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Au(III) compounds as HIV nucleocapsid protein (NCp7)-nucleic acid antagonists

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Abstract

The HIV nucleocapsid NCp7-SL2 RNA interaction is interrupted in the presence of a formally substitution-inert gold(dien)-nucleobase/N-heterocycle AuN₄ compound where the N-heterocycle serves the dual purposes of a template for “non-covalent” molecular recognition of the essential tryptophan of the protein, mimicking the natural reaction and subsequent “fixation” by Au-Cys bond formation providing a chemotype for a new distinct class of nucleocapsid-nucleic acid antagonist.

In this communication we show that the metallated nucleobase [Au(dien)(9-EtGua)]³⁺ (**I**, 9-EtG = 9-ethylguanine) inhibits the nucleocapsid protein HIVNCp7 zinc finger-RNA interaction in an antagonist fashion and expands the chemistry of this important HIV target in hitherto unrecognized directions. HIVNCp7 (NC) is a small basic protein containing two zinc finger (or zinc knuckle) CysCysHisCys (CCHC) motifs [1,2]. This motif is highly conserved in all known retroviruses, and mutation of the zinc-chelating residues results in noninfectious viruses. NC is critically involved in both the early and late steps of the HIV-1 cycle, mainly through its ability to chaperone nucleic acids toward their most stable conformation [1,2]. The rearrangement of nucleic acids is essential for many viral replication processes including reverse transcription and recombination. Two general approaches to NCp7 inactivation are (i) electrophilic attack on the cysteinate residues of the zinc fingers and (ii) zinc chelation, resulting in both cases in loss of tertiary structure [2,3].

The aromatic amino acids tryptophan (Trp, W) and phenylalanine (Phe) are critical for the NC-nucleic acid molecular recognition. The mutation of even one of these residues significantly decreases NC's nucleic acid chaperone activity, and correlates with inhibition of viral replication [4]. Nucleic acids bind exclusively to the nucleocapsid domain fixing the orientation of the two Zn knuckles relative to one another [5]. The NMR-based structure of the HIV-1 RNA packaging signal confirmed the importance of exposed or weakly paired

Incubation of $[\text{Au}(\text{dien})(9\text{-EtGua})]^{3+}$ (**I**) with intact NC resulted in the detection of species produced by ejection of both Zn^{2+} ions and incorporation of up to 3 Au ions – AuF , Au_2F and Au_3F , Figure 2. This is the first demonstration of a coordination compound reacting on a “multiple (>1)” zinc finger peptide. The multiple Au_n species are observed due to the number of cysteines and histidines capable of Au binding. The results are consistent with biophysical studies of Au(I) and Au(III) compounds on zinc fingers showing rapid displacement of the central Zn(II) with all ligands attached to the gold center being displaced [20–25].

The CD spectrum of NC is characterized by a positive maximum observed at ~215 nm. The reaction with the Au compound caused a time-dependent decrease in intensity of this band and a significant increase in the negative ellipticity with a slight blue shift of the 195–200 nm band, which are indicative of conformational changes from ordered structure to random coil, Figure 3 [26,27]. The pronounced changes are consistent with a loss of structural integrity in NC upon treatment with **I**. The combined MS and CD data confirm results using the C-terminal finger of NC (34–52 F2) and ejection of Zn^{2+} [18]. Given the demonstrated strong association of **I** with N-AcTrp [18], the combined results are consistent with molecular recognition of the NC through the predicted stacking interaction with the subsequent rapid loss of ligands upon covalent reaction with peptide.

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complex for 30 min, or by adding **I** to the preformed NC-SL2 complex. In the former case, the use of the reactive Au(III) species resulted only in observation of free unbound SL2, thus indicating that the major structural changes induced by Zn^{2+} - Au^{3+} replacement abrogated the binding capabilities of NC (Figure S1). In the latter case, a direct comparison of data obtained from NC-SL2 in the presence/absence of **I** showed a significant increase of free SL2 in solution (Figure 4 and Figure S2), thus suggesting that the gold compound is capable of inducing dissociation of the peptide from its cognate RNA. In addition, minor signals corresponding to [(Au,ZnNC)-SL2] and [(Au₂NC)-SL2] were detected, which are consistent with direct displacement of Zn^{2+} from the intact NC-SL2 complex (Figure S3).

Electrophoretic mobility shift assays confirm the general trend of these results. In Figure 5A, the control experiment shows that an increasing amount of NC protein incubated with ³²P-end-labelled SL2 RNA results in a concentration-dependent increase in NC-SL2 complex formation and a decrease in free SL2. To determine the effect of **I** on NC-SL2 binding, NC was incubated with increasing concentrations of compound prior to addition of SL2. Results show a concentration-dependent decrease in NC-SL2 complex formation suggesting that [Au(dien)(9-EtGua)]³⁺ effectively inhibits SL2 binding to NC, Figure 5B. An expected increase in unbound SL2 is not observed, however. Upon further investigation, drug-SL2 complexes could be detected. This species is shown in control experiments as a slow migrating band near the top of the gel, Figure S4. The generality of these results was corroborated using fluorescence polarization, Figure S5. Fluorescence polarization control experiments showed that SL2 was displaced from the high affinity fluorescein-labelled SL2-NCp7 complex in presence of [Au(dien)(9-EtGua)]³⁺. In this case the analog [Au(dien)(DMAP)]³⁺, **II**, which displays intrinsically higher π - π stacking with tryptophan, [18], showed similar behavior. The extrapolated IC₅₀ values were 22.0 and 29.0 μM for the 9-EtGua and DMAP compounds respectively.

In summary, the overall results are consistent with the ability of [Au(dien)(9-EtGua)]ⁿ⁺ and congeners to act as antagonist of the NC-SL2 interaction, Figure 6. Both components of the NC-nucleic acid chaperone activity have been targeted [30–35]. A study of approximately 2,000 small molecules from the NCI Diversity Set showed a good correlation between tryptophan quenching and inhibition of NC-nucleic acid binding [31]. Of five selected hits, from a total of 4800 compounds screened for inhibition of NC-mediated destabilization of the stem-loop structure of cTAR DNA (a sequence complementary to the transactivation response element) 4 of the 5 correlated with their ability to compete with the nucleic acid for binding to NC [32].

Development of NC-nucleic acid antagonists is an attractive strategy for AIDS treatment complementary to HAART. A small number of compounds such as S-acyl 2-mercaptobenzamide thioesters (SAMTs), [36,37], have undergone clinical trials as putative NC-interacting drugs [reviewed in 2,38]. Exploratory cellular studies on inhibition of infectivity of HIV-1 strains in peripheral mononuclear blood cells (PMBC) showed that both Au compounds were moderately effective inhibitors in the 20–50 μM range, Table 1. The Au(I) compound Auranofin was also compared as control given its demonstrated efficacy in restricting the viral reservoir in the monkey AIDS model and inducing containment of viral load following Anti-Retroviral Therapy (ART) suspension [39]. Auranofin was very

effective at submicromolar concentrations but was also quite cytotoxic. The origin of auranofin's activity is likely to lie in an immunological response but nevertheless the fact that the Au(III)N₄ compounds are significantly less cytotoxic (IC₅₀ > 100 μM) than auranofin suggest the possibility of enhancement of viral selectivity with these Au(III) compounds. A series of Au(I) and Au(III) compounds has been tested for inhibition of HIV activity based on reverse transcriptase and protease and viral entry as targets [40]. The results presented here suggest that the NCp7 nucleocapsid protein is also a valid rational target for optimization.

Zinc fingers in general are attractive targets for drug intervention by coordination compounds [41]. Au(I) and Au(III) compounds have also been shown to inhibit the zinc-finger PARP-1 (polyADP-ribose polymerase) action [23,24,42,43]. The question remains how to infer selectivity for specific zinc finger intervention, let alone zinc enzymes in general. The concept of "weak electrophiles" is an attractive proposition to impart selectivity toward the highly nucleophilic cysteines of zinc finger coordination spheres [44,45]. We have proposed that the formally substitution-inert MN₄ chemotype is the coordination chemistry Lewis acid equivalent of a weak "organic" electrophile, significantly less reactive than that of a MClN₃ unit, containing the more substitution-labile M-Cl bond [46]. With respect to NC inhibition, and considering the most general MN₄ structure as [M(chelate)(Nucleobase/heterocycle)]ⁿ⁺ the purine nucleobase or heterocycle serves the dual purposes of a template for "non-covalent" molecular recognition mimicking the natural reaction and subsequent "fixation" by purine/heterocycle substitution with thiolate or zinc-thiolate, Figure 6. The results reported here complement recent reports on optimization of the [Pt(dien)(Nucleobase)]²⁺ structure for inhibition of zinc finger peptide-DNA/RNA interactions [47] and the comparison of steric and electronic effects of the chelate by effect of chelate by use of bis(2-pyridylmethyl)amine (bpma) in Pt(II) and Au(III) compounds [48]. Substitution inertness is relative and the Au(III)N₄ structure is still more reactive than its Pt(II)N₄ analogs. We have observed now a general trend of reactivity of Au(bpma) > Au(dien) > Pt(bpma) > Pt(dien) in reactions with C-terminal finger of NCp7 [47,48]. This is exemplified by going from [Pt(dien)(Nucleobase)]²⁺, for example, which interacts with NC in an initially non-covalent manner without Zn²⁺ release, [47], to the [Au(dien)(Nucleobase/heterocycle)]³⁺ reported here where the displacement of Zn²⁺ by Au³⁺ can be controlled by suitable ligand modification [18]. The concept in general has significant potential for optimization for enhancement of intrinsic selectivity for this important target and as a structurally discrete new class of agents capable of disrupting the chaperone activity of NC.

Supplementary Material

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References

1. Levin JG; Guo J; Rouzina I; Musier-Forsyth K In Progress in Nucleic Acid Research and Molecular Biology; Moldave Kivie, Ed.; Academic Press, 2005, 80, 217–286. [PubMed: 16164976]
2. Mori M; Kovalenko L; Lyonnais S; Antaki D; Torbett BE; Botta M; Mirambeau G; Mély Y Curr. Top. Microbiol. Immunol 2015, 389, 53–92. [PubMed: 25749978]
3. Darlix J-L; Garrido JL; Morellet N; Mély Y; deRocquigny H Adv. Pharmacol 2008, 55, 297–347.
4. Wu H; Mitra M; McCauley MJ; Thomas JA; Rouzina I; Musier-Forsyth K; Williams MC; Gorelick RJ Virus Research 2013, 171, 263–277. [PubMed: 22814429]
5. Deshmukh L; Ghirlando R; Clore GM Angew. Chem. Intl. Ed 2014, 53, 1025–1028.
6. Keane SC; Heng X; Lu K; Kharytonchik S; Ramakrishnan V; Carter G; Barton S; Hosic A; Florwick A; Santos J; Bolden NC; McCowin S; Case DA; Johnson BA; Salemi M; Telesnitsky A; Summers MF Science, 2015, 348, 917–921. [PubMed: 25999508]
7. De Guzman RN; Wu ZR; Stalling CC; Pappalardo L; Borer PN; Summers MF Science 1998, 279, 384–388. [PubMed: 9430589]
8. Morellet N; Demene H; Teilleux V; Huynh-Dinh T; De Rocquigny H; Fournie-Zaluski M-C; Roques BP J. Mol. Biol 1998, 283, 419–434. [PubMed: 9769215]
9. Bourbigot S; Ramalanjaona N; Boudier C; Salgado GF; Roques BP; Mély Y; Bouaziz S; Morellet N J Mol. Biol 2008, 383, 1112–1128. [PubMed: 18773912]
10. Bazzi A; Zargarian L; Chaminade F; Boudier C; De Rocquigny H; Rene B; Mély Y; Fosse P; Mauffret O; Nuc. Acid. Res 2011, 39, 3903–3916.
11. Yamagata Y, Kato M, Odawara K, Tokuno Y, Nakashima Y, Matsushima N, Yasumura K, Tomita K, Ihara K and Fujii Y, Cell, 1996, 86, 311–319. [PubMed: 8706135]
12. Labahn J, Schärer OD, Long A, Ezaz-Nikpay K, Verdine GL and Ellenberger TE, Cell, 1996, 86, 321–329. [PubMed: 8706136]
13. Hodel AE, Gershon PD, Shi X, Wang SM and Quirocho FA, Nat. Struct. Biol, 1997, 4, 350–354. [PubMed: 9145102]
14. Marcotrigiano J, Gingras A, Sonenberg N and Burley SK, Cell, 1997, 89, 951–961. [PubMed: 9200613]
15. Anzellotti AI; Bayse CA; Farrell NP Inorg. Chem 2008, 47, 10425–10431. [PubMed: 18939818]
16. Anzellotti AI; Liu Q; Bloemink MJ; Scarsdale JN; Farrell N Chem. Biol 2006, 13, 539–548. [PubMed: 16720275]
17. Spell SR; Farrell NP Inorg. Chem 2015, 54, 79–86. [PubMed: 25531886]
18. Spell SR; Farrell NP Inorg. Chem 2014, 53, 30–32. [PubMed: 24328316]
19. Lee Y-M, Lim CJ Mol. Biol 2008, 379, 545–553.
20. de Paula QA; Mangrum JB; Farrell NP J. Inorg. Biochem 2009, 103, 1347–1354. [PubMed: 19692125]
21. Larabee JL; Hocker JR; Hanas JS Chem. Res. Toxicol 2005, 18, 1943–1954. [PubMed: 16359185]
22. Franzman MA; Barrios AM Inorg. Chem 2008, 47, 3928–3930. [PubMed: 18426199]
23. Jacques A; Lebrun C; Casini A; Kieffer I; Proux O; Latour JM; Sénèque O Inorg Chem. 2015, 54, 4104–4113. [PubMed: 25839236]
24. Laskay ÜA; Garino C; Tsybin YO; Salassa L; Casini A J. Chem. Soc. Chem. Comm 2015, 51, 1612–1615.
25. Abbehausen C; Peterson EJ; de Paiva REF; Corbi PP; Formiga ALB; Qu Y; Farrell NP Inorg. Chem 2013, 52, 11280–11287. [PubMed: 24063530]
26. Loo JA; Holler TP; Sanchez J; Gogliotti R; Maloney L; Reily MD J. Med. Chem 1996, 39, 4313–4320. [PubMed: 8863808]
27. Omichinski JG; Clore GM; Sakaguchi K; Appella E; Gronenborn AM FEBS. 1991, 292, 25–30.
28. Loo J Int. J. Mass Spectrom 2001, 204, 113–123.
29. Fabris D Anal. Chem 2011, 83, 5810–5816. [PubMed: 21651236]
30. Garg D; Torbett BE Virus Res. 2014, 193, 135–143. [PubMed: 25026536]

31. Stephen AG; Worthy KM; Towler E; Mikovits JA; Sei S; Roberts P; Yang Q-E; Akee RK; Klausmeyer P; McCloud TG; Henderson L; Rein A; Covell DG; Currens M; Shoemaker RH; Fisher RJ *Biochem. Biophys. Res. Comm* 2002, 296, 1228–1237. [PubMed: 12207905]
32. Shvadchak V; Sanglier S; Rocle S; Villa P; Haiech J; Hibert M; Van Dorsselaer A; Mely Y; de Rocquigny H *Biochimie* 2009, 91, 916–923. [PubMed: 19401213]
33. Goudreau N; Hucke O; Faucher A-M; Grand-Maitre C; Lepage O; Bonneau PR; Mason SW; Titolo S *J. Mol. Biol* 2013, 425, 1982–1998. [PubMed: 23485336]
34. Turner KB; Hagan NA; Fabris D *Nuc. Acids Res* 2006, 34, 1305–1316.
35. Warui DM; Baranger AM *J. Med. Chem* 2012, 55, 4132–4141. [PubMed: 22480197]
36. Schito ML; Soloff AC; Slovit D; Trichel A; Inman JK; Appella E; Turpin JA; Barratt-Boyes SM *Curr. HIV Res* 2006, 4, 379–386. [PubMed: 16842089]
37. Schito ML; Goel A; Song Y; Inman JK; Fattah RJ; Rice WG; Turpin JA; Sher A; Appella E *AIDS Res. Hum. Retroviruses* 2003, 19, 91–101. [PubMed: 12639244]
38. Bhattacharya S; Osman HJ *Infect.* 2009, 59, 377–386.
39. Lewis MG; DaFonseca S; Chomont N; Palamara AT; Tardugno M; Mai A; Collins M; Wagner WL; Yalley-Ogunro J; Greenhouse J; Chirullo B; Norelli S; Garaci E; Savarino A *AIDS* 2011, 25, 1347–1356. [PubMed: 21505294]
40. Fonteh PN; Keter FK; Meyer D *Biometals*, 2010, 23, 185–196. [PubMed: 20127392]
41. Quintal SM; dePaula QA and Farrell NP *Metallomics*, 2011, 3, 121–139. [PubMed: 21253649]
42. Mendes F; Groessel M; Nazarov AA; Tsybin OO; Sava G; Santos I; Dyson PJ and Casini AJ *Med. Chem* 2011, 54, 2196–2206.
43. Serratrice M; Edafe F; Mendes F; Scopelliti R; Zakeeruddin SM; Grätzel M; Santos I; Cinellu MA and Casini A *Dalton Trans.* 2012, 41, 3287–3293. [PubMed: 22289927]
44. Maynard AT; Covell DG *J. Am. Chem. Soc* 2001, 123, 1047–1058. [PubMed: 11456658]
45. Wang LH; Yang XY; Zhang X; Mihalic K; Fan X-Y; Xiao W; Zack Howard OM; Appella E; Maynard AT; Farrar WL *Nature Medicine* 2004, 10, 40–47.
46. de Paula QA; Tsotsoros SD; Qu Y; Bayse CA; Farrell NP *Inorg. Chim. Acta* 2012, 393, 222–229.
47. Tsotsoros SD; Lutz PB; Daniel AG; Peterson EJ; da Paiva REF; Qu Y; Bayse CA; Farrell NP *Chem Sci.* 2016, DOI: c6sc03445d.
48. Bernardes VHF; Qu Y; Du Z; Beaton J; Vargas MD; and Farrell NP *Inorg. Chem* 2016 DOI: 10.1021/acs.inorgchem.6b01925.

Figure 1. Structures of A) $[\text{Au}(\text{dien})(\text{L})]^{3+}$ (L = 9-EtGua, **I**; DMAP, **II**), B) SL-2 RNA sequence (SL2) and C) 1–55 HIVNCp7 zinc finger (NC). Red box is C-terminal finger used in previous studies.

Figure 2. ESI-FTICRMS spectra (positive ion mode to probe protein) of a 1:1 reaction of NC: [Au(dien)(9-EtGua)]³⁺ showing the 5+ and 6+ charge states. F is apo-peptide without Zn. ApoNC is NC-Zn

Figure 3. CD spectra of the reaction of NC and $[\text{Au}(\text{dien})(9\text{-EtGua})]^{3+}$ with time. Final spectrum is consistent with Zn^{2+} displacement.

Figure 4. FT-ICRMS spectra (negative ion mode to probe RNA) of 2:1 NC•SL2 complex upon addition of 1 eq. $[Au(dien)(9-EtGua)]^{3+}$ (top $t=0$, run immediately; bottom after 15 min reaction)

Figure 5. Effect of $[\text{Au}(\text{dien})(9\text{-EtGua})]^{3+}$ (**I**) on SL2 RNA-NCp7 protein interaction. A. NCp7-SL2 binding control. SL2 was incubated with increasing concentrations of NC for 30 min. Lane 1, SL2 only; Lanes 2–8, SL2 + 1000, 500, 250, 125, 62.5, 31.3, and 15.6 nM NC respectively. B. **I** incubated with 250 nM NCp7 for 30 min. prior to addition of SL2. Lane 1, SL2 only; Lane 2, SL2 + NC; Lanes 3–8 NC, SL2, and 2000, 1000, 500, 250, 125, and 62.5 μM **I**. Representative experiments of two independent repeats is shown.

Figure 6.
Interaction of $[\text{Au}(\text{dien})(9\text{-EtGua})]^{3+}$ abrogates NC binding to cognate SL2 sequence by Zn^{2+} displacement by Au.

Table 1.

In Vitro Inhibition of HIV Infectivity of Au Compounds. IC₅₀: Concentration to inhibit infectivity. TC₅₀: Cytotoxicity. TI: Therapeutic (Selectivity) Index.

Compound (TC ₅₀ (μM))	Virus	IC ₅₀ (μM)	TI (TC ₅₀ /IC ₅₀)
Auranofin (1.72)	Ba-L ^a	0.15	11.6
	NL4-3 ^b	0.20	8.44
	91US001 ^c	0.24	7.29
[Au(dien)(9-EtG)] ³⁺ (> 100)	Ba-L	38.3	> 2.61
	NL4-3	19.1	> 5.24
	91US001	46.5	> 2.15
[Au(dien)(DMAP)] ³⁺ (> 100)	Ba-L	27.6	> 3.62
	NL4-3	19.2	> 5.20
	91US001	20.3	> 4.93
AZT (>1.0)	Ba-L	.0026	> 377
	NL4-3	.0050	> 199
	91US001	.0016	> 614

^aCCR5-tropic, Group M Subtype B, lab-adapted isolate

^bCXCR4-tropic, Group M Subtype B, molecular clone

^cCCR5-tropic, Group M Subtype B, clinical isolate. See SI for full details.