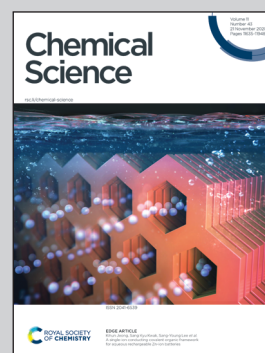


Showcasing research from Professor Hongzhe Sun's laboratory, Department of Chemistry, The University of Hong Kong, Hong Kong, P.R.China.

Atomic differentiation of silver binding preference in protein targets: *Escherichia coli* malate dehydrogenase as a paradigm

Understanding how metallodrugs interact with their protein targets is of vital importance. This work unveiled, at the atomic resolution, a process of silver ion binding to the multiple sites of its authentic protein target MDH *via* a series of Ag-bound MDH structures. Silver-binding preference to the multiple sites of MDH was differentiated as Cys113 > Cys251 > Cys109 > Met227. Besides, silver exhibits preferences to the donor atoms and residues in the order of S > N > O and Cys > Met > His > Lys > Val, respectively.

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# Atomic differentiation of silver binding preference in protein targets: *Escherichia coli* malate dehydrogenase as a paradigm†

Haibo Wang,<sup>1</sup> Xinming Yang,<sup>a</sup> Minji Wang,<sup>ab</sup> Menglong Hu,<sup>c</sup> Xiaohan Xu,<sup>a</sup> Aixin Yan,<sup>d</sup> Quan Hao,<sup>c</sup> Hongyan Li<sup>a</sup> and Hongzhe Sun<sup>1\*</sup>

Understanding how metallodrugs interact with their protein targets is of vital importance for uncovering their molecular mode of actions as well as overall pharmacological/toxicological profiles, which in turn facilitates the development of novel metallodrugs. Silver has been used as an antimicrobial agent since antiquity, yet there is limited knowledge about silver-binding proteins. Given the multiple dispersed cysteine residues and histidine–methionine pairs, *Escherichia coli* malate dehydrogenase (*EcMDH*) represents an excellent model to investigate silver coordination chemistry as well as its targeting sites in enzymes. We show by systematic biochemical characterizations that silver ions ( $\text{Ag}^+$ ) bind *EcMDH* at multiple sites including three cysteine-containing sites. By X-ray crystallography, we unravel the binding preference of  $\text{Ag}^+$  to multiple binding sites in *EcMDH*, i.e.,  $\text{Cys113} > \text{Cys251} > \text{Cys109} > \text{Met227}$ . Silver exhibits preferences to the donor atoms and residues in the order of  $\text{S} > \text{N} > \text{O}$  and  $\text{Cys} > \text{Met} > \text{His} > \text{Lys} > \text{Val}$ , respectively, in *EcMDH*. For the first time, we report the coordination of silver to a lysine in proteins. Besides, we also observed argentophilic interactions ( $\text{Ag} \cdots \text{Ag}$ , 2.7 to 3.3 Å) between two silver ions coordinating to one thiolate. Combined with site-directed mutagenesis and an enzymatic activity test, we unveil that the binding of  $\text{Ag}^+$  to the site IV (His177–Ag–Met227 site) plays a vital role in  $\text{Ag}^+$ -mediated MDH inactivation. This work stands as the first unusual and explicit study of silver binding preference to multiple binding sites in its authentic protein target at the atomic resolution. These findings enrich our knowledge on the biocoordination chemistry of silver(I), which in turn facilitates the prediction of the unknown silver-binding proteins and extends the pharmaceutical potentials of metal-based drugs.

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## Introduction

Despite the wide application of metallodrugs for the treatment of various diseases clinically,<sup>1–5</sup> the molecular targets mediating their biological effects remain largely unknown.<sup>3,6,7</sup> It is very important to understand how metal-based drugs interact with their molecular targets, which defines the overall pharmacological and toxicological profiles of metallo-medicines.<sup>8–11</sup> However, there is a paucity of detailed structural information on the metallodrug–protein adducts so far, which hinders the

exploration of metallodrug targeting sites in proteins and their pharmacological potentials.

Silver ions and silver nanoparticles (AgNPs) are nowadays being extensively used as antimicrobial agents in healthcare and food industry owing to their broad-spectrum antimicrobial properties and less chance of resistance, yet their molecular targets remain largely unknown.<sup>4,12–18</sup> Proteins are deemed to be vital targets of silver and the widespread assumption of silver toxicity is that silver binds to the thiolate or thioester of cysteine or methionine residues and subsequently substitutes the cognate ions in the catalytic sites of metalloenzymes.<sup>16,19,20</sup> Particularly, being isoelectronic with  $\text{Cu}^+$ ,  $\text{Ag}^+$  has been reported to mimic  $\text{Cu}^+$  at its native binding sites in copper proteins by adopting various coordination numbers and geometries.<sup>21–23</sup> However, to date, the binding of  $\text{Ag}^+$  to proteins, especially the authentic silver-binding targets, has not been structurally extensively investigated. To the best of our knowledge, azurin,<sup>24</sup> multicopper oxidase CueO,<sup>21</sup> urease,<sup>25</sup> and trypanothione reductase<sup>26</sup> are the only silver inhibited proteins *in vitro* with available silver-bound crystal structures.

<sup>a</sup>Department of Chemistry, CAS-HKU Joint Laboratory of Metallomics on Health and Environment, The University of Hong Kong, Pokfulam Road, Hong Kong, P. R. China. E-mail: haibo\_wang@connect.hku.hk; hsun@hku.hk

<sup>b</sup>School of Chemistry and Molecular Engineering, East China Normal University, No. 3663 Zhongshan Road North, Shanghai, 200062, P. R. China

<sup>c</sup>School of Biomedical Sciences, The University of Hong Kong, Laboratory Block, 21 Sassoon Road, Pokfulam, Hong Kong, P. R. China

<sup>d</sup>School of Biological Sciences, The University of Hong Kong, Pokfulam Road, Hong Kong, P. R. China

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Considering the diversity of silver-binding proteins and the multifaceted factors defining the silver–protein interaction, the silver targeting sites in proteins should be richly varied and exquisitely attuned to different proteins and residues.<sup>3,16,27</sup> However, limited information on silver-binding proteins restrained the thorough examination, particularly at the atomic level, on which sites or residues of proteins are targeted by silver and what is the preferred coordination geometry of silver in proteins. In the case of multiple sites present, how silver differentiates the binding priority to these sites and what is the driving force for the binding preferences. These long-standing issues remain to be addressed.<sup>16</sup>

Until very recently, we resolved and identified 34 silver-binding proteins ( $\text{Ag}^+$ -proteome) in *E. coli* using our home-made two-dimensional approach of liquid chromatography coupled with GE-ICP-MS (LC-GE-ICP-MS).<sup>27,28</sup> We unveiled that silver eradicates *E. coli* through targeting multiple proteins and disrupting multiple pathways, including glycolysis, TCA cycle, glyoxylate shunt and ROS defense system. *E. coli* malate dehydrogenase (*EcMDH* or MDH) is the only enzyme involved in both TCA cycle and glyoxylate shunt, indicating its vital role as a silver target.

*EcMDH* contains three cysteine residues and several methionine–histidine pairs, while all these cysteine sites are far away from the active sites. These properties of *E. coli* MDH render it as an exceptional model system for investigating the long-standing issues between silver and its protein target. In this work, we found that silver binds to MDH at multiple sites based on systematic characterizations. Subsequently, we demonstrate a diversified coordination behavior of  $\text{Ag}^+$  to MDH by using an array of  $\text{Ag}$ -bound MDH X-ray structures. For the first time, we unravel the  $\text{Ag}^+$  binding preference to its multiple binding sites of its protein target at the atomic level. This work highlights the first successful attempt to capture the binding preference of silver to its protein target, enriching the bio-coordination chemistry of metals.

## Results and discussions

To study how  $\text{Ag}^+$  binds to MDH *in vitro*, we overexpressed and purified the MDH protein (Tables S1 and S2†). The purity was confirmed by size exclusion chromatography (Fig. S1A†) and SDS-PAGE (Fig. S1B†). Binding of  $\text{Ag}^+$  to MDH (monomer) was examined by matrix assisted laser desorption ionization-time of flight mass spectrometry (MALDI-TOF MS). The peak at  $m/z$  of 32 732.4 was assignable to native MDH monomer, while an intense peak and a weak peak at  $m/z$  of 33 053.6 and 33 160.4 (calcd  $m/z$  of 33 052.6 and 33 159.5 respectively) were assignable to MDH with three and four  $\text{Ag}^+$  ions bound respectively (Fig. 1A). Further increasing  $\text{Ag}^+$  concentration to 8 eq. led to neither higher intensity of the peak at  $m/z$  of 33 160.4 nor more peak at higher  $m/z$  (Fig. 1A), verifying that the binding stoichiometry of  $\text{Ag}/\text{MDH}$  is 4 : 1. The MALDI-TOF MS data indicate that the first three  $\text{Ag}^+$  ions might bind MDH with much higher affinity than the fourth one.

We therefore further measured the binding affinity of  $\text{Ag}^+$  towards MDH by isothermal titration calorimetry (ITC). ITC

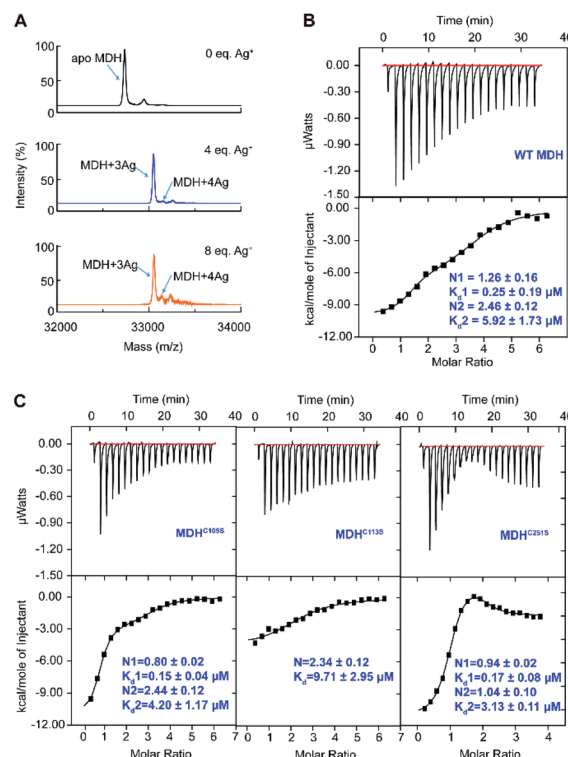


Fig. 1 Silver ions bind to MDH at three cysteine sites and an unknown site. (A) MALDI-TOF mass spectra of apo-MDH and MDH after incubation with 4 and 8 eq. of  $\text{Ag}^+$ . Isothermal titration calorimetry (ITC) results of  $\text{Ag}^+$  binding to wild-type MDH (B) and MDH mutants MDH<sup>C109S</sup>, MDH<sup>C113S</sup> and MDH<sup>C251S</sup> ( $n = 3$ ). The titrations were carried out at 25 °C in 35 mM Tris– $\text{HNO}_3$  and 100 mM  $\text{NaNO}_3$  buffer at pH 7.4. Nitrate rather than other anions is used to avoid potential precipitation of  $\text{Ag}^+$  ions. The data were fitted to a two-set-of-sites binding model using the Origin software. One representative of three replicates is shown.

data exhibited two binding modes and were fitted with a two-set binding model, which gave rise to apparent dissociation constants ( $K_d$ ) of  $0.25 \pm 0.09$  and  $5.92 \pm 1.73 \mu\text{M}$  for the first ( $N_1 = 1.26 \pm 0.16$ ) and second ( $N_2 = 2.46 \pm 0.12$ )  $\text{Ag}^+$ -binding modes respectively (Fig. 1B and Table S3†). These values generally agree well with our previous studies on the binding affinity between silver ions and cysteines (at sub-micro molar level).<sup>29,30</sup> Given that silver is highly thiophilic, we hypothesize that cysteine residues are involved in silver binding and constructed three MDH mutants with cysteines being mutated into serine and then examined the binding affinity of  $\text{Ag}^+$  to MDH mutants under identical conditions. Two binding modes were still observed for MDH<sup>C109S</sup> and MDH<sup>C251S</sup>. In line with the wild-type MDH, MDH<sup>C109S</sup> and MDH<sup>C251S</sup> retained the high binding affinity to  $\text{Ag}^+$  with the binding capacity of *ca.* 0.97 ( $K_d = 0.15 \pm 0.04 \mu\text{M}$ ) and 0.94 ( $K_d = 0.17 \pm 0.08 \mu\text{M}$ ) respectively, indicating that Cys109 and Cys251 are not the strongest binding sites (Fig. 1C and Table S3†). While for the second binding mode, MDH<sup>C109S</sup> and MDH<sup>C251S</sup> bound *ca.* 1.44 and 0.94 eq. of  $\text{Ag}^+$  with the  $K_d$  of  $4.20 \pm 0.17$  and  $3.13 \pm 0.11 \mu\text{M}$  respectively (Fig. 1C and Table S3†). The loss of one weak binding site of MDH<sup>C109S</sup> and MDH<sup>C251S</sup> demonstrates that  $\text{Ag}^+$  binds to





Cys109 or Cys251 with relatively lower affinity. While MDH<sup>C113S</sup> bound  $2.34 \pm 0.12$  molar equivalents of  $\text{Ag}^+$  with the  $K_d$  of  $9.71 \pm 2.75$   $\mu\text{M}$ , and the loss of the strong binding site in this mutant suggests that  $\text{Ag}^+$  binds to Cys113 with relatively higher affinity (Fig. 1C and Table S3†). Collectively, these data demonstrate that  $\text{Ag}^+$  binds to the three cysteine sites of MDH, with Cys113 exhibiting the highest affinity, followed by Cys109 and Cys251.

Our combined data from MALDI-TOF MS and ITC show that silver ions bind to MDH mainly *via* cysteine residues and each cysteine site binds one silver ion under physiological condition. Nevertheless, the coordination geometries as well as the forces driving the binding preference of silver ions to the three cysteine sites and the fourth site remain unclear.

To further explore the local coordination geometry of silver in its binding sites at the atomic level, we attempted to crystallize the apo-form MDH. Tetragonal-shaped single crystals were formed in three days and reached a size suitable for X-ray diffraction after one week (Fig. S2A†) and the structure was solved at the resolution of 1.54 Å (Fig. S2B†). To obtain the  $\text{Ag}^+$ -bound form crystals, we soaked the crystals into cryo-protectant solutions containing 2 mM  $\text{Ag}^+$  for 10 min (Ag-MDH-1), 30 min (Ag-MDH-2), and 1 h (Ag-MDH-3), yielding Ag-bound MDH crystals loaded with various number of  $\text{Ag}^+$  at the resolution from 2.06 Å to 2.55 Å (Table S4†). The crystallographic results showed no significant overall conformational changes among the Ag-bound MDHs and apo-form (RMSD of 0.233 Å, 0.312 Å, and 0.309 Å when superimposing Ag-MDH-1, Ag-MDH-2, and Ag-MDH-3 (Chain A) to apo-MDH, respectively). For all Ag-MDHs, the identification of Ag was made on the significant positive peaks ( $\geq 20\sigma$ ) in the mFo-DFc (difference Fourier) map (Table S5†).

In the crystal structure of Ag-MDH-1, only one  $\text{Ag}^+$  bound to the Cys113 site (Site I), with  $\text{Ag}^+$  coordinating to the side chain of the Sγ of Cys113 and a water molecule (Fig. 2A and B). The

bond lengths of  $\text{Ag}^+$  to Cys113-Sγ or  $\text{H}_2\text{O}$  are the same (2.60 Å) with a pseudo-linear S–Ag–O angle of  $159.1^\circ$  (Tables S6 and S7†). When the crystal soaking duration was extended to 30 min (Ag-MDH-2), four silver ions were observed at Cys113 site (Site I) and Cys251 site (Site II) with two silver ions observed at each site (Fig. 2C). Unlike Ag-MDH-1, the side-chain of Cys113 bound two silver ions with Ag···Ag distance of 2.7 Å in Ag-MDH-2 (Fig. 2D). The first silver ion (Ag401) at this site further coordinated to a water molecule whilst the other (Ag402) to the Nζ of Lys142. Superimposing the Chain A of the two structures showed that the silver bound to Cys113 of Ag-MDH-1 is located in the center of the two silver ions of Ag-MDH-2 (Fig. S3A†). In Ag-MDH-2, Cys251 also bound two silver ions (Ag···Ag distance of 3.1 Å) with one being further coordinated by a water molecule and another by main-chain nitrogen and oxygen of Val173 respectively (Fig. 2E). For the silver ions observed in Ag-MDH-1 and Ag-MDH-2, most of them share a similar quasi-linear coordination geometry with the angle of ligand–Ag–ligand ranging from  $153.8^\circ$  to  $177.7^\circ$  except Ag403 (in Ag-MDH-2) (Table S7†), which has three coordinations with Cys251 and Val173 in nearly trigonal planar geometry.

In the structure of Ag-MDH-3, four  $\text{Ag}^+$ -binding sites with seven silver ions were identified in each chain of MDH. Consistent with the biochemical studies, three of the  $\text{Ag}^+$ -binding sites contain Cys residues, *i.e.*, Cys113 (Site I), Cys251 (Site II), and Cys109 (Site III), with two silver ions found in each thiolate of three cysteine residues (Fig. 3A). Similarly, argentophilic interactions were observed between the two silver ions that bound to the thiolate group of the same cysteine (Ag···Ag, 2.8–3.3 Å) (Fig. 3B–D and Table 1).<sup>31</sup> In addition, each silver ion bound to a nitrogen atom from either a lysine side-chain (Site I)

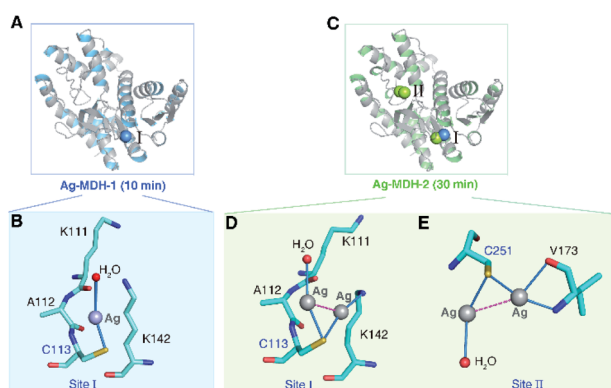


Fig. 2 X-ray crystallographic analysis unveils the coordination geometry of  $\text{Ag}^+$  at Cys113 and Cys251 sites of MDH. (A) Overall structure of Ag-bound MDH at the Cys113 site (Ag-MDH-1). (B) The coordination geometry of  $\text{Ag}^+$  at Cys113 site with  $\text{Ag}^+$  shown as a grey sphere, water as a red sphere. (C) Overall structure of Ag-bound MDH at both Cys113 and Cys251 sites (Ag-MDH-2). The coordination geometry of  $\text{Ag}^+$  at Cys113 site (D) and Cys251 site (E) with  $\text{Ag}^+$  ions shown as grey spheres, water as red spheres and the argentophilic interactions of adjacent Ag in purple dash line.

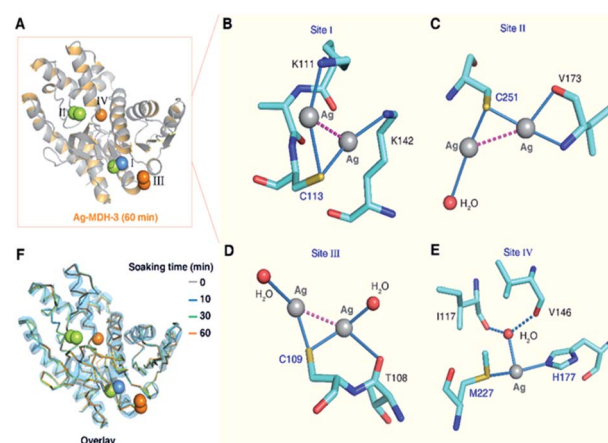


Fig. 3 Crystallographic analysis reveals the coordination geometries of the four silver-binding sites of MDH. (A) Overall structure of Ag-MDH-3. The coordination geometry of silver at the Cys113 site (B), Cys251 site (C), Cys109 site (D) and Met227/His177 site (E).  $\text{Ag}^+$  ions are shown as grey spheres, water as red spheres, the argentophilic interactions in purple dash line and weak interactions between the O atom from Ile117/Val146 and  $\text{Ag}^+$  ions shown as blue dash line. (F) Superimposition of Ag-MDH-1 (blue), Ag-MDH-2 (green), Ag-MDH-3 (yellow) with apo-MDH (grey) with RMSD of 0.233 Å, 0.312 Å, and 0.309 Å respectively. Structural alignment was done over  $\text{C}_\alpha$  residues using Dalilite.



Table 1 Metal–ligand distances (Å) in Ag-MDH-3 structure

Metal	Ligand	Distance (Å)
Ag401	Cys109-Sγ	2.5
Ag402	Cys109-Sγ	2.4
Ag402	Thr108-O	2.9
Ag401	HOH512	2.9
Ag402	HOH511	2.3
Ag401	Ag402	3.0 <sup>a</sup>
Ag403	Cys113-Sγ	2.9
Ag404	Cys113-Sγ	2.5
Ag403	Lys111-Nζ	2.6
Ag404	Lys142-Nζ	2.5
Ag403	Ag404	2.8 <sup>a</sup>
Ag405	Cys251-Sγ	2.3
Ag406	Cys251-Sγ	2.4
Ag405	Val173-N	2.2
Ag405	Val173-O	2.5
Ag406	HOH513	2.9
Ag405	Ag406	3.3
Ag407	Met227-Sδ	2.7
Ag407	His177-Nε2	2.5
Ag407	HOH509	2.7

<sup>a</sup> Only weak interactions exist.

or a valine backbone (Site II), or an additional water molecule (Site III), with most of them forming quasi-linear coordination geometry (Fig. 3B–D and Table 2). In the Cys109 and Cys251 sites, a backbone oxygen atom formed an additional coordination to one of the silver atoms (L–Ag–S, 102–104°). The four silver ions bound to Cys113 and Cys251 in Ag-MDH-3 overlap with those in Ag-MDH-2, albeit that one of the silver ion bound to Cys113 in Ag-MDH-3 is coordinated by the Nζ of Lys111 rather than a water molecule (Fig. S3A and B†).

As a soft metal, Ag<sup>+</sup> exhibits higher affinity to nitrogen than oxygen atoms, the higher binding affinity of Ag<sup>+</sup> to the Cys113 and Cys251 sites than the other sites could be ascribed to the coordination of Ag<sup>+</sup> to the adjacent nitrogen atoms of the lysine side chains (Lys111 and Lys142) and the nitrogen atom of the valine main chain (Val173), respectively (Fig. 3B and C). At the fourth

silver-binding site, Met227, His177 and a water molecule (HOH 509) formed a trigonal pyramidal geometry (the S–Ag–N angle of 146.6°) for Ag<sup>+</sup> binding with lower occupancy than the previous three Ag<sup>+</sup>-binding sites (Fig. 3E). Additionally, the water molecule was further stabilized by two hydrogen bonds (O⋯O, 2.8–3.0 Å) from the backbone oxygen atoms of Ile117 and Val146.

To verify these four sites are indeed silver binding sites, we further constructed a triple cysteine mutant MDH<sup>3CS</sup> as well as two variants with four residues mutated (MDH<sup>3CS/M227S</sup> and MDH<sup>3CS/H177S</sup>) and examined their silver binding capability by ITC and MALDI-TOF MS. For the triple Cys mutant MDH<sup>3CS</sup>, a weak signal of Ag-bound MDH<sup>3CS</sup> was detected by MALDI-TOF MS (MDH<sup>3CS</sup>, *m/z* = 32 681.2; Ag-MDH<sup>3CS</sup>, *m/z* = 32 787.6) (Fig. S4A†), whereas, no obvious Ag<sup>+</sup> binding was observed by ITC (Fig. S4B†), indicating that the affinity of the fourth Ag<sup>+</sup>-binding might be too weak to be detected by ITC. Consistently, no peak corresponding to Ag<sup>+</sup>-binding protein was observed for MDH<sup>3CS</sup> after treatment with 4 eq. of Ag<sup>+</sup> by GE-ICP-MS, for which the weakly bound metal ions are dissociated during the gel electrophoresis separation (Fig. S4C†). For MDH<sup>3CS/M227S</sup> and MDH<sup>3CS/H177S</sup>, no MW change was observed for the former one upon incubation with silver ions, whereas a weak signal for one Ag<sup>+</sup> bound to MDH<sup>3CS/H177S</sup> was observed in MALDI-TOF MS (Fig. S5A and B†), indicating that Ag<sup>+</sup> binds to Met227 with higher affinity than His177. Taken together, these data demonstrate that mutation of the major silver-binding residues could completely abrogate silver binding to MDH, verifying the authenticity of the silver-binding sites in MDH resolved by crystallographic structures.

Compared with MALDI-TOF MS and ITC results (*vide supra*), our crystallographic structures collectively illustrated identical silver-binding sites and a more detailed silver-binding preference in MDH. Despite only one Ag<sup>+</sup> binds to each cysteine site in solution, two Ag<sup>+</sup> were observed in each thiolate with weak argentophilic interactions and seven Ag<sup>+</sup> were found in one MDH molecule in the Ag-MDH-3 crystal structure. The observation of two Ag<sup>+</sup> bound to each cysteine site in the crystal structure is likely due to the use of a high concentration of AgNO<sub>3</sub> (2 mM) and a long duration (1 h) for crystal soaking. Argentophilic interactions between silver ions in the form of di-nuclear,<sup>25,26</sup> tetra-nuclear<sup>32</sup> and even hepta-nuclear<sup>33</sup> were observed in urease, ferredoxin and metallothionein as well. Nevertheless, the observed two Ag<sup>+</sup> ions in each thiolate indeed provide two potential silver binding modes at each cysteine site, which may represent novel silver shuttle systems under physiological conditions.

To investigate the effect of Ag<sup>+</sup>-binding on the enzyme function, we further measured the activity of MDH after incubation with different amounts of Ag<sup>+</sup>. The result showed a dose-dependent inhibition of Ag<sup>+</sup> on MDH activity with 3 eq. and 6 eq. of Ag<sup>+</sup> inhibiting over 50% and 90% of MDH activity respectively (Fig. 4A). To uncover the binding of Ag<sup>+</sup> to which site contribute to MDH inhibition, we compared the structures of Ag-MDH-3 and previously reported structure of EcMDH in ternary complex with citrate (as substrate analog) and cofactor NAD<sup>+</sup> (PDB: 1EMD).<sup>34</sup> Superimposition of 1EMD with Ag-MDH-3 demonstrates that Ag<sup>+</sup>-binding site IV is the one being most adjacent to the active site (Fig. 4B and C). A close look at the comparison of Ag-MDH-3 and NAD-bound MDH shows that the

Table 2 Ligand–Ag–ligand angles (°) in Ag-MDH-3 structure<sup>a</sup>

Atom1	Atom2	Atom3	Angle (°)
HOH512	Ag401	Cys109-Sγ	136.3
HOH511	Ag402	Cys109-Sγ	145.5
Thr108-O	Ag402	Cys109-Sγ	104.6
Lys111-Nζ	Ag403	Cys113-Sγ	154.0
Lys142-Nζ	Ag404	Cys113-Sγ	151.7
Val173-N	Ag405	Cys251-Sγ	168.9
Val173-O	Ag405	Cys251-Sγ	102.7
HOH513	Ag406	Cys251-Sγ	170.3
His177-Nε2	Ag407	Met227-Sδ	146.6
HOH509	Ag407	His177-Nε2	88.3
HOH509	Ag407	Met227-Sδ	89.3

<sup>a</sup> All Ag atoms and water molecules are assigned their universal chain IDs. The measurements are based on the atomic coordination for the polymeric Chain A and Ag atoms or water molecules.



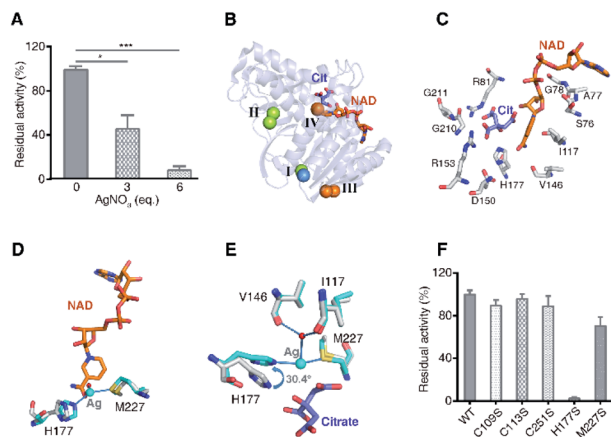


Fig. 4 Binding of Ag<sup>+</sup> to site IV (His177-Met227 site) contributes to Ag<sup>+</sup>-mediated MDH inactivation. (A) Dose dependent inhibition of MDH by Ag<sup>+</sup> ( $n = 3$ ). (B) Superimposition of native MDH (PDB: 1EMD) with Ag-bound MDH (Ag-MDH-3). (C) Active site of MDH with key residues shown in sticks (PDB: 1EMD). (D) An overlay image comparing the relative position of Ag<sup>+</sup> (cyan sphere) with the cofactor NAD<sup>+</sup> (PDB: 1EMD). (E) An overlay image comparing the relative position of Ag<sup>+</sup> (cyan sphere) with the substrate analog citrate (PDB: 1EMD). (F) Normalized residual activity of WT MDH and MDH mutants ( $n = 3$ ). The results are shown as mean  $\pm$  SEM (A, F). \* $P < 0.05$ , \*\* $P < 0.01$  and \*\*\* $P < 0.001$ . Structural alignment was done over C $_{\alpha}$  residues using DaliLite (B–E).

binding of silver to His177 and Met227 form a steric hindrance to the nicotinamide carboxamide of NAD<sup>+</sup> (Fig. 4D). In comparison with citrate-bound MDH, the side chain of His177 in Ag-MDH-3 rotates 30.4° to the opposite direction of citrate for silver binding. Such a rotation results in 2.3 Å shift of the His177-Nε2 away from citrate binding (Fig. 4E).

To further unveil which residue is pivotal for Ag<sup>+</sup> inhibition on MDH activity, we examined the enzyme activities of WT MDH and a series of MDH mutants, including MDH<sup>C109S</sup>, MDH<sup>C113S</sup>, MDH<sup>C251S</sup>, MDH<sup>H177S</sup>, and MDH<sup>M227S</sup>. As shown in Fig. 4F, individual site-directed mutation of Cys109, Cys113, and Cys251 to Ser led to no significant change of MDH activity. In contrast, the mutation of Met227 to Ser resulted in a decrease of ~30% of MDH activity and mutation of His177 to Ser completely inactivated the enzyme (Fig. 4F), demonstrating that these two residues play critical roles for the catalytic activity of MDH. Indeed, previous studies reported that Met227 is involved in the binding of the ribose from NAD<sup>+</sup><sup>34</sup> and the His177–Asp150 pair acts as an essential proton relay system *via* hydrogen bonds during the catalytic reaction.<sup>35–37</sup> Thus, the binding of Ag<sup>+</sup> to the site IV as shown by the crystal structure could potentially disrupt the optimal orientation of NAD<sup>+</sup> to the active site and abrogate the proton transfer capability of catalytic His177, leading to the MDH inactivation.

Taken together, the X-ray studies show that silver ions bind to four sites of MDH with higher affinity to the three Cys sites and relatively lower affinity to the fourth Met/His site. Although the three cysteine sites are all located at the solvent accessible surface, they exhibit different affinities to silver ions due to slightly different coordination spheres of silver ions, *i.e.*, the nitrogen

atoms of side-chains of Lys111/Lys142 and the main-chain of Val173 at Cys113 and Cys251 sites respectively. The distance between Val173/N and Ag405 is shorter (0.7 Å) than that between Thr108/O and Ag402, which might contribute to the higher affinity of silver ions to Cys251 than to Cys109. The crystallographic data are generally coherent with the biochemical results that Ag<sup>+</sup> binds to four sites of MDH albeit the different stoichiometry. Importantly, the binding of Ag<sup>+</sup> to the site IV (His177–Ag–Met227) plays a vital role in Ag<sup>+</sup>-mediated MDH inactivation. For the three cysteine sites, despite the long distance between each cysteine to the active site (>20 Å), whether silver binding to these sites contributes to MDH inhibition, *e.g.*, through unusual allosteric regulation, warrants further investigation.

## Conclusion

We unveil, at the atomic resolution, a process of silver ion binding to the multiple sites of MDH *via* a series of Ag-bound MDH structures. Consistent with the biochemical characterizations, Ag<sup>+</sup> ions bind to the multiple sites of MDH with the binding preference in the order of Cys113 > Cys251 > Cys109 > Met227/His177. Generally, silver ions show a diversified coordination geometry through binding to S, N and O from Cys/Met, His/Lys/Val and Val/H<sub>2</sub>O respectively. Coordination of silver to Lys was observed for the first time in protein structures. The coordination numbers of silver generally range from 2 to 4 and angles from 130° to 178° with quasi-linear as preferred geometry, followed by trigonal planar. We found that, in the case of silver coordination to a single residue, it exhibits preferences to the atoms and residues in the order of S > N > O and Cys > Met > His > Lys > Val respectively. When silver coordinates to the sites containing the same dominant residue of cysteine, the flexibility of surrounding residues together with the solvent accessibility confer the site specific avidity for silver. Moreover, we observed argentophilic interactions between silver ions (Ag<sup>+</sup>–Ag distance of 2.7–3.3 Å) coordinating to the same thiolate of each cysteine site, which could only be observed in soaked crystals (high concentration of Ag<sup>+</sup> being used) but not in solutions. Importantly, combined with enzymatic activity test on the site-directed MDH mutants, we uncover that the binding of Ag<sup>+</sup> to the site IV (His177–Ag–Met227 site) plays a pivotal role in Ag<sup>+</sup>-mediated MDH inactivation.

Collectively, using MDH as a showcase, we demonstrate a diversified bio-coordination chemistry of silver to its authentic protein targets, which are exquisitely attuned to different sites and residues. Our work represents for the first unusual and explicit study of silver-binding to the multiple sites in its authentic protein target at the atomic resolution. Recently, we demonstrated that Ag<sup>+</sup> targets Cys149 and His176 at the active site of GAPDH (an essential enzyme in glycolysis pathway without using metal ion as cofactor), leading to abolishment of its enzymatic activity *via* a mix-type inhibition.<sup>38</sup> The diversity of silver-binding proteins and the multiple properties defining the silver–ligand interactions may provide novel insights into pharmacological and toxicological profiles of antimicrobial silver, thereby opening a new horizon for the study of the interactions between metallo-based medicines and their protein targets. The atomistic knowledge of silver–





ligand interaction could also facilitate the development of novel metalloenzyme inhibitors as metallodrug candidates.<sup>39,40</sup>

## Additional information

The coordinates and structure factors of Ag-MDH-1, Ag-MDH-2 and Ag-MDH-3 are deposit at Protein Data Bank (wwPDB) with accession codes of 5Z3W, 7CGC, and 7CGD, respectively.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 K. J. Franz and N. Metzler-Nolte, *Chem. Rev.*, 2019, **119**, 727–729.
- 2 J. J. Soldevila-Barreda and N. Metzler-Nolte, *Chem. Rev.*, 2019, **119**, 829–869.
- 3 J. A. Lemire, J. J. Harrison and R. J. Turner, *Nat. Rev. Microbiol.*, 2013, **11**, 371–384.
- 4 R. J. Turner, *Microb. Biotechnol.*, 2017, **10**, 1062–1065.
- 5 Y. C. Ong, S. Roy, P. C. Andrews and G. Gasser, *Chem. Rev.*, 2019, **119**, 730–796.
- 6 N. P. E. Barry and P. J. Sadler, *Chem. Commun.*, 2013, **49**, 5106–5131.
- 7 K. D. Mjos and C. Orvig, *Chem. Rev.*, 2014, **114**, 4540–4563.
- 8 H. Wang, Y. Zhou, X. Xu, H. Li and H. Sun, *Curr. Opin. Chem. Biol.*, 2020, **55**, 171–179.
- 9 H. Li, R. Wang and H. Sun, *Acc. Chem. Res.*, 2019, **52**, 216–227.
- 10 M. P. Sullivan, H. U. Holtkamp and C. G. Hartinger, *Met. Ions Life Sci.*, 2018, **18**, 351–386.
- 11 C. G. Hartinger, M. Groessl, S. M. Meier, A. Casini and P. J. Dyson, *Chem. Soc. Rev.*, 2013, **42**, 6186–6199.
- 12 R. J. Turner, *Antibiotics*, 2018, **7**, 112.
- 13 J. J. Harrison, H. Ceri and R. J. Turner, *Nat. Rev. Microbiol.*, 2007, **5**, 928–938.
- 14 J. R. Morones-Ramirez, J. A. Winkler, C. S. Spina and J. J. Collins, *Sci. Transl. Med.*, 2013, **5**, 190ra181.
- 15 J.-Y. Maillard and P. Hartemann, *Crit. Rev. Microbiol.*, 2013, **39**, 373–383.
- 16 S. Eckhardt, P. S. Brunetto, J. Gagnon, M. Priebe, B. Giese and K. M. Fromm, *Chem. Rev.*, 2013, **113**, 4708–4754.
- 17 S. Chernousova and M. Eppele, *Angew. Chem., Int. Ed.*, 2013, **52**, 1636–1653.
- 18 K. M. Fromm, *Nat. Chem.*, 2011, **3**, 178.
- 19 S. Medici, M. Peana, G. Crisponi, V. M. Nurchi, J. I. Lachowicz, M. Remelli and M. A. Zoroddu, *Coord. Chem. Rev.*, 2016, **327**, 349–359.
- 20 J. L. Clement and P. S. Jarrett, *Met.-Based Drugs*, 1994, **1**, 467–482.
- 21 S. K. Singh, S. A. Roberts, S. F. McDevitt, A. Weichsel, G. F. Wildner, G. B. Grass, C. Rensing and W. R. Montfort, *J. Biol. Chem.*, 2011, **286**, 37849–37857.
- 22 M. Meury, M. Knop and F. P. Seebeck, *Angew. Chem., Int. Ed.*, 2017, **56**, 8115–8119.
- 23 F. Long, C. C. Su, M. T. Zimmermann, S. E. Boyken, K. R. Rajashankar, R. L. Jernigan and E. W. Yu, *Nature*, 2010, **467**, 484–U140.
- 24 M. J. Panzner, S. M. Bilinovich, J. A. Parker, E. L. Bladholm, C. J. Ziegler, S. M. Berry and T. C. Leeper, *J. Inorg. Biochem.*, 2013, **128**, 11–16.
- 25 L. Mazzei, M. Cianci, A. Gonzalez Vara and S. Ciurli, *Dalton Trans.*, 2018, **47**, 8240–8247.
- 26 P. Baiocco, A. Ilari, P. Ceci, S. Orsini, M. Gramiccia, T. Di Muccio and G. Colotti, *ACS Med. Chem. Lett.*, 2011, **2**, 230–233.
- 27 H. Wang, A. Yan, Z. Liu, X. Yang, Z. Xu, Y. Wang, R. Wang, M. Koohi-Moghadam, L. Hu, W. Xia, H. Tang, Y. Wang, H. Li and H. Sun, *PLoS Biol.*, 2019, **17**, e3000292.
- 28 L. Hu, T. Cheng, B. He, L. Li, Y. Wang, Y.-T. Lai, G. Jiang and H. Sun, *Angew. Chem., Int. Ed.*, 2013, **52**, 4916–4920.
- 29 X. Liao, F. Yang, R. Wang, X. He, H. Li, R. Y. T. Kao, W. Xia and H. Sun, *Chem. Sci.*, 2017, **8**, 8061–8066.
- 30 X. Liao, F. Yang, H. Li, P.-K. So, Z. Yao, W. Xia and H. Sun, *Inorg. Chem.*, 2017, **8**, 8061–8066.
- 31 H. Schmidbaur and A. Schier, *Angew. Chem., Int. Ed.*, 2015, **54**, 746–784.
- 32 M. Martic, I. N. Jakab-Simon, L. T. Haahr, W. R. Hagen and H. E. Christensen, *J. Biol. Inorg. Chem.*, 2013, **18**, 261–276.
- 33 C. W. Peterson, S. S. Narula and I. M. Armitage, *FEBS Lett.*, 1996, **379**, 85–93.
- 34 M. D. Hall and L. J. Banaszak, *J. Mol. Biol.*, 1993, **232**, 213–222.
- 35 J. K. Bell, H. P. Yennawar, S. K. Wright, J. R. Thompson, R. E. Viola and L. J. Banaszak, *J. Biol. Chem.*, 2001, **276**, 31156–31162.
- 36 M. Lemaire, M. Miginiac-Maslow and P. Decottignies, *Eur. J. Biochem.*, 1996, **236**, 947–952.
- 37 J. J. Birktoft and L. J. Banaszak, *J. Biol. Chem.*, 1983, **258**, 472–482.
- 38 H. Wang, M. Wang, X. Yang, X. Xu, Q. Hao, A. Yan, M. Hu, R. Lobinski, H. Li and H. Sun, *Chem. Sci.*, 2019, **10**, 7193–7199.
- 39 L. Riccardi, V. Genna and M. De Vivo, *Nat. Rev. Chem.*, 2018, **2**, 100–112.
- 40 R. Wang, T. P. Lai, P. Gao, H. Zhang, P. L. Ho, P. C. Woo, G. Ma, R. Y. Kao, H. Li and H. Sun, *Nat. Commun.*, 2018, **9**, 439.

