Anti-tumour active gold(I), palladium(II) and ruthenium(II) complexes with thio- and selenoureaato ligands: a comparative study†

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We report the results of a comparative study of the biological activity of a series of gold(I), palladium(II) and ruthenium(II) complexes containing deprotonated thio- and selenoureaato ligands. A library of compounds was prepared and characterised by spectroscopic methods and the solid-state structures of several derivatives were determined by single crystal X-ray diffraction. The in vitro activity of these compounds was evaluated in mammary and ovarian carcinoma, acute lymphatic and acute and chronic myeloid leukaemia cell lines. At lower concentrations Ru- and Pd-containing compounds displayed stronger anti-cancer effects than the gold compounds. In all cases, the selenium derivatives proved to be more active than the corresponding sulfur compounds.

Introduction

Acylselenoureas and acylthioureas of the type ArC(O)NHC(E)NR₂ (E = S, Se) have been known since the early 19th century1,2 and their metal complexes, particularly those with acylthiourea ligands, have been thoroughly investigated for a number of applications.3–6 For example, some acylthio- and acylselenoureas have been used for the extraction of gold from aqueous solutions.7 Gold(i) complexes, especially those with phosphine ligands, are known to display biological activity.8–14 A cytotoxicity study of a series of gold(i) triethylphosphine complexes [Au(X)PEt₃] (X = Cl, Br, CN, SCN, or thiolato ligand) revealed that the sulfur containing complexes exhibited a higher activity compared to those containing halide ligands.10 We have observed similar results in several gold(i) phosphate or N-heterocyclic carbene complexes containing heterocyclic sulfur ligands.15,16 There are several reports describing studies of acylthiourea palladium complexes focusing on their anti-microbial,17 fungicidal,18 thermal19 or mesomorphic20 properties. Palladium(II) complexes with phenantrone or its derivatives can intercalate with DNA thus exhibiting significant in vitro anti-tumour activity.8,14,21–23

The anticancer activity of palladium complexes has recently been subject of a detailed review.24 Apart from some homoleptic bis(selenoureaato) palladium(II) complexes, there are only very few examples of heteroleptic Pd(II) derivatives containing monoaionic acylselenoureas.6 Ruthenium compounds, in particular organometallic arene ruthenium(II) species, are emerging as a new class of promising anti-tumour active substances.8,14,23 Ruthenium complexes with selenoureaato ligands are unknown and only a few complexes with monoanionic thiourea to ligands have been reported, although none have been examined for their biological properties.33–38 Given our interest in the study of biologically active metal complexes with chalcogen ligands,15,16,39–44 we wished to carry out a comparative study of a family of compounds containing metal centres with known anti-cancer activity [Au(PTA), Pd(1,10-Phen) and Ru(2-cymol)] in combination with both acylthio- and acylselenoureaato ligands.

Results and discussion

Synthesis and crystal structures

The new selenourea N₂N-diethyl-N′-4-nitrobenzoylselenourea (denoted hereafter as HLSe) was prepared from KSeCN, 4-nitrobenzoyl chloride and Et₂NH in CH₂Cl₂ in the presence of PEG-400 as phase-transfer catalyst following a literature procedure.45 Orange crystals of the pure material were obtained in about 35% yield after recrystallisation from EtOH. Alternatively, the compound can also be obtained in similar yield using the original method of Douglass.1 The known
thiourea \(N,N\)-diethyl-\(N\)′-4-nitrobenzoylthiourea (denoted hereafter as \(\text{HL}^S\)) was prepared analogously from KSCN, 4-nitrobenzoyl chloride and Et\(_2\)NH in acetone. The identity and purity of \(\text{HL}^{Se}\) was confirmed by \(^1\)H, \(^{13}\)C and \(^{77}\)Se NMR spectroscopy as well as by single crystal X-ray diffraction. The molecular structure of the compound is shown in Fig. 1. Overall, the bond distances and angles of \(\text{HL}^{Se}\) (Table S1†) are similar to those reported for PhC(O)NHC(Se)NEt\(_2\) with some minor differences, which are detailed in the following section. The C=O and C≡Se units in \(\text{HL}^{Se}\) lie almost perpendicular to each other (dihedral angle Se–C–N–C = 90.2°), whilst in the Ph-derivative this angle is approximately 108°. Furthermore, the Ph-derivative forms dimers through intermolecular H-bonds between the NH group and the selenium atom in the solid-state, whereas \(\text{HL}^{Se}\) forms an infinite chain through N–H⋯O hydrogen bonds [N(H)⋯O = 2.905(3) Å]. In \(\text{HL}^S\) similar features can be observed: infinite chain formation through N–H⋯O hydrogen bonds [N(H)⋯O = 2.915(2) Å] and perpendicular C=O and C≡S units (dihedral angle S–C–N–C = 91.1°).

The seleno- or thioureas \(\text{HL}^{Se}\) and \(\text{HL}^S\) react with [AuCl\(_2\)(PTA)] (PTA: 1,3,5-triaza-7-phosphaadamantane) in the presence of base to give neutral gold(I) PTA complexes containing the deprotonated seleno- or thioureato ligands (Scheme 1). Similarly, the reaction of \(\text{HL}^{Se}\) and \(\text{HL}^S\) with \(\text{cis-[PdCl}_2(1,10\text{-Phen})]\) in the presence of base affords the cationic Pd(II) species \(\text{cis-[Pd}[4-O_2NC_6H_4C(O)NC(\text{E})NEt_2](1,10\text{-Phen})]\)\(^+\) (E = S, Se), which were isolated as their PF\(_6\) salts as shown in Scheme 1. The organoruthenium(II) complexes \([\text{Ru}[4-O_2NC_6H_4C(O)NC(\text{E})NEt_2)(p\text{-cymol})(PPh_3)]\)PF\(_6\) (E = S, Se) were prepared from the reaction of [RuCl\(_2(p\text{-cymol})\)] with the corresponding ligand in the presence of Ph\(_3\)P, base and NH\(_4\)PF\(_6\) (Scheme 1).

The compounds were characterised by \(^1\)H, \(^{13}\)C, \(^{77}\)Se and \(^{31}\)P NMR spectroscopy, electrospray mass spectrometry and several derivatives also by X-ray crystallography. In the gold(i) complexes the monoanionic thio- or selenourea ligands coordinate only via the sulfur or selenium atom to the gold(i) centre. In case of the ruthenium and palladium species, bidentate coordination of both chalcogen atoms of the thio- and selenourea ligands is observed. In all the compounds reported here, the thio- or selenourea ligands are deprotonated, which is evident from the disappearance of the signal for the N–H proton in the \(^1\)H NMR spectra of the compounds. For the gold derivatives, a shift of the position of the singlet due to the PTA ligand in the \(^{31}\)P NMR spectra is observed upon coordination of the seleno- or thioureato ligands. This resonance is observed at −52.0 ppm in [AuCl\(_2\)(PTA)] and at −51.7 and −49.8 ppm in the thio- and selenourea derivatives, respectively. In addition, the \(^{13}\)C NMR signals of the quaternary carbon atom bound to the sulfur or selenium atom are shifted to 176.8 (S) or 167.1 (Se) ppm to higher field upon coordination. Whilst the \(^{77}\)Se NMR signal for the unbound selenourea is found at 502 ppm, that of the gold selenourea is shifted to 492 ppm (Scheme 1).

![Fig. 1](image-url) Molecular structure of \(N,N\)-diethyl-\(N\)′-4-nitrobenzoylselenourea \(\text{HL}^{Se}\). Ellipsoids show 30% probability levels. Hydrogen atoms have been omitted for clarity.

![Scheme 1](image-url)
The palladium complexes (Fig. 3) show square-planar coordination with one bidentate chalcogenoureato and a 1,10-phenanthroline ligand bound to the metal. In both the ruthenium and the palladium compounds, the chalcogenoureato ligands are nearly planar due to electron delocalisation in the ligand backbone. A comparison of the two C–N distances as well as the C–O and C–Se bond lengths of the free and coordinated ligand, shows that upon coordination to the metal atom the NH–C=N–Se moiety is transformed into an N=C=Se moiety (the selenolato tautomer). The C–Se bond distance increases from 1.82 Å to 1.88 Å (Pd) or 1.90 Å (Ru) which is in agreement with literature data for ruthenium-µ-phenylselenido compounds. Due to the slightly smaller radius of the sulfur atom, the analogous bond distances in the sulfur complexes are correspondingly shorter. The C–S bond distance increases from 1.66 Å (HL) to 1.73 Å in the complex. In contrast, the C=O and C(O)N bond distances remain virtually unchanged upon coordination when compared with those of the free seleno- or thiourea, respectively. A table listing important
bond distances and angles of the metal complexes and the thio- and selenourea ligands is included in the ESI.†

**Biological studies**

The biological activity of HLS, HLSe as well as selected Au(i), Pd(ii) and Ru(ii) complexes including their chlorido-precursors (Chart 1) and cisplatin as further control was determined by their effect on proliferation, metabolic activity, programmed cell death and induction of oxidative stress of various tumor and leukemia cell lines.

All Au(i), Pd(ii) and Ru(ii) compounds dose-dependently reduced the proliferation of mammary and ovarian carcinoma, acute lymphatic as well as acute and chronic myeloid leukemia cell lines (ESI Fig. S1A–S1F†), whereby, in general, the selenium derivatives induced stronger effects than the sulfur compounds. For example, a concentration of 5 µM Pd–S significantly reduced the proliferation of SUP-B15 cells from 100% (without compound; data not shown) to 44.5 ± 8.4% and a concentration of 10 µM to 18.7 ± 3.1% (p < 0.05; Fig. 4) so that the IC50 was calculated as 4.9 ± 1.2 µM (Table 1). Interestingly, in the mammary carcinoma as well as the myeloid leukemia cell lines Pd–S was almost ineffective. In contrast, Pd–Se induced strong anti-proliferative effects and both Ru(ii) compounds completely abolished the proliferation in all cell lines tested. Weaker anti-proliferative activity (again in all cell lines) was observed for the two Au(i) complexes. Cisplatin dose-dependently reduced the proliferation of the tumor cell lines, but did not show any activity on the leukemia cell lines. Whereas [RuCl2(p-cymol)]2 was completely ineffective in all cell lines, [AuCl(PTA)] and the thio- and selenoureas HLSe and HLS induced anti-proliferative effects at higher concentrations (25 µM and 50 µM, ESI Fig. S1A–S1F†).

![Chart 1](image)

**Fig. 4** Proliferation of SUP-B15 cells after a three-day incubation with the Au(i), Ru(ii), Pd(ii) complexes, HLS, HLSe, the chlorido complexes [AuCl(PTA)], [RuCl2(p-cymol)]2, cis-[PdCl2(L,10-Phen)] as well as cisplatin. Proliferation in the absence of the compounds was set to 100%. Mean proliferation and standard error of four experiments is shown. Statistical significance was determined with the Wilcoxon Rank Sum Test (*p < 0.05 against the untreated control).

**Table 1** IC50 values of the metal complexes in SUP-B15 cells

<table>
<thead>
<tr>
<th>Compound</th>
<th>IC50 (µM)</th>
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<tbody>
<tr>
<td></td>
<td>Proliferation</td>
</tr>
<tr>
<td>Au–S</td>
<td>17.9 ± 3.2</td>
</tr>
<tr>
<td>Au–Se</td>
<td>9.1 ± 0.5</td>
</tr>
<tr>
<td>Ru–S</td>
<td>3.5 ± 0.2</td>
</tr>
<tr>
<td>Ru–Se</td>
<td>2.6 ± 0.5</td>
</tr>
<tr>
<td>Pd–S</td>
<td>4.9 ± 1.2</td>
</tr>
<tr>
<td>Pd–Se</td>
<td>3.7 ± 0.4</td>
</tr>
</tbody>
</table>

IC50 values are calculated as the concentration at which the compounds induce 50% inhibition of proliferation or metabolic activity of SUP-B15 cells in comparison to untreated cells. The IC50 values are presented as means ± standard error of four experiments.
[PdCl2(1,10-Phen)] dose-dependently reduced the proliferation of all cell lines almost as effective as Pd-Se which may be explained by the ability of cis-[PdCl2(1,10-Phen)] to intercalate with DNA.

The anti-proliferative activity of the substances was also examined on A-2780 cells resistant to cisplatin and revealed that the Au(i), Pd(II) and Ru(II) compounds have the potential to overcome cisplatin resistance (Table 2).

To evaluate if the reduced cell growth was also accompanied by cell death, the metabolic activity of the cell lines after compound incubation was analysed. Metabolic activity is determined by the reduction of tetrazolium salts into formazan derivatives. Since this process requires functional mitochondria, the assay discriminates living and dead cells whose mitochondria are deactivated within a few minutes after cell death. As shown in Fig. 5 and ESI Fig. S2A–S2F,† the metabolic activity of the cell lines was also dose-dependently diminished after 72 hours of incubation with the test compounds and cis-[PdCl2(1,10-Phen)]. Again, the Ru(II) compounds and Pd-Se were the most active compounds with an IC50 of ∼4 µM (Table 1), whereas the chlorido precursors [AuCl(PTA)] and [RuCl2(p-cymol)], HL5, HLSe as well as cisplatin were almost ineffective.

To confirm the cytotoxicity of the compounds, their capacity to induce programmed cell-death was determined by ssDNA apoptosis ELISA, which is based on the selective denaturation of DNA in apoptotic cells by formamide and further detection of denatured DNA with a monoclonal antibody to single-stranded DNA (ssDNA). Compounds which reduced proliferation and metabolic activity also induced apoptosis of the leukemia and tumor cells, whereby the sensitivity of the cell lines to programmed cell-death induction varied (ESI Fig. S3A–S3E†). For example, apoptosis of the cell line SUP-B15 was 3.7 ± 0.8 times higher after addition of 5 µM Pd-Se and 5.2 ± 0.6 times higher when 10 µM Pd-Se was added to the cultures in comparison to cells without compound incubation (Fig. 6).

A similar pro-apoptotic effect was observed for both Ru(II) compounds, whereas 10 µM Au-Se and 5 µM as well as 10 µM Pd-Se caused a lower, but significant apoptosis rate. Consistent with the weak anti-proliferative and anti-metabolic activity, incubation with Au-S did not lead to programmed cell death in any of the cell lines. [AuCl(PTA)], [RuCl2(p-cymol)], and cisplatin were also ineffective in inducing apoptosis. Interestingly, despite no activity on proliferation and metabolism, HL5 and HLSe significantly induced apoptosis in all leukemic cell lines, but not in the carcinoma cell lines (Fig. 6 and ESI Fig. S3A–S3E†). A similar effect was observed for cis-[PdCl2(1,10-Phen)], which decreased proliferation and metabolic activity of the cell lines, however, without inducing their cell death. These rather contradictory results are currently under investigation.

The pro-apoptotic effect of Ru-Se (Fig. 7A) and the lacking effect of [RuCl2(p-cymol)]2 (Fig. 7B) can be readily detected by morphological changes of the cisplatin-resistant A-2780 cells.

One mechanism involved in the induction of apoptosis is oxidative stress triggered by the formation of reactive oxygen...
species (ROS) due to deregulated mitochondrial activity. It has been shown for a variety of gold,50–52 ruthenium31,32,53–55 and palladium56 complexes that their anti-tumour activity is mediated by an enhanced ROS production. Therefore, ROS levels were measured after a two-hour incubation of SUP-B15 cells with 10 µM of the respective compound. As shown in Fig. 8, both Ru(II) compounds and Pd–Se significantly induced oxidative stress in this cell line, in agreement with an increase of programmed cell death. Interestingly, the pro-apoptotic effect of HLS and HLSe cannot be explained by an increase in ROS formation. Neither the chlorido compounds nor cisplatin did induce oxidative stress of SUP-B15 cells.

Conclusions

In summary, gold(I), palladium(II) and ruthenium(II) compounds containing anionic nitro-substituted acylseleno- or acylthioureas display strong anti-tumor and anti-leukemia effects. The ligands HLS, HLSe as well as the chlorido complexes [AuCl(PTA)] and [RuCl2(p-cymol)]2 induced only weak effects or only at the highest concentrations tested, whereas cis-[PdCl2(1,10-Phen)] reduced the proliferation and the metabolic activity almost as effective as the test compounds. The mechanisms of action include inhibition of proliferation and metabolic activity as well as induction of apoptosis and oxidative stress by ROS formation. The strongest effects were observed for the Ru(II) compounds followed by the Pd(II) complexes whereby the selenium-containing compounds displayed stronger biological activity than the sulfur derivatives. The Au(I) complexes were only effective at higher concentrations.

Materials and methods

Chemistry

Unless otherwise stated, all manipulations were carried out without taking precautions to exclude air and moisture. All chemicals and solvents (HPLC quality) were sourced commercially and used as received. [AuCl(PTA)] was prepared from the reaction of [AuCl(tht)]57 with the appropriate amount of the phosphine. The other metal precursors cis-[PdCl2(1,10-phen)]58 [Ru(p-cymol)Cl]259 as well as N,N-diethyl-N’-4-nitrobenzoylthiourea (HLs)60 were prepared by standard procedures described in the literature. The purity of all compounds was confirmed to be >95% by elemental analysis or high-resolution mass spectrometry.

Instrumentation

NMR spectroscopy. 1H, 13C, 77Se and 31P{1H} NMR spectra were recorded on 400 Bruker Avance or 600 MHz Bruker Avance II spectrometers. Chemical shifts are quoted relative to external SiMe4 (1H, 13C), Me2Se or KSeCN (77Se) and 85% H3PO4 (31P).

IR spectroscopy. IR spectra were run as KBr pellets on a Bruker Tensor 27 instrument.
Elemental analyses. Elemental analyses were performed by staff of the in-house microanalytical laboratory using an Elementar Vario EL analyser.

Mass spectrometry. High-resolution electrospray mass spectra were recorded on Bruker MicroTOF spectrometer in positive ion mode using solutions of the samples in MeCN.

X-ray crystallography. Diffraction data were collected at 150 K using a Rigaku Oxford Diffraction Gemini E Ultra diffractometer, equipped with an EOS CCD area detector and a four-circle kappa goniometer. For the data collection the Mo Kα source emitting graphite-monochromated Mo-Kα radiation (λ = 0.71073 Å) was used. Data integration, scaling and empirical absorption correction was carried out using the Crysalis Pro program package. The structures were solved using Direct Methods or Patterson Methods and refined by Full-Matrix-Least-Squares against F². The non-hydrogen atoms were refined anisotropically allover and hydrogen atoms were placed at idealized positions and refined using the riding model. All calculations were carried out using the program Olex2. Full crystallographic and refinement details as well as tables with bond lengths and angles are included in the ESI.†

Syntheses

**N,N-Diethyl-N′-4-nitrobenzoylselenourea (HLSe).** To a suspension of KSeCN (2.88 g, 20 mmol) and PEG-400 (0.28 mL) in CH₂Cl₂ (10 mL) was added a solution of 4-nitrobenzoyl chloride (3.71 g, 20 mmol) in CH₂Cl₂ (20 mL). After ca. 15 min at room temperature Et₂NH (2.07 mL, 20 mmol) was added, and the resulting mixture was stirred for 1 h protected from light. The red-orange mixture was filtered and the filtrate sub-sequently evaporated to dryness. The resulting mixture was stirred for 20 min at room temperature, then [AuCl₂(C₆H₄CO)NC(S)NEt₂]PF₆ was added, and the orange suspension was cooled to 0°C under argon protection. Addition of diethyl ether afforded a yellow solid, which was isolated by filtration and dried in air. Yield: 0.150 g (86%).¹ H NMR (400 MHz, CDCl₃): δ = 1.28 (br. s, 6H, NCH₂CH₃), 3.66 (br. s, 2H, NCH₂CH₃), 3.81 (br. s, 2H, NCH₂CH₃), 4.02 (q, J_H-H = 13.2 Hz, 2H, PCH₃), 8.21 (d, J = 8.9 Hz, 2H, H-3), 8.28 (d, J = 8.9 Hz, 2H, H-2) ppm. ¹³C{¹H} NMR (101 MHz, CDCl₃): δ = 13.0 (NCH₂CH₃), 13.4 (NCH₂CH₃), 46.0 (NCH₂CH₃), 48.4 (NCH₂CH₃), 52.3 (d, J_p-c = 18.7 Hz, PCH₃N), 72.9 (d, J_p-c = 7.7 Hz, NCH₂N), 123.1 (C-2), 130.2 (C-3), 143.68 (C-4), 149.2 (C-1), 163.1 (C-O), 167.1 (C-Se) ppm. ⁷⁷Se NMR (76 MHz, CDCl₃, Me$_2$Se): 244 ppm. IR (KBr disk) ν: 1342 (s, NO₂), 1520 (s, NO₂), 1567 (s, C=O), 2932 (m, Csp³–H) cm⁻¹. Elemental analysis calc (%) for C₁₂H₁₅N₃O₃Se: 31.64, H 4.16, N 11.83.

Preparation of the palladium(II) complexes

cis-[Pd(4-NO₂C₆H₄CO)(NC(S)NEt₂)]PF₆. A mixture containing [PdCl₂(1,10-Phen)] (50.0 mg, 0.14 mmol), NaOMe (12.6 mg, 0.25 mmol) and [AuCl₂(PTA)] (100 mg, 0.26 mmol) was added to a saturated DMSO solution of the complex. The mixture was heated to reflux for 30 min. The orange suspension was cooled to room temperature and concentrated in vacuum. The resulting solid was filtered and washed with water and diethyl ether to give an orange solid. Yield: 0.098 g (93%).¹ H NMR (400 MHz, DMSO-d₆, 32 °C): δ = 1.28 (t, J = 7.1 Hz, 3H, NCH₂CH₃), 1.42 (t, J = 7.2 Hz, 3H, NCH₂CH₃), 3.94 (q, J = 7.1 Hz, 2H, NCH₂CH₃), 3.97 (q, J = 7.2 Hz, 2H, NCH₂CH₃), 7.98 (dd, J = 8.1, 5.3 Hz, 1H, H₃– or H₈-ph), 8.20 (dd, J = 7.9 Hz, 1H, H₉– or H₈-ph), 8.30 (m, 4H, H₂–H₂), 8.62 (dd, J = 5.3, 0.9 Hz, 1H, H₂– or H₉-ph), 8.91 (dd, J = 8.1, 0.6 Hz, 1H, H₄– or H₇-ph), 8.97 (m, 2H, H₄– or H₇-ph) ppm. ¹³C{¹H} NMR (243 MHz, DMSO-d₆, 25 °C): δ = 114.2 (sept, J = 711.2 Hz, PFF₆) ppm. HR-ESI-MS: m/z [obs., calc.] = 613.9920, 613.9923 [M⁺]. X-ray quality crystals were obtained by slow diffusion of diethyl ether into a saturated DMSO solution of the complex.

cis-[Pd(4-NO₂C₆H₄CO)(NC(S)NEt₂)]PF₆. This was prepared as above from [PdCl₂(1,10-Phen)] (50.0 mg, MeCN).
0.14 mmol), HL^5 (39.3 mg, 0.14 mmol), NaOAc (12.6 mg, 0.15 mmol) and NH_4PF_6 (25.1 mg, 0.15 mmol). The product was obtained as an orange solid in 79% yield (0.078 g). \(^1\)H NMR (600 MHz, DMSO-d_6): \(\delta = 1.28\) (t, \(J = 7.1\) Hz, 3H, NCH_2CH_3), 1.41 (t, \(J = 7.1\) Hz, 3H, NCH_2CH_3), 3.92 (q, \(J = 7.1\) Hz, 2H, NCH_2CH_3), 3.97 (q, \(J = 7.1\) Hz, 2H, NCH_2CH_3), 8.06 (dd, \(J = 8.1, 5.3\) Hz, 1H, H3- or H4-phen), 8.22 (dd, \(J = 8.8\) Hz, 1H, H5- or H6-phen), 8.25 (dd, \(J = 8.2, 5.1\) Hz, 1H, H8- or H9-phen), 8.27 (dd, \(J = 8.8\) Hz, 1H, H5- or H6-phen), 8.30 (d, \(J = 9.0\) Hz, 2H, H-2), 8.33 (d, \(J = 9.0\) Hz, 2H, H-3), 8.66 (dd, \(J = 5.3, 1.0\) Hz, 1H, H4- or H7-phen), 8.99 (dd, \(J = 8.2, 1.0\) Hz, 1H, H4- or H7-phen), 9.02 (dd, \(J = 5.1, 1.1\) Hz, 1H, H2- or H9-phen) ppm. \(^31\)P{\(^1\)H} NMR (243 MHz, DMSO-d_6): \(\delta = -144.2\) (sept, \(J = 711.2\) Hz, PF_6) ppm. HR-ESI-MS: m/z (obs., calc.) = 566.0474, 566.0478 [M^+]. X-ray quality crystals were obtained by slow diffusion of diethyl ether into a saturated acetone solution of the complex.

Preparation of the ruthenium complexes

\[\text{[Ru(p-cym)][4-NO_2C_6H_4C_6O(NC_5Se)(NPh)]PF}_6\] A mixture containing [Ru(p-cym)]Cl_2 (50 mg, 0.082 mmol), HL^5 (53.6 mg, 0.163 mmol), Et_3N (23 \(\mu\)l, 0.165 mmol), Ph_3P (43 mg, 0.164 mmol) and NH_4PF_6 (29 mg, 0.179 mmol) in methanol (15 ml) was heated to reflux for 15 min. Addition of water to the red solution gave an orange precipitate. This was isolated by filtration and was washed with water and diethyl ether and subsequently dried in air. Yield: 0.132 g (84%). \(^1\)H NMR (400 MHz, CDCl_3): \(\delta = 1.11\) (d, \(J = 6.9\) Hz, 3H, CH(CH_3)_2), 1.20 (d, \(J = 6.9\) Hz, 3H, CH(CH_3)_2), 1.28 (t, \(J = 7.1\) Hz, 3H, NCH_2CH_3), 1.40 (t, \(J = 7.2\) Hz, 3H, NCH_2CH_3), 1.91 (s, 3H, CCH_3), 2.54 (sept, \(J = 6.9\) Hz, 1H, CH(CH_3)_2), 3.67 (dq, \(J = 13.2, 7.1\) Hz, 1H, 1NHCH_2CH_3), 3.92 (m, 2H, NCH_2CMe), 4.35 (dq, \(J = 14.2, 7.1\) Hz, 1H, 1NHCH_2CH_3), 5.17 (dd, \(J = 6.1, 1.0\) Hz, 1H, H-3'cy), 5.65 (m, 2H, H-2cy, H-3'cy), 5.72 (dd, \(J = 6.2, 1.0\) Hz, 1H, H-2'cy), 7.33-7.46 (m, 15H, PPh_3), 7.61 (d, \(J = 9.0\) Hz, 2H, H-2), 8.03 (d, \(J = 8.9\) Hz, 2H, H-3') ppm. \(^31\)P{\(^1\)H} NMR (162 MHz, CDCl_3): 39.5 (PPh_3), \(-144.2\) (sept, \(J_p^c = 716.1\) Hz, PF_6) ppm. \(^13\)C{\(^1\)H} NMR (101 MHz, CDCl_3): \(\delta = 13.0\) (NCH_2CH_3), 13.4 (NCH_2CH_3), 18.0 (CCH_3), 22.0 (CH(CH_3)_2), 22.2 (CH(CH_3)_2), 31.2 (CH), 46.4 (NCH_2CH_3), 49.5 (NCH_2CH_3), 89.8 (d, \(J_p^c = 3.2\) Hz, C-3cy), 90.0 (d, \(J_p^c = 7.1\) Hz, C-2cy, C-90y), 90.9 (C-9'cy), 92.5 (d, \(J_p^c = 1.5\) Hz, C-2'cy) ppm. Biological activity

Compounds. All compounds were dissolved in dmsol, aliquoted and stored at \(-20^\circ\)C. Prior to use solutions of the respective compound were diluted at least 1:1000 with cell culture medium. Dmsol in the appropriate concentration did not induce any effect on the cell lines.

Cell lines. The leukemic cell lines were purchased from DSMZ – German Collection of Microorganisms and Cell Cultures, Braunschweig, Germany. The mammary carcinoma cell lines were kindly provided by the Institute of Pharmacy, Free University of Berlin. The ovarian carcinoma cell lines were kindly provided by the Department of Gynecology, Medical University Innsbruck. The acute myeloid leukemia cell line HL-60 as well as the chronic myeloid leukemia cell line K-562, the mammary carcinoma cell lines MCF-7 and MDA-MB231 as well as the ovarian carcinoma cell lines A2780 (cisplatin-sensitive and cisplatin-resistant) were grown in RPMI 1640 without phenol red (PAA Laboratories, Pasching, Austria), supplemented with glutamine (2 mM), penicillin (100 U ml^-1), streptomycin (100 \(\mu\)g ml^-1) and fetal bovine serum (FBS; 10%; all from Invitrogen Corporation, Gibco, Paisley, Scotland) at 37 °C in a 5% CO_2/95% air atmosphere and fed twice weekly. Cisplatin (1 \(\mu\)M) was added to the cisplatin-resistant A-2780 cell line every third passage. The acute lymphatic leukemia cell line SUP-B15 was grown in Mc Coy's 5A medium (Invitrogen) and supplemented with 20% FBS.

Analysis of metabolic activity and proliferation. Logarithmically growing cells were resuspended in culture medium at 1 \(\times\) 10^6 cells per ml and plated in triplicate in U-bottomed or flat-bottomed microtiter plates (50 \(\mu\)l; Falcon, Becton Dickinson, Franklin Lakes, NJ, USA). Various concent-
trations of compounds were added one hour thereafter. After a three-day culture at 37 °C in a 5% CO2/95% air atmosphere cultures were analyzed for metabolic activity using a modified 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay (EZ4U kit; Biomedica, Vienna, Austria) according to the manufacturer’s instructions and for cellular proliferation using [methyl-3H]-thymidine uptake. Metabolic activity and proliferation in the absence of the compound was set to 100%.

Analysis of apoptosis. The plates for apoptosis were prepared analogously as described above and were evaluated with an enzyme-linked immunosorbent assay (ELISA) using the ssDNA apoptosis kit (Chemicon International, Hofheim, Germany) according to the manufacturer’s instructions. Apoptosis rate was evaluated by dividing the optical density of the wells measured in the plates for ssDNA apoptosis by the optical density of the wells for metabolic activity. Apoptosis in the absence of the compounds was set to 1.

Detection of morphological changes. 3 × 10^5 cells were adjusted to 1 × 10^5 cells per ml, placed in a 12-well plate and incubated at 37 °C in a humidified 5% CO2/95% air atmosphere for 24 hours to let the cells adhere. After 24 hours 10 µM compound was added and the cells were cultivated for another 24 hours. Every 30 minutes a picture was made by JuLi™ Live cell imaging system (NanoEnTek, Seoul, Korea).

Measurement of intracellular ROS formation. SUP-B15 cells (1 × 10^5 cells per well) were incubated with 10 µM of the respective compound for two hours at 37 °C in a humidified 5% CO2/95% air atmosphere, washed and treated for further 15 minutes with 5 µL of a 5 mM dihydroethidium solution (Sigma, St Louis, MO, USA) in 1 ml phosphate-buffered saline (PBS) + 1% bovine albumin (Ortho-Clinical Diagnostics, High Wycombe, UK). After another wash cells were resuspended in 500 µL PBS + 1% bovine albumin and analyzed immediately by flow cytometry (FACSCalibur, Becton Dickinson, San José, CA, USA).

Statistical analysis. The Wilcoxon Rank Sum Test was used to analyze the differences between proliferation, metabolic activity, apoptosis and ROS production in the absence and the presence of a variable concentration of the test compounds (NCSS software, Kaysville, UT, USA).

**Conflicts of interest**

There are no conflicts to declare.

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**References**
