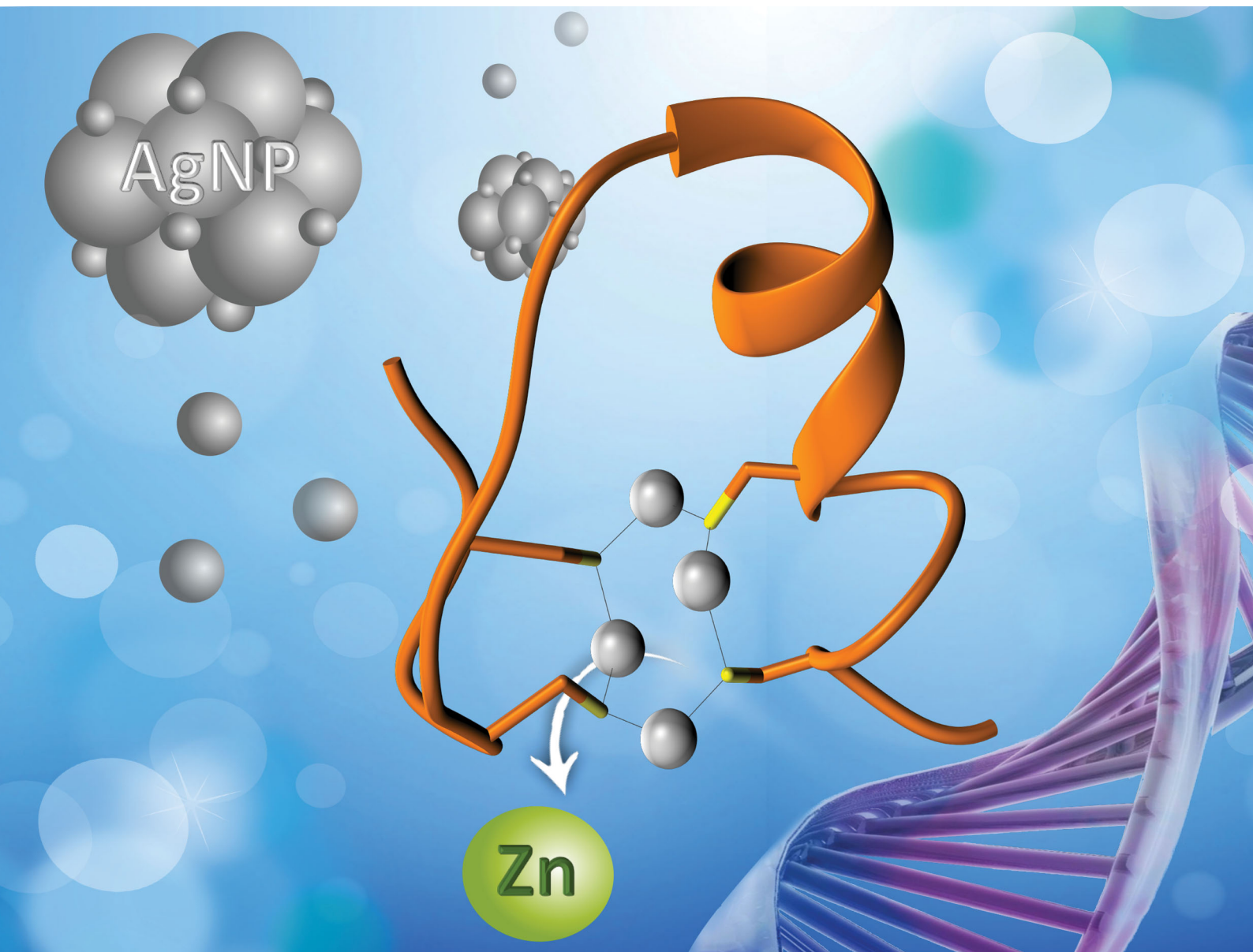


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ISSN 1359-7345

COMMUNICATION

Wojciech Bał, Artur Krężel *et al.*
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Cite this: *Chem. Commun.*, 2020, 56, 1329

Received 4th December 2019,
Accepted 19th December 2019

DOI: 10.1039/c9cc09418k

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Formation of highly stable multinuclear Ag_nS_n clusters in zinc fingers disrupts their structure and function†

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Silver (Ag(I)) binding to consensus zinc fingers (ZFs) causes Zn(II) release inducing a gradual disruption of the hydrophobic core, followed by an overall conformational change and formation of highly stable Ag_nS_n clusters. A compact eight-membered Ag_4S_4 structure formed by a CCCC ZF is the first cluster example reported for a single biological molecule. Ag(I) -induced conformational changes of ZFs can, as a consequence, affect transcriptional regulation and other cellular processes.

Zinc finger (ZF) domains are one of the most abundant structural motifs found in proteins, with a wide range of physiological roles including transcriptional regulation, signal transduction, DNA repair, cell migration, *etc.*^{1–5} Although ZFs contain Zn(II) in their native structures, they provide an impressive range of binding properties for monovalent (Cu(I) and Au(I)), divalent (Cd(II) , Pb(II) , Hg(II) , Pt(II) , Fe(II) , Co(II) , and Ni(II)) and trivalent (Sb(III) and As(III)) metal ions. These cations can compete with Zn(II) , impacting the native domain structure (Fig. S1a, ESI†) and as a consequence influencing the biological functions of ZFs due to their various coordination geometries, ionic radii and ligand specificities.^{4–8} Moreover, they can promote multinuclear species formation, cysteinyl sulfur oxidation or incomplete coordination.^{9–11} All of these outcomes are important in the case of toxic metal ions, which can be absorbed into the body from diet, drinking water, by inhalation, dermal exposure or medical intervention, eventually finding their way to cells to exert toxicity by abolishing the ZF functions.

Extensive research has been devoted to the effects of heavy metals, like Pb(II) , Hg(II) or Cd(II) , on ZF domains, but much less is known about the impact of Ag(I) ions. Various Ag(I) complexes have been vigorously tested for antimicrobial,^{12,13}

anti-inflammatory^{14,15} or anticancer^{16,17} properties. The use of elemental silver nanoparticles (AgNPs) is high and rising rapidly (30% of the products declared to contain NPs are AgNPs). They can be found in multiple daily life applications including food and textile industries, electronics and personal care. Moreover, the AgNP coatings are becoming popular in medical devices and bone implants as antibacterial additives.^{13,18,19} This rapid expansion of AgNPs in various fields elevates the population exposure to metallic nanoparticulates and ionic silver due to the high reactivity of Ag in aerobiosis.²⁰

Animal studies have shown that exposure to Ag in any form leads to its accumulation mostly in the liver and spleen,²¹ while occupational health studies in humans reported deposition of insoluble silver sulfide or selenide precipitates in the dermis and cornea, causing argyria and argyrosis, respectively.^{22–24} Besides, a very slow clearance of Ag has been measured in the brain and testis, two crucial organs.²¹ Cellular experiments using hepatocytes, fibroblasts, and other human cell lines showed that endocytosed AgNPs undergo lysosome-assisted dissolution to Ag(I) ,^{25,26} which is the actual intracellular chemically reactive form of silver. This process leads to the subsequent formation of intracellular Ag(I) -thiolate complexes,^{20,25,27,28} strongly suggesting that Ag(I) ions target cysteine rich proteins and peptides including metallothioneins and glutathione that are abundant. Other proteins with solvent-exposed cysteines such as ZFs could also bind to Ag(I) , with toxic consequences. This issue has not been properly explored, however.

In order to shed more light on how Ag(I) ions affect ZFs, we examined their interactions with consensus peptide 1 (CP1) ZF model peptides containing two (CCHH), three (CCCH) and four (CCCC) Cys residues using UV-Vis and CD spectroscopies, mass spectrometry (MS)-electrospray ionization (ESI) and ion-mobility (IM), quantum mechanics/molecular mechanics (QM/MM), and molecular dynamics (MD) and well-tempered metadynamics (WT-MetaD) simulations. CP1 ZF models have been extensively studied in terms of the coordination chemistry and thermodynamics.^{29,30} We used the recently updated CP1 sequence which represents natural ZF features even better (Fig. S1b, ESI†).³¹

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9cc09418k



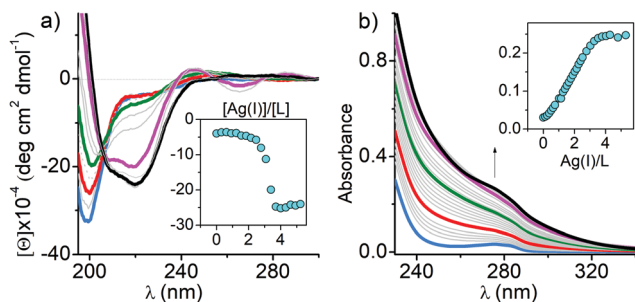


Fig. 1 CD (a) and UV-Vis (b) titrations of 25 μM CCCC ZF in 20 mM TES buffer (pH 7.0) with Ag(I). Blue, red, green, magenta, and black indicate 0, 1, 2, 3, and 4 Ag(I) eq., respectively. The insets show the changes in molar ellipticity and absorption at 220 nm and 279 nm over the range of 0–4 eq. of Ag(I), respectively.

The CD spectra were used to determine whether the Ag(I) binding can induce conformational changes of the ZF peptides. First, 0 to 4 equivalents of Ag(I) were titrated into each ZF. In the CCCC ZF the Ag_4L species (L denotes a given ZF) was formed predominantly, as evidenced by the presence of isodichroic points and the overall stoichiometry (Fig. 1a). Its CD spectrum features two partially overlapping intensive negative bands at 210 and 225 nm indicating the presence of an α -helix with some contribution from a β -sheet. Ag(I) binding also induced a new band at 265 nm, assigned to a $\text{S} \rightarrow \text{Ag(I)}$ charge transfer (CT) band.^{32–34} Ag(I) titrations of other peptides (Fig. S2a, ESI†) yielded various negative and positive bands in the 200–225 nm range indicating the formation of multiple Ag(I) complexes. The comparison with Zn(II) induced CD spectra clearly demonstrates the fundamental alteration of the ZF conformation by Ag(I) coordination in all cases, but the far-UV features observed for CCHH and CCCH ZFs were significantly weaker, indicating a mostly random coil character of the formed Ag(I) complexes.³⁵ The absorption spectra of the same titrations confirmed the formation of $\text{S} \rightarrow \text{Ag(I)}$ CT bands at 240–260 nm. Their molar absorption coefficients increase with the number of Cys residues (Fig. S2b (ESI†) and Fig. 1b).^{32,33} Interestingly when plotted against the Ag(I)-to-ZF molar ratio, the CT intensity linearly corresponds to the number of Cys residues in ZFs, confirming that predominant/final complexes formed for CCHH, CCCH and CCCC ZF peptides are Ag_2L , Ag_3L and Ag_4L , respectively (see below). The course of titrations shows, however, that lower stoichiometry complexes were also formed. Very similar UV-Vis spectral patterns and complex stoichiometries observed in Ag(I) titrations of CCAA and CCCA ZFs (His residues replaced with alanines) indicate the absence of imidazole coordination to Ag(I) ions in CCHH and CCCH (Fig. S3, ESI†). The above observations, together with QM/MM MD simulations (Fig. S4–S6, ESI†), strongly support the formation of Ag_2Cys_2 , Ag_3Cys_3 and Ag_4Cys_4 clusters in Ag_2L , Ag_3L and Ag_4L species, respectively, with sulfur donors bringing two Ag(I) ions. The simulations indicated the absence of secondary structures in Ag_2L and Ag_3L , but the highly regular octagonal Ag_4Cys_4 cluster in Ag_4L maintained the α -helix and a distorted β -sheet, in excellent agreement with the CD results. The Ag_nS_n rings with $n = 3–14$ are known for low

molecular weight thiolates,³⁴ and recently a similar Ag_4Cys_4 cluster was found in the Cu(I) site of dimeric Atox1 copper chaperone (PDB: 5F0W), but the cluster formed by a CCCC ZF is the first one reported for a single biological molecule.

The above results were corroborated by positive-ion ESI-MS-monitored Ag(I) titrations of ZFs performed in pH neutral solutions (Fig. S7, ESI†). The observed 3+ ions of metal free ZFs turn to mono- and then to multinuclear Ag(I) complexes in a stepwise manner, with Ag_2L , Ag_3L and Ag_4L complexes being the most saturated ones for CCHH, CCCH and CCCC ZFs, respectively, confirming the spectroscopic results (Table S1, ESI†). These gas phase data, however, have only a qualitative character.³⁶

The tight structure of the Ag_4L complex of the CCCC ZF was additionally confirmed by MS/MS experiments, in which no molecular fragment could be obtained for this species, as opposed to other studied complexes (Tables S2–S6, ESI†).³⁷ The $[\text{Ag}_4\text{L}]^{3+}$ molecular ion (Fig. S8 (ESI†) for the experimental and simulated isotopic patterns) was subjected to the IM-MS experiment, along with the $[\text{ZnL}]^{3+}$ and L^{3+} ions. The shorter drift time (DT) and the narrower peak of $[\text{Ag}_4\text{L}]^{3+}$, compared to $[\text{ZnL}]^{3+}$, perfectly agree with its simulated compacted structure (Fig. 2 and Fig. S8, ESI†). To further investigate the nature of these conformational changes, the average collision cross-sections (CCSs, Ω) of the ZF complexes in the 3+ charge state were calculated from the DT data (Fig. 2). The Ag_4L complex was more compact ($\Omega = 439 \text{ \AA}^2$) than the ZnL complex ($\Omega = 461 \text{ \AA}^2$) and the apo-peptide L ($\Omega = 512 \text{ \AA}^2$) (Fig. 2). The CCS values calculated with IMPACT software from MD simulations are comparable to those obtained experimentally (Fig. S9, ESI†). To examine if Ag(I) ions could affect stable Zn(II)-ZF complexes, Zn(II) saturated ZF peptides were titrated with Ag(I) ions as monitored by CD and UV-Vis (Fig. S10, ESI†). In all three peptides Zn(II) ions were easily substituted by Ag(I) ions. The comparison of Ag(I) titrations of free and Zn(II)-complexed ZFs shows some differences in the course of signal changes, suggestive of low amounts of mixed metal intermediates, but at stoichiometric Ag(I) concentrations the Ag_2L , Ag_3L and Ag_4L species were formed almost exclusively for peptides with two, three and four cysteinyl residues, respectively. The spectra recorded at those ratios and at a slight Ag(I) excess are identical to those recorded for free peptide titrations. This indicated that the Ag(I) affinity to the respective complexes is at least about two orders of magnitude higher than that of Zn(II). In order to

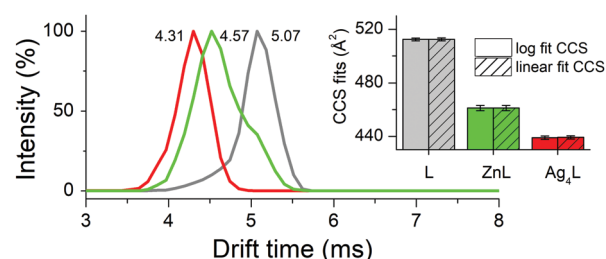


Fig. 2 The arrival time distributions (ATDs) of the CCCC ZF species in the 3+ charge state derived from the IM-MS experiment. The inset shows average collision cross-sections (CCSs) of the CCCC ZF species in the 3+ charge state.



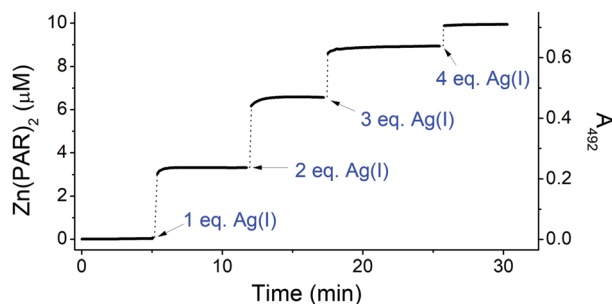


Fig. 3 Zn(II) transfer from 10 μM CCCC ZF to 100 μM PAR during titration with 0–4 eq. of Ag(I). The measurement was performed in 20 mM TES buffer (pH 7.0).

correlate Zn(II) release with Ag(I) binding, the chromophoric Zn(II) chelating probe 4-(2-pyridylazo)resorcinol (PAR) was added to the Zn(II) saturated ZFs, followed by stepwise addition of single mol equiv. of Ag(I) ions, up to four per ZF for consistency.³⁸ Firstly, to show that Ag(I) binding to PAR does not induce absorption at 492 nm, the control experiments (without peptides) were performed (Fig. S11, ESI[†]). After that, the concomitant Zn(II) release steps were determined in a time-dependent mode at 492 nm for Zn(PAR)₂ (Fig. 3 and Fig. S12, ESI[†]). The Zn(II) release was rapid in all cases, with the apparent rate positively correlated with the number of Cys residues. A similar higher reactivity of ZnCys₄ cores has been observed previously for a human/chicken estrogen receptor, *E. coli* N-Ada, HIV-1 nucleocapsid and FOG-1 protein.^{4,39–41} The addition of 1 mol equiv. of Ag(I) to the CCHH ZF caused the release of ca. 60% of Zn(II) from this peptide (Fig. S12, ESI[†]). The 1st Ag(I) effect was less pronounced for CCCH and CCCC ZFs, at ca. 40% and 30%, respectively (Fig. 3 and Fig. S12, ESI[†]). This trend reflects the increased cooperativity of the formation of Ag₃Cys₃ and Ag₄Cys₄ clusters. Furthermore, as showed by us previously, a Cys substitution to other noncoordinating residues weakens the Zn(II) binding more strongly than a His substitution.^{42,43} Individual Cys residues in ZnCys₄ have comparable reactivities and energetic contributions to the Zn(II) complex.^{4,44}

The Ag(I) complexes are too strong for the direct determination of affinity constants; therefore, we performed CD-monitored Ag(I) titration experiments in the presence of 150 μM cyanides (CN[−]), the strongest known simple Ag(I) ligand, used as a competitor.⁴⁵ The binding isotherms were significantly shifted towards higher Ag(I) concentrations (Table S7 and Fig. S13, ESI[†]). This effect was used to estimate the constant for the CCCC ZF, for which the Ag₄L species predominates at all concentrations of added Ag(I). Using $K_{\text{Ag}(\text{CN})_2} = 20.9$ we obtained $\log K_{\text{Ag}_4\text{L}} = 54.9$ corresponding to $\log K = 13.7$ per one Ag(I) for pH 7.0 (see the ESI[†] for details of calculations).⁴⁵ This value is indeed more than 2 log units higher than that for the Zn(II)–CCCC complex ($\log K_{\text{ZnL}} = 11.26$ at pH 7.0) (Fig. S14 and Table S8, ESI[†]).⁴⁶ Due to the significant overlap of intermediate and final complexes for CCHH and CCCH ZFs, their Ag(I) affinities could not be determined precisely, but the ratio of ca. 100 of Ag(I)/Zn(II) affinities appears to remain valid (Fig. S14 and Table S8, ESI[†]). This means that Zn(II) replacement in ZFs by Ag(I) can occur even at low silver exposure, and at higher Ag(I) concentrations the cooperative cluster formation will make this process essentially

thermodynamically irreversible. Taking into account that exposure of AgNPs may load a cell with millions of silver atoms,⁴⁷ our results indicate that ZFs may indeed be targeted intracellularly.

To illustrate the Ag(I)-coupled folding process of the CCCC ZF, a 1.3 μs long WT-MetaD simulation was performed along with two collective variables (CVs) capturing an α -helix and hydrophobic core formation (Tyr1, Phe10 and Phe18).⁴⁸ To fully characterize how the Ag(I) cluster influences the folding, unbiased probability distributions were obtained for a set of CVs through a reweighting procedure (details in the ESI[†], Fig. S15).⁴⁹ The binding of the first Ag(I) ion (two Ag–S bonds) induces structures with the α -helix either folded or unfolded but the hydrophobic core always unpacked (Fig. 4a, Fig. S16a, b and S17, ESI[†]). The modulated formation of the α -helix depends on whether Ag(I) binds to Cys3 and Cys6 or Cys19 and Cys23, the latter located at the α -helix C-terminus (intermediate I2, Fig. S17a and b, ESI[†]). In agreement, the analysis of Ag(I) binding order demonstrated that either Cys6–Ag(I)–Cys3 or Cys23–Ag(I)–Cys19 could be formed first as shown in Fig. S18 (ESI[†]). The binding of the second Ag(I) (four Ag–S bonds) occurs through intermediate I4 forming the Cys3–Ag(I)–Cys19 bonds. This partially folds the α -helix and stabilizes the hydrophobic core (Fig. 4b, Fig. S16a–c, S17 and S18, ESI[†]). The Ag₃L complex (six Ag–S bonds) is formed by the Cys19–Ag(I)–Cys23 coordination (Fig. 4c and Fig. S16a–c, ESI[†]). A basin for this intermediate I5 was not found, suggesting a transient character of this species (Fig. S16b and S17b, ESI[†]). The fourth Ag(I) ion binds *via* Cys6–Ag(I)–Cys23 and stabilizes the hydrophobic core by packing the Tyr1 residue (native state N, Fig. 4d, Fig. S16a–c, S17 and S18, ESI[†]). In the metal-coupled folding mechanism, proposed elsewhere,⁵⁰ partial α -helix formation was observed for the CCHH ZF in the absence of a Zn(II) ion. In contrast, in the Ag(I)-coupled folding mechanism, proposed herein for the CCCC ZF, there is no α -helix formation in the absence of Ag(I). Therefore, stabilization of the α -helix and packing of the hydrophobic core are modulated by the number of Ag(I)–S bonds formed.^{4,50} Such a cooperative stabilization might point towards compact Ag₄Cys₄ cluster formation and indicate that breaking up any of these components will destabilize the CCCC ZF.

Our study demonstrates that Ag(I) ions can directly replace Zn(II) in ZFs of all kinds, forming complexes of various stoichiometries and inducing extensive conformational changes in the native ZF structures. They include the loss of secondary

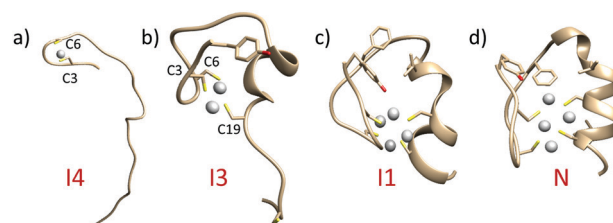


Fig. 4 Average structures for the intermediate and native basins extracted from FES obtained by WT-MetaD. (a) Intermediate 4 (I4); (b) intermediate I3; (c) intermediate I1; and (d) native state (N) (see Fig. S16, ESI[†] for more details).



structure elements in CCHH and CCCH ZFs and formation of an alternative secondary structure in the CCCC ZF. The maximum number of Ag(I) ions bound per ZF is equal to the number of Cys residues, with the highly ordered eight-membered Ag₄Cys₄ structure formed in the CCCC ZF. The Ag(I)-induced conformational changes of biologically active ZF proteins might enhance or inhibit their binding to cognate DNA sequences. Very recently the ability of low dose AgNP exposure to inhibit the activity of hepatocyte nuclear receptors (NR) was demonstrated.⁴⁷ NR are CCCC ZF transcription factors and thus our study provides the molecular basis for the understanding of this and other manifestations of silver toxicity. Our results warrant further research on the chemistry and biology of silver reactivity towards ZF proteins and other cellular thiols in the context of widespread AgNP exposure.

The study was sponsored by the National Science Centre of Poland, grant no. 2016/21/B/NZ1/02847. The authors thank Wrocław Centre for Networking and Supercomputing from Wrocław University of Technology for access to computational resources. The equipment used was sponsored in part by the Centre for Preclinical Research and Technology (CePT), a project co-sponsored by European Regional Development Fund and Innovative Economy, The National Cohesion Strategy of Poland.

Conflicts of interest

There are no conflicts to declare.

Notes and references

- J. H. Laity, B. M. Lee and P. E. Wright, *Curr. Opin. Struct. Biol.*, 2001, **11**, 39–46.
- J. M. Berg, *J. Biol. Chem.*, 1990, **265**, 6513–6516.
- J. H. Laity and G. K. Andrews, *Arch. Biochem. Biophys.*, 2007, **463**, 201–210.
- K. Kluska, J. Adamczyk and A. Krężel, *Coord. Chem. Rev.*, 2018, **367**, 18–64.
- M. Padjasek, A. Kocyla, K. Kluska, O. Kerber, J. B. Tran and A. Krężel, *J. Inorg. Biochem.*, 2020, **204**, 110955.
- Y. Shi, R. D. Beger and J. M. Berg, *Biophys. J.*, 1993, **64**, 749–753.
- A. Hartwig, M. Asmuss, I. Ehleben, U. Herzer, D. Kostelac, A. Pelzer, T. Schwerdtle and A. Bürkle, *Environ. Health Perspect.*, 2002, **110**, 797–799.
- M. Padjasek, M. Maciejczyk, M. Nowakowski, O. Kerber, M. Pyrka, W. Koźmiński and A. Krężel, *Chemistry*, 2020, DOI: 10.1002/chem.201904942.
- S. M. Quintal, Q. A. dePaula and N. P. Farrell, *Metallomics*, 2011, **3**, 121–139.
- A. Witkiewicz-Kucharczyk and W. Bal, *Toxicol. Lett.*, 2006, **162**, 29–42.
- W. Bal, T. Schwerdtle and A. Hartwig, *Chem. Res. Toxicol.*, 2003, **16**, 242–248.
- S. M. Modak, L. Sampath and C. L. Fox Jr, *J. Burn Care Rehabil.*, 1988, **9**, 359–363.
- S. Eckhardt, P. S. Brunetto, J. Gagnon, M. Priebe, B. Giese and K. M. Fromm, *Chem. Rev.*, 2013, **113**, 4708–4754.
- P. L. Nadworny, J. Wang, E. E. Tredget and R. E. Burnell, *Nanomedicine*, 2008, **4**, 241–251.
- K. C. Bhol and P. J. Schechter, *Dig. Dis. Sci.*, 2007, **52**, 2732–2742.
- S. J. Allison, M. Sadiq, E. Baronou, P. A. Cooper, C. Dunnill, N. T. Georgopoulos, A. Latif, S. Shepherd, S. D. Shnyder, I. J. Stratford, R. T. Wheelhouse, C. E. Willans and R. M. Phillips, *Cancer Lett.*, 2017, **403**, 98–107.
- D. F. Shen, S. S. Wu, R. R. Wang, Q. Zhang, Z. J. Ren, H. Liu, H. D. Guo and G. G. Gao, *Small*, 2016, **12**, 6153–6159.
- K. Chaloupka, Y. Malam and A. M. Seifalian, *Trends Biotechnol.*, 2010, **28**, 580–588.
- R. A. Bapat, T. V. Chaubal, C. P. Joshi, P. R. Bapat, H. Choudhury, M. Pandey, B. Gorain and P. Kesharwani, *Mater. Sci. Eng., C*, 2018, **91**, 881–898.
- M. Marchioni, P. H. Jouneau, M. Chevallet, I. Michaud-Soret and A. Deniaud, *Coord. Chem. Rev.*, 2018, **364**, 118–136.
- M. van der Zande, R. J. Vandebriel, E. Van Doren, E. Kramer, Z. Herrera Rivera, C. S. Serrano-Rojero, E. R. Gremmer, J. Mast, R. J. Peters, P. C. Hollman, P. J. Hendriksen, H. J. Marvin, A. A. Peijnenburg and H. Bouwmeester, *ACS Nano*, 2012, **6**, 7427–7442.
- K. D. Rosenman, A. Moss and S. Kon, *J. Occup. Med.*, 1979, **21**, 430–435.
- G. D. DiVincenzo, C. J. Giordano and L. S. Schriever, *Int. Arch. Occup. Environ. Health*, 1985, **56**, 207–215.
- P. L. Drake and K. J. Hazelwood, *Ann. Occup. Hyg.*, 2005, **49**, 575–585.
- G. Veronesi, A. Deniaud, T. Gallon, P. H. Jouneau, J. Villanova, P. Delangle, M. Carriere, I. Kieffer, P. Charbonnier, E. Mintz and I. Michaud-Soret, *Nanoscale*, 2016, **8**, 17012–17021.
- V. De Matteis, M. A. Malvindi, A. Galeone, V. Brunetti, E. De Luca, S. Kote, P. Kshirsagar, S. Sabella, G. Bardi and P. P. Pompa, *Nanomedicine*, 2015, **11**, 731–739.
- X. Jiang, T. Miclăuș, L. Wang, R. Foldbjerg, D. S. Sutherland, H. Autrup, C. Chen and C. Beer, *Nanotoxicology*, 2014, **9**, 181–189.
- G. Veronesi, C. Aude-Garcia, I. Kieffer, T. Gallon, P. Delangle, N. Herlin-Boime, T. Rabilloud and M. Carriere, *Nanoscale*, 2015, **7**, 7323–7330.
- B. A. Krizek, B. T. Amann, V. J. Kilfoil, D. L. Merkle and J. M. Berg, *J. Am. Chem. Soc.*, 1991, **113**, 4518–4523.
- B. A. Krizek, D. L. Merkle and J. M. Berg, *Inorg. Chem.*, 1993, **32**, 937–940.
- A. N. Besold, L. R. Widger, F. Namuswe, J. L. Michalek, S. L. Michael and D. P. Goldberg, *Mol. Biosyst.*, 2016, **12**, 1183–1193.
- A. J. Żelazowski, Z. Gasyna and M. J. Stillman, *J. Biol. Chem.*, 1989, **264**, 17091–17099.
- A. J. Żelazowski and M. J. Stillman, *Inorg. Chem.*, 1992, **31**, 3363–3370.
- B. O. Leung, F. Jalilvand, V. Mah, M. Parvez and Q. Wu, *Inorg. Chem.*, 2013, **52**, 4593–4602.
- G. Park, Z. N. Amaris, M. K. Eiken, K. V. Baumgartner, K. A. Johnston, M. A. Williams, J. G. Marckwordt, J. E. Millstone, K. E. Splanc and K. E. Wheeler, *Environ. Sci.: Nano*, 2019, **6**, 2367–2378.
- H. Mattapalli, W. B. Monteith, C. S. Burns and A. S. Danell, *J. Am. Soc. Mass Spectrom.*, 2009, **20**, 2199–2205.
- Z. Du, R. E. F. de Paiva, K. Nelson and N. P. Farrell, *Angew. Chem., Int. Ed.*, 2017, **56**, 4464–4467.
- A. Kocyla, A. Pomorski and A. Krężel, *J. Inorg. Biochem.*, 2015, **152**, 82–92.
- Y. M. Lee and C. Lim, *J. Am. Chem. Soc.*, 2011, **133**, 8691–8703.
- C. Abbehausen, *Metallomics*, 2019, **11**, 15–28.
- L. H. Wang, X. Y. Yang, X. Zhang, K. Mihalic, Y. X. Fan, W. Xiao, O. M. Howard, E. Appella, A. T. Maynard and W. L. Farrar, *Nat. Med.*, 2004, **10**, 40–47.
- K. Kluska, J. Adamczyk and A. Krężel, *Metallomics*, 2018, **10**, 248–263.
- A. Miłoch and A. Krężel, *Metallomics*, 2014, **6**, 2015–2024.
- T. Kočańczyk, M. Nowakowski, D. Wojewska, A. Kocyla, A. Ejchart and W. Koźmiński, *Sci. Rep.*, 2016, **6**, 36346.
- M. Beck, *Pure Appl. Chem.*, 1987, **59**, 1703–1720.
- O. Sénéque and J. M. Latour, *J. Am. Chem. Soc.*, 2010, **132**, 17760–17774.
- V. Tardillo-Suarez, E. Karepina, B. Gallet, C. Cottet-Rousselle, M. Chevallet, P. Charbonnier, C. Moriscot, I. Michaud-Soret, W. Bal, A. Fuchs, R. Tucoulou, P.-H. Jouneau, G. Veronesi and A. Deniaud, *BioRxiv*, 2019, DOI: 10.1101/825919.
- A. Laio and M. Parrinello, *Proc. Natl. Acad. Sci. U. S. A.*, 2002, **99**, 12562–12566.
- P. Tiwary and M. Parrinello, *J. Phys. Chem. B*, 2015, **119**, 736–742.
- W. Li, J. Zhang, J. Wang and W. Wang, *J. Am. Chem. Soc.*, 2008, **130**, 892–900.

