  $CO_2$  for 1, 4 or 8 h. The cell pellets were isolated by trypsinisation and centrifugation (room temperature, 3500 g, 5 min), resuspended in double distilled water, lysed by sonication and appropriately diluted using double distilled water. An aliquot was removed for the purpose of protein quantification by the Bradford method. The determination of the gold content of the samples was performed by high resolution continuum source atomic absorption spectroscopy (see below). Results were calculated from the data of 2–3 independent experiments and are given as nmol gold per mg cellular protein.

### **Atomic absorption spectroscopy (AAS)**

Gold contents were measured with a graphite furnace high resolution continuum source atomic absorption spectrometer (contraAA®700, Analytik Jena AG) at 242.795 nm according to a recently described method with minor modifications.<sup>32, 33</sup> In short: to 200  $\mu\text{L}$  aliquots of the diluted lysates 20  $\mu\text{L}$  Triton X-100 (1%) and 20  $\mu\text{L}$  ascorbic acid (1%) were added. The gold content of the samples was accessed using freshly prepared matrix-matched solutions of test compounds for calibration. Probes were injected at a volume of 25  $\mu\text{L}$  into standard graphite wall tubes. The mean absorbance of triplicate injections was used throughout the study. Drying, atomization and tube cleaning steps were performed as outlined in more detail in the literature.<sup>10</sup>

### **Concentration Dependent Effects on drug resistant Human Leukemia Cells.**

The following human cell line and its chemoresistant subline were used in this study: leukemic B-cell precursor (Nalm-6) and its vincristin (Nalm/VCR) resistant subline. All cell lines were cultured in RPMI 1640 supplemented with 10% fetal calf serum. Apoptotic cell death was determined by a modified cell cycle analysis, which detects DNA fragmentation at the single cell level. For measurement of DNA fragmentation cells were seeded at a density of  $1 \times 10^5$  cells/mL and treated with different concentrations of compound **2a** or **2d** (stock solutions were prepared in DMSO in these experiments). After 72 h of incubation, cells were collected by centrifugation at 300 g for 5 min, washed with PBS at 4 °C, and fixed in PBS/2% (v/v) formaldehyde on ice for 30 min. After fixation, cells were incubated with ethanol/PBS (2:1, v/v) for 15 min, pelleted, and resuspended in PBS containing 40  $\mu\text{g}/\text{mL}$  RNase A. After incubation for 30 min at 37 °C, cells were pelleted again and finally resuspended in PBS containing 50  $\mu\text{g}/\text{mL}$  propidium iodide. Nuclear DNA fragmentation was then quantified by flow cytometric determination of hypodiploid DNA. Data were collected and analyzed using FACScan (Becton Dickinson, Heidelberg, Germany) equipped with the CELLQuest software. Data are given in % hypoploidy (subG1), which reflects the number of apoptotic cells.

### **Acknowledgements**

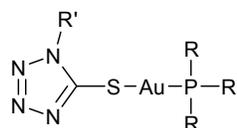
The postdoctoral fellowships for T.V.S by German Academic Exchange Service (DAAD) and Saint Petersburg State University (12.50.1560.2013) as well as the financial support by Dr. Kleist-Foundation (Berlin) are gratefully acknowledged. Mass-spectrometry was performed at the Center for chemical analysis and material research of St. Petersburg State University.

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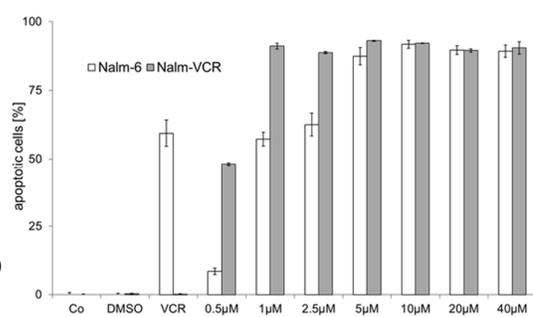
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## Graphical Abstract



- 2a** (R = Ph, R' = Me)  
**2b** (R = Ph, R' = Ph)  
**2c** (R = Ph, R' = (CH<sub>2</sub>)<sub>2</sub>NMe<sub>2</sub>)  
**2d** (R = 2-furyl, R' = Me)  
**2e** (R = 2-furyl, R' = Ph)



Gold(I) complexes with thiotetrazolate ligands are potent TrxR inhibitors with activity in drug resistant Nalm-6 cells