Structure and metal binding properties of ZnuA, a periplasmic zinc transporter from *Escherichia coli*

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Received: 21 August 2007 / Accepted: 1 November 2007 © SBIC 2007

**Abstract**  ZnuA is the periplasmic Zn$^{2+}$-binding protein associated with the high-affinity ATP-binding cassette ZnuABC transporter from *Escherichia coli*. Although several structures of ZnuA and its homologs have been determined, details regarding metal ion stoichiometry, affinity, and specificity as well as the mechanism of metal uptake and transfer remain unclear. The crystal structures of *E. coli* ZnuA (Eco-ZnuA) in the apo, Zn$^{2+}$-bound, and Co$^{2+}$-bound forms have been determined. Zn$^{2+}$ZnuA binds at least two metal ions. The first, observed previously in other structures, is coordinated tetrahedrally by Glu59, His60, His143, and His207. Replacement of Zn$^{2+}$ with Co$^{2+}$ results in almost identical coordination geometry at this site. The second metal binding site involves His224 and several yet to be identified residues from the His-rich loop that is unique to Zn$^{2+}$ periplasmic metal binding receptors. Electron paramagnetic resonance and X-ray absorption spectroscopic data on CoZnuA provide additional insight into possible residues involved in this second site. The second metal binding site involves His224 and several yet to be identified residues from the His-rich loop that is unique to Zn$^{2+}$ periplasmic metal binding receptors. Electron paramagnetic resonance and X-ray absorption spectroscopic data on CoZnuA provide additional insight into possible residues involved in this second site. The second site is also detected by metal analysis and circular dichroism (CD) titrations. Eco-ZnuA binds Zn$^{2+}$ (estimated $K_d \approx 20 \text{ nM}$), Co$^{2+}$, Ni$^{2+}$, Cu$^{2+}$, Cu$^{+}$, and Cd$^{2+}$, but not Mn$^{2+}$. Finally, conformational changes upon metal binding observed in the crystal structures together with fluorescence and CD data indicate that only Zn$^{2+}$ substantially stabilizes ZnuA and might facilitate recognition of ZnuB and subsequent metal transfer.

**Keywords**  ZnuA · Zinc-specific uptake system · Zinc binding · ATP binding cassette transporter

**Abbreviations**

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
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<tbody>
<tr>
<td>ABC</td>
<td>ATP-binding cassette</td>
</tr>
<tr>
<td>ANS</td>
<td>8-Anilino-1-naphthalenesulfonic acid</td>
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<tr>
<td>apoZnuA</td>
<td>Apo form of ZnuA protein from <em>Escherichia coli</em></td>
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<td>CD</td>
<td>Circular dichroism</td>
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<tr>
<td>CoZnuA</td>
<td>Co$^{2+}$-loaded form of ZnuA protein from <em>Escherichia coli</em></td>
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<tr>
<td>Eco-ZnuA</td>
<td>ZnuA protein from <em>Escherichia coli</em></td>
</tr>
<tr>
<td>EPR</td>
<td>Electron paramagnetic resonance</td>
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</table>
EXAFS	Extended X-ray absorption fine structure
HEPES	N-(2-Hydroxyethyl)piperazine-N'-(2-Hydroxyethanesulfonic acid
Hin-Pzp1	Zn$^{2+}$-binding protein from *Haemophilus influenzae*
ICP-AES	Inductively coupled plasma atomic emission spectroscopy
MBR	Metal-binding receptor
MF	Mag-Fura-2
MOPS	3-(N-Morpholino)propanesulfonic acid
PDB	Protein Data Bank
PEG	Poly(ethylene glycol)
PLBP	Periplasmic ligand-binding proteins
RMSD	RMS deviation
Spn-PsaA	Mn$^{2+}$-binding protein from *Streptococcus pneumoniae*
Syn-MntC	Mn$^{2+}$-binding protein from *Synechocystis PCC sp. 6803*
Syn-ZnuA	Zn$^{2+}$-binding protein from *Synechocystis PCC sp. 6803*
Tpa-TroA	Mn$^{2+}$-binding protein from *Treponema pallidum*
Tris	Tris(hydroxymethyl)aminomethane
Znu	Zn$^{2+}$-specific uptake system
ZnZnuA	Zn$^{2+}$-loaded form of ZnuA protein from *Escherichia coli*

**Introduction**

Zinc serves as an essential cofactor for many enzymes involved in diverse biological processes but can be toxic at high concentrations. Cells regulate the uptake, distribution, and excretion of zinc through several systems, including the Zn$^{2+}$-specific uptake system (Znu). Znu, first identified in *Escherichia coli*, belongs to the ATP-binding cassette (ABC) transporter family [1, 2]. Expressed under low extracellular zinc concentrations in both prokaryotes and eukaryotes, ZnuABC includes three components: the ZnuA periplasmic Zn$^{2+}$-binding protein, the ZnuB integral membrane transporter responsible for Zn$^{2+}$ transport across the cytoplasmic membrane, and the ZnuC ATPase that couples ATP hydrolysis to the transport process. In *E. coli*, cells with mutations in the genes encoding ZnuA and ZnuB exhibit a growth defect when grown on Zn$^{2+}$-depleted medium, which is compensated by the addition of Zn$^{2+}$. In addition, these cells are unable to take up $^{65}$Zn$^{2+}$ [3].

Mutants deficient in the expression of ZnuABC in pathogens such as *Haemophilus influenzae* and *Brucella abortus* exhibit decreased virulence as well as retarded transformation and growth rates [4, 5]. Thus, ZnuABC might play an important role in bacterial survival during host infection.

The extended family of bacterial periplasmic ligand-binding proteins (PLBPs) is divided into nine clusters on the basis of sequence homology and ligand identity. The ZnuA protein from *E. coli* (Eco-ZnuA) together with 47 other putative metal-binding receptors (MBRs) belongs to the newly defined cluster 9. This cluster comprises two MBR families with primary specificities for Zn$^{2+}$ and Mn$^{2+}$ [6]. PLBPs from other clusters transport ions, amino acids, oligopeptides, and sugars. Fourteen MBRs that belong to the proposed Zn$^{2+}$-binding subcluster contain a charged loop rich in acidic and His residues (His-rich loop), located near the metal binding site in the *Synechocystis* PCC sp. 6803 ZnuA (Syn-ZnuA) crystal structure [7]. This loop, which is not found in the Mn$^{2+}$-specific MBRs, is proposed to play a role in periplasmic metal acquisition [7], in regulation of the ABC permease activity through zinc sensing [8], and/or in ZnuA/ZnuB interactions [6]. There are limited experimental data to support these hypotheses, however. Notably, a number of Zn$^{2+}$/Cd$^{2+}$ P1B-type ATPases such as *E. coli* ZntA and *Anabaena* PCC sp. 7120 AztA [9] contain a similar loop, implying a common role for this motif in Zn$^{2+}$ transport.

Metal-bound structures of Syn-ZnuA [7], *Streptococcus pneumoniae* PsaA (Spn-PsaA) [10], *Treponema pallidum* TroA (Tpa-TroA) [11], *Synechocystis* PCC sp. 6803 MntC (Syn-MntC) [12], and Eco-ZnuA [13, 14] as well as the metal-free (apo) structures of Tpa-TroA [15] and mutant Syn-ZnuA (without the His-rich loop) [8] have been determined by X-ray crystallography. The overall architecture of these MBRs is similar to that of other PLBPs from ABC transporter systems. A substrate binding cleft is located between two domains connected by a flexible hinge region. PLBPs are proposed to function by a “Venus-flytrap” mechanism with an open, solvent-accessible ligand-free state and a closed ligand-bound state that are in kinetic equilibrium. Binding of the substrate shifts the equilibrium toward the closed conformation [16–18]. In the cluster 9 MBRs, the hinge region is replaced by a long z-helix that restricts domain motion upon metal exchange. In principle, this z-helix could function as a hinge since only a slight conformational change might be required for the binding of a zinc ion. The structural characterization of apo Tpa-TroA reveals a more closed structure than the Zn$^{2+}$-loaded form, however, with two loops at the C-terminus collapsed into the Zn$^{2+}$-binding pocket, reducing its volume twofold [15]. On the basis of this structure, apo Tpa-TroA was proposed to adopt two states, a nonpermissive form that prevents binding of Zn$^{2+}$ and a hydrated form that is geometrically poised for Zn$^{2+}$ coordination. It is unclear whether this mechanism might be generally applicable to metal binding by all cluster 9 MBRs.
Despite the availability of crystallographic data, detailed biochemical characterization of cluster 9 MBRs has not been reported. Although the structures of Spn-PsaA [10] and Tpa-TroA [11] both contain Zn$^{2+}$, the physiological ligand for these proteins is likely Mn$^{2+}$ [2]. The in vitro metal binding affinities of Tpa-TroA for Zn$^{2+}$ and Mn$^{2+}$ are similar, with $K_d$ values of 1.27 and 0.95 $\mu$M, respectively [19]; that of Syn-ZnuA is substantially higher, approximately 10 nM [8]. Metal binding affinities of other MBRs have not been measured. The stoichiometry of metal binding properties by circular dichroism (CD), optical, electron paramagnetic resonance (EPR), and X-ray absorption spectroscopies. We have also investigated the effect of Zn$^{2+}$ binding on Eco-ZnuA stability and have estimated its Zn$^{2+}$ binding affinity.

**Materials and methods**

Cloning, expression, and purification of Eco-ZnuA

The *E. coli* znuA gene was amplified from genomic DNA (strain W3110) using the forward primer 5′-AAAAAAA-CATATGTTACATAAAAAACGCTT-3′ and the reverse primer 5′-AAAAAGCTTAATCTCCTTTCAGGCAGC-3′, which introduce unique *Nde*I and *Hind*III restriction sites flanking the znuA gene. The 930-bp PCR product was digested with *Nde*I and *Hind*III and ligated into the pET26b vector (Novagen). The correct sequence was confirmed by DNA sequencing. The plasmid was then transformed into *E. coli* BL21(DE3) cells.

The transformants were cultured at 310 K to an optical density at 600 nm of 0.8–1.0 in Luria–Bertani medium supplemented with 25 $\mu$g mL$^{-1}$ kanamycin, and protein production was induced with 0.25 mM isopropyl-β-D-thiogalactopyranoside. Cells were harvested 3 h after induction by centrifugation for 15 min at 8,000g, resuspended in 40 mL of cold 30 mM tris(hydroxymethyl) aminomethane (Tris), pH 7.0, containing 1 mM EDTA and 1 mM phenylmethanesulfonyl fluoride, and lysed by passage through a French press three times at 18,000 lb in$^{-2}$. Cell debris was removed by centrifugation at 125,000g for 25 min, and the supernatant was applied to a Q-Sepharose column (16 × 20 mm) preequilibrated with 30 mM Tris, pH 7.0. Eco-ZnuA was eluted with a linear gradient at approximately 200–400 mM NaCl. Fractions containing Eco-ZnuA were concentrated and treated with 10 mM EDTA for 2 h or more. EDTA and protein impurities were removed by gel filtration on a Superdex 75 column (HiLoad 16/60 prep grade, Pharmacia). The absence of metal in apoZnuA was confirmed by inductively coupled plasma atomic emission spectroscopy (ICP-AES; Varian). To prepare metallated ZnuA for crystallization, apo protein was treated with 0.5 mM zinc acetate (ZnZnuA) or 0.5 mM cobalt(II) chloride (CoZnuA). Excess metal was removed on a Superdex 75 column or by extensive dialysis. The protein concentration was determined using an extinction coefficient at 280 nm of 24,750 M$^{-1}$ cm$^{-1}$ obtained by amino acid analysis. CoZnuA was light pink in color.

Eco-ZnuA crystallization and structure determination

The purified proteins were concentrated to 2.5–10 mg mL$^{-1}$ and exchanged into 20 mM 3-(N-morpholino)propanesulfonic acid (MOPS), pH 7.0 containing 20 mM NaCl and 5% glycerol. Eco-ZnuA was crystallized using both hanging-drop and sitting-drop vapor diffusion methods. ZnZnuA crystallized in two different crystal forms. ZnZnuA crystals belonging to the space group $P2_1_2_1_2_1$ (form I) were obtained at 293 K from a precipitant solution containing 30% (w/v) poly(ethylene glycol) (PEG) 4000, 5% 2-propanol, 0.1 M succinate, pH 4.9, mixed 1:1 with 10 mg mL$^{-1}$ ZnZnuA or 1:2 with 2.5 mg mL$^{-1}$ ZnZnuA. Block-shaped crystals with average dimensions of 0.05 × 0.1 × 0.2 mm$^3$ appeared after 4 h and grew for the next 3 days. For data collection, crystals were transferred to a cryosolution containing 25% (w/v) PEG 4000, 3.5% 2-propanol, 0.1 M N-(2-hydroxyethyl)piperazine-N′-ethanesulfonic acid (HEPES), pH 7.5, 15% glycerol and were flash-cooled in liquid nitrogen. ZnZnuA crystals that belong to the space group $C2_1$ (form II) grew from 30% (w/v) PEG 4000, 5% 2-propanol, 0.1 M HEPES, pH 7.5 and were flash-cooled using the same cryoprotectant. For crystallization of apoZnuA, 5 mg mL$^{-1}$ protein in 20 mM MOPS, pH 7.0, 5 mM EDTA, 20 mM NaCl, 5% glycerol was mixed 1:1 with a cryoprotectant solution containing 35.5% (w/v) PEG 4000, 0.2 M ammonium acetate, 9% glycerol, 0.1 M sodium acetate, pH 4.6. These crystals belong to the space group $P2_1_2_1_2_1$. Finally, 7.25 mg mL$^{-1}$ CoZnuA in buffer H (50 mM HEPES, pH 7.5, 200 mM NaCl) was mixed 1:1 with 25% (w/v) PEG 4000, 3.5% 2-propanol, 0.1 M HEPES, pH 7.5. Light pink crystals of CoZnuA belonging to the space group $P2_1_2_1_2_1$ were obtained and were flash-cooled using the same cryoprotectant as mentioned above.

Data sets were collected at 100 K at the Advanced Photon Source (Argonne National Laboratory) beamlines.
5ID, 19ID, and 23ID, which are operated by DND-CAT, SBC-CAT, and GM/CA-CAT, respectively. X-ray fluorescence was used to verify the presence of zinc in ZnZnuA and cobalt in CoZnuA as well as the absence of metals in apoZnuA. All data were integrated using HKL2000 and scaled with SCALEPACK [20]. Each crystal form contains two molecules in the asymmetric unit. The initial phases were determined by single anomalous dispersion using data collected on crystal form I at a wavelength of 1.2828 Å (Table 1) in 10° segments by the inverse beam technique. Two zinc positions were identified with SHELX [21] and HySS [22], and were refined using SHARP [23]. Solvent flattening yielded an excellent map into which most of the protein was automatically modeled using ARP/wARP [24].

**Table 1** Crystallographic statistics

<table>
<thead>
<tr>
<th>Data collection</th>
<th>ApoZnuA</th>
<th>ZnZnuA (form I)</th>
<th>ZnZnuA (form II)</th>
<th>CoZnuA</th>
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<tbody>
<tr>
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<td>0.9792</td>
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<td>45.00–2.00 (2.07–2.00)</td>
<td>86.00–1.70 (1.76–1.70)</td>
<td>50.00–2.15 (2.23–2.15)</td>
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<td>P2₁2₁2₁</td>
<td>P2₁2₁2₁</td>
<td>C2</td>
<td>P2₁2₁2</td>
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<td></td>
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<td>b = 86.27</td>
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<td></td>
<td>c = 91.56</td>
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<td>262,168</td>
<td>235,145</td>
<td>210,028</td>
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<tr>
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<td>38,461</td>
<td>64,618</td>
<td>32,326</td>
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<td>Completeness (%)</td>
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<td>99.8 (98.4)</td>
<td>97.2 (85.5)</td>
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<td>Redundancy (%)</td>
<td>6.6 (4.4)</td>
<td>6.8 (5.8)</td>
<td>3.6 (3.1)</td>
<td>6.5 (4.7)</td>
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<td>% &gt; 3σ(I)</td>
<td>81.0 (48.9)</td>
<td>81.0 (49.3)</td>
<td>77.8 (44.7)</td>
<td>70.1 (48.3)</td>
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<td>Rsym</td>
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<td>0.082 (0.289)</td>
<td>0.053 (0.323)</td>
<td>0.086 (0.371)</td>
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<tr>
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<td>Rwork (%)</td>
<td>23.96</td>
<td>19.88</td>
<td>21.30</td>
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<td></td>
<td>Rfree (%)</td>
<td>30.25</td>
<td>25.00</td>
<td>25.05</td>
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<td>2</td>
<td>2</td>
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<td></td>
<td>Total atoms</td>
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<td>Metal ion (Å²)</td>
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<td>35.9</td>
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<td></td>
<td>RMS ΔB main chain bond (Å²)</td>
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<td>0.51</td>
<td>0.57</td>
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<tr>
<td></td>
<td>RMS ΔB side chain bond (Å²)</td>
<td>0.98</td>
<td>1.23</td>
<td>1.30</td>
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</tbody>
</table>

*a.u.* asymmetric unit

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Model refinement

The initial model of ZnZnuA form I was built using XtalView [25] and Coot-0.1 [26]. Refinement of the structure was initiated with CNS [27] and completed with Refmac5 [28]. The two ZnZnuA molecules in the form I asymmetric unit are designated molecules A and B. The side chains of residues Asn74, Glu164 and Asn228 in molecule A and of Gln90, Glu181, and Ser186 in molecule B were visible in two conformations, which were modeled appropriately at half occupancy. Two metal sites were identified in each molecule by strong peaks in the anomalous difference Fourier and F_o – F_c difference maps. The main zinc site was fully occupied, whereas the second zinc site was refined to an occupancy of 0.35 in each molecule.
The His-rich loop, residues 117–137, is disordered in the crystal and was partially modeled. The final form I ZnZnuA model consists of residues 27–117 and 138–310 for molecule A and residues 27–118 and 139–309 for molecule B, four zinc ions, and 394 water molecules. A Ramachandran plot calculated with PROCHECK [29] indicates that 93.6% of the residues are within the most favorable regions, with the remainder in additionally allowed regions.

The structures of ZnZnuA form II, CoZnuA, and apoZnuA were solved by molecular replacement with Phaser [30] using ZnZnuA form I without metal as a search model. The two molecules in the asymmetric unit are assigned as molecules C and D for ZnZnuA form II, molecules E and F for CoZnuA, and molecules G and H for apoZnuA. The final model of ZnZnuA form II includes residues 27–116 and 138–310 for molecule C and residues 27–118 and 139–310 for molecule D, three zinc ions, 444 water molecules, and one molecule of 2-propanol. Two zinc sites were identified in molecule C (occupancies of 1.0 and 0.35) and one fully occupied metal site was identified in molecule D. Residues Val69, Ser186, Val273, Met275, Asn284, Ile285, and Lys286 from molecule C and residues Gln101, Arg152, Glu164, Gln183, Thr220, Val253, Met275, and Leu299 from molecule D were modeled in two conformations, each at half occupancy. A Ramachandran plot calculated with PROCHECK [29] shows that 94.0% of the residues are within the most favorable regions, with 5.8% in additionally allowed regions and 0.2% in generously allowed regions.

The final model of CoZnuA consists of residues 27–118 and 139–309 for molecule E and residues 27–120 and 139–309 for molecule F, two cobalt ions, and 291 water molecules. One cobalt site was identified in each molecule. The side chains of residues Gln258 and Lys286 from molecule C and residues Met275, and Leu299 from molecule D were modeled in two occupancies, at half occupancy. A Ramachandran plot shows that 93.2% of residues are within the most favorable regions, with the remainder in additionally allowed regions and one amino acid in the disallowed region. The final model of apoZnuA consists of residues 27–117 and 138–310 for molecule G and residues 27–117 and 138–310 for molecule H, and 49 water molecules. A Ramachandran analysis calculated with PROCHECK [29] indicates that 89.4% of the residues are within the most favorable regions, 10.4% in additional allowed regions, and 0.2% in generously allowed regions. RMS deviations (RMSD) for superposition of different structures were calculated with LSQMAN [31] using the brute force function (fragment length 10, fragment function 2). The coordinates of Eco-ZnuA have been deposited in the Protein Data Bank (PDB) with accession codes 2PS0, 2PRS, 2PS9, and 2PS3 for ZnZnuA form I, ZnZnuA form II, CoZnuA, and apoZnuA, respectively.

Sample preparation for spectroscopic studies

All buffers and water were treated with Chelex 100 and tested for metal impurities by ICP-AES. Before each set of experiments, Eco-ZnuA protein samples were incubated with 10 mM EDTA on ice for more than 2 h followed by removal of EDTA via gel filtration chromatography on a Superdex 75 column or multiple rounds of dilution/concentration. The metal content of samples treated in this way was below the detection limit of ICP-AES. Glassware was soaked for more than 24 h with trace metal grade sulfuric acid and thoroughly rinsed with Chelex-treated water. Between measurements, cuvettes were soaked in 0.2 M EDTA for 10–30 min or in trace metal grade sulfuric acid overnight to prevent metal cross-contamination.

CD spectroscopy

For CD experiments, apoZnuA was exchanged into 20 mM potassium phosphate, pH 7.5, 20 mM NaF buffer and adjusted to approximately 5 μM final protein concentration. CD spectra were recorded for apoZnuA alone and in the presence of various metal ions (Zn²⁺, Mn²⁺, Cu²⁺, and Co²⁺) at room temperature using a JASCO J-715 spectropolarimeter with a 10-nm band width, 50 nm min⁻¹ scan speed, and 1-nm step in a 0.2-cm cell. All spectra were corrected for a buffer background absorbance and converted to mean residue ellipticity. Thermal denaturation of 1.4 μM ZnuA in the apo form and in the presence of 10 μM Zn²⁺, Cu²⁺, Ni²⁺, Co²⁺, and Mn²⁺ was followed by monitoring the CD absorbance at 222 nm (far-UV) in 20 mM potassium phosphate, pH 7.5, 20 mM NaF buffer in a 1-cm cell. The spectropolarimeter was equipped with a Peltier heating unit (single, RTEon). The accuracy of the internal temperature probe was ±0.2 K, and the heating rate was 1 K min⁻¹ in the temperature range 285–363 K. CD spectra were collected between 190 and 280 nm before and after the temperature scan to determine the reversibility of the denaturation process. In most cases, 90% or more of the protein remained folded after the temperature scan. The data analysis [32] yielded the transition temperature, Tm, the enthalpy of unfolding, ΔH, and the free energy of unfolding at room temperature, ΔG_u(298 K). Fits were performed assuming a two-state equilibrium between the native and the unfolded state and constant ΔH (ΔC_p = 0). Fits improved slightly upon including a nonzero ΔC_p.

Spectroscopic analysis of Zn²⁺ binding to Eco-ZnuA

Zinc-binding experiments were conducted in the presence of the indicator Mag-Fura-2 (MF) (Molecular Probes,
Absorption spectra were collected from 250 to 800 nm with a Hewlett-Packard 8452A spectrophotometer at ambient temperature. The spectrometer was blanked against buffer H. ApoZnuA was added to buffer H to a final concentration of 20–30 μM, and a reference spectrum was collected. Freshly prepared MF was then added to a final concentration of approximately 16 μM. The actual concentration of MF was calculated from its absorbance at 366 nm using the extinction coefficient of 29,900 M⁻¹ cm⁻¹. Aliquots (4 μL) of 1 mM Zn²⁺ prepared from the zinc atomic absorption standard (Aldrich, 1,008 μg mL⁻¹) in buffer H were added, mixed well, incubated for 2 min, and spectra were collected. Equilibrium was established within 1–1.5 min after addition of the metal. The absorbance of apoZnuA was subtracted from each spectrum. Data reduction was performed following reported procedures [33–36], and data fitting was performed with the program DynaFit [37]. Each spectrum was corrected for dilution and absorbance at 800 nm.

To estimate the relative affinity of ZnuA and MF for Zn²⁺, competition experiments between fully metallated ZnZnuA and metal-free MF were conducted. First, apoZnuA was incubated overnight with 8–10 equiv of zinc acetate. Excess metal was removed by five successive rounds of concentration/dilution using Amicon Ultra-15 centrifuge filter units with Ultracel-10 membranes. The filter was preequilibrated with thoroughly demetallated buffer H (the zinc content of the buffer was below the detection limit of ICP-AES). In each round, the sample was concentrated to 2 mL and subsequently diluted to 15 mL with buffer H. ZnZnuA was then mixed with increasing concentrations of MF (1–16 equiv) and incubated for 1 h. ZnuA/MF mixtures were separated by gel filtration on Superdex 75 preequilibrated with chelaxed buffer H. In all cases, the protein concentration was measured by the bicinchoninic acid assay [38, 39] and the zinc concentration was measured by ICP-AES.

Spectroscopic studies of Co²⁺-substituted Eco-ZnuA

ApoZnuA was prepared by treating the protein with 10 mM EDTA followed by two cycles of dialysis against 30 mM Tris, pH 7.0 for 12 h, and then by two more cycles of dialysis against 30 mM Tris, pH 6.9 for 12 h. The pH was maintained below 7.0 to slow air oxidation of Co²⁺, which is faster at alkaline pH. The residual Zn²⁺ content was below the detection limit of ICP-AES. Co²⁺-substituted ZnuA was prepared by the direct addition of 1 or 2 equiv of CoCl₂, turning the protein light pink. Spectra were collected at room temperature using an Agilent 8453 UV/vis spectrophotometer. EPR spectroscopy was carried out at 5–13 K, 9.63 GHz, and 2–50-mW microwave power with a Bruker EleXsys E500 spectrometer equipped with an ER4116DM TE₁₀₂/TE₀₁₂ cavity and an Oxford Instruments ESR900 helium-flow cryostat, using 100-kHz magnetic field modulation with 12-G (1.2-mT) modulation amplitude. Simulations were performed using XSophe (Bruker), explicitly taking S = 3/2.

Extended X-ray absorption fine structure studies of Zn²⁺- and Zn²⁺Co²⁺-substituted Eco-ZnuA

Samples for extended X-ray absorption fine structure (EXAFS) studies (1–2 mM) were prepared with 20% (v/v) glycerol as a cryoprotectant, preloaded into Lucite cuvettes with 6-μm polypropylene windows, and frozen rapidly in liquid nitrogen. X-ray absorption spectroscopic data were measured at the National Synchrotron Light Source, beamline X3B, using a Si(111) double-crystal monochromator and a bickel focusing mirror for harmonic rejection. Data collection and reduction were performed according to published procedures [40]. X-ray absorption spectra for ZnuA loaded with 1 equiv of Zn²⁺ were measured on two individually prepared samples (approximately eight scans per sample). Since both datasets gave comparable results, they were averaged, and the average is presented here. All other datasets represent single measurements, eight scans each.

The resulting χ(k) EXAFS data were fit (approximately 11 degrees of freedom for single-scattering and 37 for multiple-scattering fits) using the nonlinear least-squares algorithm contained in the program IFEFFIT, interfaced with SixPack (available free of charge from http://www-ssrl.slac.stanford.edu/~swebb/index.htm). In all fits, the coordination number was varied in integer steps, while the scale factor, S₀, and ΔE₀ were held fixed at calibrated values (for Zn, ΔE₀ = –21 eV, S₀ Zn–N = 0.78; for Co, ΔE₀ = –16 eV, S₀ Co–N = 0.74), allowing only the distance (R₀) and the Debye–Waller factor (σ²) to float. The fits presented in Fig. 10 were generated for Fourier-filtered data; fits to unfiltered data gave similar results (Figs. S4, S5, Table 3).

Multiple-scattering contributions from His ligands were approximated by fitting FEFF calculated paths to the experimental EXAFS, as described previously [40, 41]. Briefly, the imidazolate ring outer-shell scattering is approximated using the four most prominent scattering paths derived from the structure of Zn(imidazole)₄, as calculated by FEFF [42]. The coordination number employed for a given path is adjusted to reflect the total number of scattering paths of similar length. These four “combined” paths are refined independently (varying only R₀ and σ²), and the number of coordinated imidazolates is estimated on the basis of the lowest fit residual in an...
incremental series of fits. The uncertainty in the number of imidazolate ligands determined in this way is estimated to be ±0.5.

Fluorescence with 8-anilino-1-naphthalenesulfonic acid

Conformational changes of Eco-ZnuA upon addition of various metal ions were investigated using the fluorescent dye 8-anilino-1-naphthalenesulfonic acid (ANS; Sigma). A mixture of 0.5 mM ANS and 5 μM apoZnuA in buffer H (initial total volume 800 μL) was placed in a 1.2-mL cuvette (1-cm pathlength). Fluorescence spectra were collected at 296 K using an excitation wavelength of 380 nm (slit width of 8 nm) and an emission wavelength of 420–660 nm (slit width of 8 nm), with ten iterations and a 2-nm step size with an ISS PC1 photon counting spectrofluorometer. The excitation wavelength was determined by collecting an excitation spectrum between 300 and 450 nm with a detected emission at 510 nm. Stock solutions of metals (10 and 1 mM Zn2+, Cd2+, Co2+, Ca2+, Mn2+, Ni2+, Cu2+, and Cu+) were prepared from atomic absorption standards (Aldrich) diluted with buffer H (estimated final concentration of HNO3 is 0.6% by weight). For Cu1+, the oxidation state was maintained by the addition of 20 mM ascorbic acid (1:2 Cu-to-ascorbate ratio). Two sets of experiments were conducted. In the first, metal ions were added to the mixture of 0.5 mM ANS and 5 μM apoZnuA from the stock solutions to 5 and 10 μM final concentrations. In the second set of experiments, the metal ions were added to 0.1, 0.5, and 1.0 mM final concentrations. Fluorescence spectra were collected after each addition. Each initial addition of a new metal ion was followed by a 10-min time scan. Since the time scans showed no significant changes in the emission spectra, all spectra were acquired immediately upon metal addition after thorough mixing. All the data were corrected for dilution.

In addition, the following control experiments were performed. First, stock metal solutions were added to 0.5 mM ANS alone under conditions identical to those described above. No change in ANS fluorescence was observed, indicating the specificity of fluorescence response in the presence of ZnuA. Second, to ensure that dilution of the protein sample with 0.6% HNO3 in buffer H during metal addition had no effect on protein conformation, the experiments were repeated with additions of buffer volumes that corresponded to the volumes of metal solutions. For Cu1+, two additional control experiments were performed to ensure that ascorbate had no effect on ANS fluorescence either directly (Cu/ascorbate mixture was added to ANS alone) or by altering the protein conformation (ascorbate was added to the ANS/protein mixture). In both cases, ANS fluorescence was not affected by the presence of ascorbate.

Results

Overall structure of Eco-ZnuA

The crystal structure of Eco-ZnuA has been solved in the apo, Zn2+-bound (forms I and II), and Co2+-bound forms to 2.45-, 2.00-, 1.70-, and 2.15-Å resolution, respectively (Table 1). While this work was in progress, two other crystal structures of Eco-ZnuA were reported [13, 14] and are virtually identical in fold to the structure of ZnuA form I (Fig. 1). The first 26 N-terminal amino acids are not present in any Eco-ZnuA structure [13, 14] and are believed to comprise a signal peptide. The His-rich loop (residues 116–138) located in the N-terminal domain between β4 and α4 (Fig. 1) has weak electron density and was modeled only partially. In Syn-ZnuA [7] and the other Eco-ZnuA [13, 14] structures, this loop is also disordered. In all the Eco-ZnuA structures, Cys252 and Cys306 form a disulfide bond that might be important for maintaining the integrity of Eco-ZnuA structure or in regulation of Zn2+ binding as was suggested for MntC [12]. There are two molecules of Eco-ZnuA in the asymmetric unit of all crystal forms, yielding eight independent views: molecules A and B in ZnuA form I, C and D in ZnZnuA form II, E and F in CoZnuA, and G and H in apoZnuA. Whereas the two reported Eco-ZnuA structures have a single metal binding site per molecule [13, 14], molecules A, B, and C have two distinct metal binding sites occupied, molecules D, E, and F have only one metal binding site occupied, and molecules G and H contain no metal ions. The RMSD values between ZnZnuA molecule A and the other chains are 0.42 and 0.47 Å for ZnZnuA molecules B and C, respectively, and 0.74–0.91 Å for molecules D, E, F, G, and H. The N-terminal domain and the connecting β6 loop (β6 loop) can adopt both open (position 1, molecules A, E, F, G, and H) and closed (position 2, molecules A, B, C, D, E, and F) conformations, resulting in two different positions of the neighboring helices β6 and α7 (Fig. 2b). The flexibility of this region is manifested by higher B factors. For example, the overall B factor for ZnZnuA molecule C, is 31.6 Å2, whereas the average B factor for the region spanning residues 219–253 is 48.3 Å2.

The primary metal binding site

The main metal binding site was identified by a strong peak in an anomalous Fourier map (Fig. 3a). Its position and coordination are identical to those reported by Li and Jogl [14]. The Zn2+ ion is coordinated by His60, His143, His207, and Glu59 in an almost ideal tetrahedral geometry.
(Table S1). In the structure reported by Chandra et al. [13], coordinating residue Glu59 is replaced with water, resulting in a site more similar to that in Syn-ZnuA [7]. In all four ZnZnuA molecules in the form I and form II structures, the coordination geometry and Zn–N/O distances are within the experimental error of one another (Table S2) and agree well with the average Zn–N His and Zn–O Glu distances reported for tetrahedral binding sites in other proteins [43]. A superposition of molecules A, B, C, and D indicates that the positions of Glu59 and His143 are strictly maintained, the position of His60 varies somewhat (by approximately 0.3–0.8 Å), and His207 clearly occupies two distinct positions related to conformational variation in the β6φ7 loop and helices α6 and α7 (Fig. 3b). Secondary sphere interactions involving the His δ nitrogen atoms and the uncoordinated side chain oxygen of Glu59 are similar to those described for the other Eco-ZnuA structures [13, 14] and observed for Syn-ZnuA [7] and presumably fine-tune the primary ligand sphere for efficient Zn2+ binding [44]. The secondary-sphere carboxylate residues may also act as proton acceptors for the Zn2+-bound His residues if the metal binding site is solvent-inaccessible [43], as is the case for Eco-ZnuA.

The primary metal binding site can also be loaded with Co2+. The overall protein fold and details of metal binding (Fig. 3c) are the same as in the ZnZnuA structure. Ligand His207 occupies position 1 (Fig. 3b), similar to molecule D in the ZnZnuA form II structure. Different orientations of His207 in molecules D, E, and F from those in molecules A, B, and C result in a slight lengthening of the bond between the metal ion and the His207 ε nitrogen from 1.93–1.97 to 2.03–2.12 Å (Table S1). Interestingly, a water molecule is found adjacent to the metal ion in the structures in which His207 occupies position 1 and may account for the altered position of His207. In CoZnuA molecules E and F, this water molecule is 3.11–3.21 Å away from the Co2+ ion and in ZnZnuA, molecule D, it is 3.66 Å away from the Zn2+ ion. By contrast, two water molecules present in the metal-binding pocket of ZnZnuA molecules A, B, and C are substantially further from the metal center, with distances of 4.28–4.45 Å. A search of the PDB reveals a large number of Co2+-containing protein structures with water molecules approximately 3 Å from the Co2+ ion. The only changes in the metal coordination geometry that can be assigned to the substitution of Co2+ for Zn2+ are the lengthening of the distance from the metal to the ε nitrogen atom of His143 from 2.02–2.08 to 2.19–2.21 Å and small changes in the dihedral angles (Table S1). The almost complete conservation of metal coordination geometry between ZnZnuA and CoZnuA indicates that spectroscopic studies of Co2+-substituted MBRs are an appropriate approach to obtain information about Zn2+ binding sites in the absence of crystallographic data.

The apoZnuA structure is completely devoid of metal ions (Fig. 3d). Ligand His143 occupies a similar position, which is somewhat surprising since this residue is only five amino acids from the disordered His-rich loop. Residue His207 in apoZnuA occupies position 1, and His60 and Glu59 are flexible, adopting conformations in the two apoZnuA molecules that differ from one another as well as
from their conformations in the metal-loaded ZnuA structures (Fig. 3e). In the apo Tpa-TroA structure [15], the residue corresponding to Eco-ZnuA His60 flips its position and hydrogen-bonds to a directly preceding Asp residue in a DPH motif that is conserved in Syn-ZnuA and Spn-PsaA. Since His60 is preceded by Ser and Glu residues and does not undergo a similar shift in apoZnuA, the DPH motif is not critical to the metal binding site. This conclusion is supported by the recent mutant Syn-ZnuA structures where zinc release was not accompanied by hydrogen-bond formation in the DPH motif [8]. The secondary-sphere hydrogen-bonding interactions observed in ZnZnuA remain intact in apoZnuA, and additional interactions between the side chain of Glu59 and the side chains of both His143 and His207 are formed.

The second metal binding site

In contrast to the other Eco-ZnuA structures [13, 14], a second metal binding site is clearly present in our structures. Purified, recombinant Eco-ZnuA contains 0.99 ± 0.12 equiv of Zn\(^{2+}\). However, the addition of excess Zn\(^{2+}\) (approximately 5 equiv) followed by extensive dialysis or gel filtration results in 1.85 ± 0.26 Zn\(^{2+}\) ions per Eco-ZnuA molecule, suggesting the presence of a second metal binding site. This site, located approximately 12 Å from the main metal binding site, was identified in anomalous difference Fourier maps of ZnZnuA molecules A, B, and C (Fig. 4a). It is partially occupied with refined occupancies of 0.35 for all three molecules. Since this site is not fully occupied and can be partially or
completely depleted during purification, its affinity for Zn$^{2+}$ is likely lower than that of the primary site. The only visible ligand to this second Zn$^{2+}$ ion is His224, which coordinates via its $\varepsilon$ nitrogen atom (Fig. 4a). The Zn–N distances are 2.35, 2.16, and 2.10 Å in molecules A, B, and C, respectively. The $\delta$ nitrogen atom and backbone NH group of His224 interact with water molecules (distances 2.75–3.18 Å), and the carbonyl oxygen of His224 interacts with the amino group of Arg237 (distances 3.01–3.13 Å).

When the second metal binding site is empty and/or the $\beta$6z6 loop occupies position 1 (Fig. 4b), Arg237 is too far away for this interaction to occur. The position of His224 differs only slightly between the apo and holo Eco-ZnuA structures, which is surprising since His224 is only one residue away from the flexible $\beta$6z6 loop (Fig. 4b). Besides His224, some additional electron density is visible around the second Zn$^{2+}$ ion (Fig. 4a), although it is not possible to identify specific residues. These residues likely derive from the His-rich loop, however. The His-rich loop is on the same face of the protein as the second Zn$^{2+}$ ion (Fig. 1b), and contains multiple His, Asp, and Glu residues, any of which would be candidate ligands. At least one additional ligand may be a His residue because density for His119 from the His-rich loop is observed in the apoZnuA structure in close proximity to His224. EXAFS studies also support the presence of His ligands (vide infra).

**CD spectroscopy**

The metal binding properties of ZnuA were investigated by CD spectroscopy. The far-UV CD spectra of Eco-ZnuA at pH 7.5 titrated with increasing amounts of Zn$^{2+}$ are shown in Fig. 5. Spectral changes occur with the addition of up to 2 equiv of Zn$^{2+}$, indicating a 2:1 zinc-to-protein stoichiometry. This finding is in agreement with the presence of two metal binding sites in the crystal structures as well as with the measured metal stoichiometry of 1.85 ± 0.26.
However, it is possible that the binding of additional metal ions occurs without further spectral changes. Addition of Cd\(^{2+}\) and Co\(^{2+}\) results in spectral changes similar to those observed upon Zn\(^{2+}\) addition (Fig. S1a, c), suggesting that Zn\(^{2+}\), Co\(^{2+}\), and Cd\(^{2+}\) bind similarly. Addition of Mn\(^{2+}\) does not affect the CD spectrum (Fig. S1b).

CD was also used to study the thermal stability of Eco-ZnuA in the presence of various metal ions (apo, Zn\(^{2+}\), Cu\(^{2+}\), Cu\(^{+}\), Ni\(^{2+}\), Co\(^{2+}\), and Mn\(^{2+}\)). In a typical experiment, 90% or more of the protein remains folded immediately following the temperature scan (after cooling the sample to room temperature). This amount drops to approximately 60% in the case of Cu\(^{+}\). Owing to the reversibility of the thermal unfolding process, the data could be fit very well to a simple two-state model of an equilibrium between native folded and unfolded states (Fig. 6, Table 2). In all cases but Cu\(^{+}\), the thermal unfolding curves have relatively sharp transition regions. The much broader transition region of the Cu\(^{+}\)-loaded ZnuA could be related to some irreversibility of the thermal unfolding process. Addition of Zn\(^{2+}\), Co\(^{2+}\), Ni\(^{2+}\), and Cu\(^{2+}\), but not Cu\(^{+}\) and Mn\(^{2+}\), stabilizes ZnuA against thermal unfolding, which agrees with the presence (Zn\(^{2+}\) and Co\(^{2+}\)) or absence (Mn\(^{2+}\)) of CD spectroscopic changes (Figs. 5, S1). Notably, addition of Zn\(^{2+}\) stabilizes Eco-ZnuA the most significantly as judged by a substantial increase in the \(T_m\) value (\(\Delta T_m = 12.5\) K) and in \(\Delta G\) [\(\Delta G_0(298\) K) = 31.8 kJ mol\(^{-1}\)]. Including nonzero \(\Delta C_p\) in the fits resulted in a slight improvement of the fit quality (as assessed by an increase in the correlation coefficient \(R\), and a decrease in \(\chi^2/\text{DOF}\), where DOF is the number of degrees of freedom).

**Zn\(^{2+}\) binding affinity of Eco-ZnuA**

Because Zn\(^{2+}\) is spectroscopically silent and addition of Co\(^{2+}\) produces only small changes in the visible spectrum (Fig. 8, \(\varepsilon_{\text{CoZnuA}} \approx 80\) M\(^{-1}\) cm\(^{-1}\)), the Zn\(^{2+}\) binding affinity of Eco-ZnuA was measured by competition experiments with the chromophoric Zn\(^{2+}\) chelator MF [33–36, 45]. MF forms a 1:1 complex with Zn\(^{2+}\) (Zn-MF, \(K_d = 20\) nM, pH 7.0, 298 K [46]) and is widely used to study metal centers with micromolar to nanomolar binding.
Table 2  Thermodynamic parameters for ZnuA folding (1.4 μM protein, 10 μM metals, in 20 mM phosphate, pH 7.5, 20 mM NaF buffer)

<table>
<thead>
<tr>
<th></th>
<th>ΔH (kJ mol⁻¹)</th>
<th>Tm (K)</th>
<th>ΔGm (kJ mol⁻¹)</th>
<th>ΔCp (kJ mol⁻¹)</th>
<th>ΔH (kJ mol⁻¹)</th>
<th>Tm (K)</th>
<th>ΔGm (kJ mol⁻¹)</th>
<th>ΔCp (kJ mol⁻¹ K⁻¹)</th>
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<tr>
<td>ZnuA</td>
<td>499.9 ± 5.6</td>
<td>337.5 ± 0.1</td>
<td>58.5</td>
<td></td>
<td>495.7 ± 4.2</td>
<td>337.4 ± 0.1</td>
<td>57.9</td>
<td>15.7 ± 0.2</td>
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<tr>
<td>Zn²⁺</td>
<td>607.9 ± 21.0</td>
<td>350.0 ± 0.1</td>
<td>90.3</td>
<td></td>
<td>645.1 ± 30.4</td>
<td>349.6 ± 0.1</td>
<td>95.2</td>
<td>19.6 ± 0.9</td>
</tr>
<tr>
<td>Cu²⁺</td>
<td>709.3 ± 19.5</td>
<td>344.6 ± 0.1</td>
<td>95.9</td>
<td></td>
<td>711.7 ± 18.1</td>
<td>344.6 ± 0.1</td>
<td>96.2</td>
<td>22.3 ± 3.2</td>
</tr>
<tr>
<td>Cu⁺</td>
<td>215.9 ± 3.4</td>
<td>336.7 ± 0.1</td>
<td>24.8</td>
<td></td>
<td>198.2 ± 3.7</td>
<td>335.9 ± 0.1</td>
<td>22.4</td>
<td>8.1 ± 0.2</td>
</tr>
<tr>
<td>Ni²⁺</td>
<td>555.0 ± 10.1</td>
<td>342.2 ± 0.1</td>
<td>71.8</td>
<td></td>
<td>565.1 ± 7.8</td>
<td>342.3 ± 0.1</td>
<td>73.1</td>
<td>19.2 ± 0.3</td>
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<tr>
<td>Co²⁺</td>
<td>570.8 ± 11.0</td>
<td>342.9 ± 0.1</td>
<td>74.7</td>
<td></td>
<td>580.5 ± 9.6</td>
<td>342.9 ± 0.1</td>
<td>76.0</td>
<td>19.6 ± 0.4</td>
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<tr>
<td>Mn²⁺</td>
<td>497.6 ± 5.2</td>
<td>338.6 ± 0.1</td>
<td>59.7</td>
<td></td>
<td>500.2 ± 5.0</td>
<td>338.6 ± 0.1</td>
<td>60.9</td>
<td>10.7 ± 3.1</td>
</tr>
</tbody>
</table>

ΔGm is calculated for 298 K

affinities. The 366 nm absorption maximum of MF shifts to 325 nm upon Zn²⁺ binding, and the extinction coefficient at 366 nm decreases significantly from 29,900 M⁻¹ cm⁻¹ (MF) to 1,880 M⁻¹ cm⁻¹ (Zn-MF). In titrations of an apoZnuA/MF mixture with increasing concentrations of Zn²⁺ ions, the decrease in absorbance at 366 nm is quite slow until approximately 2 equiv of Zn²⁺ is added (Fig. 7). These data indicate that the affinity of Eco-ZnuA for both Zn²⁺ ions is indeed higher than that of MF. This result is in agreement with the isothermal titration calorimetric studies of Syn-ZnuA, in which Kd ~ 10 nM was measured [8].

Spectroscopic studies of CoZnuA

The UV/vis difference spectrum of CoZnuA (spectrum of CoZnuA loaded with 1 equiv of Co²⁺, 1-CoZnuA, minus that of apoZnuA) exhibits peaks at 569 nm (ε = 80 M⁻¹ cm⁻¹) and 602 nm (ε = 80 M⁻¹ cm⁻¹) and a shoulder at 540 nm (Fig. 8). An extinction coefficient between 50 and 250 M⁻¹ cm⁻¹ is consistent with five-coordinate Co²⁺ and agrees well with the crystal structure of CoZnuA, in which Co²⁺ in the main metal binding site is coordinated by three His, a Glu, and a water molecule (Fig. 3c). The UV/vis spectrum of CoZnuA containing 2 equiv of Co²⁺, 2-CoZnuA, is similar in appearance to that of 1-CoZnuA. The difference spectrum (spectrum of 2-CoZnuA minus that of 1-CoZnuA; Fig. S3) exhibits a very weak broad peak at approximately 506 nm with an extinction coefficient of approximately 10 M⁻¹ cm⁻¹, suggestive of six-coordinate Co²⁺ in the second metal binding site.

CoZnuA was also studied by EPR spectroscopy to gain further insight into the Co²⁺ coordination environment (Fig. 9). The EPR spectrum of 1-CoZnuA (Fig. 9, spectrum B) recorded under nonsaturating conditions (13 K, 2 mW) was simulated (Fig. 9, spectrum C) as a single species with the following parameters: S = 3/2, Ms = |±1/2), D ≠ gβBS (50 cm⁻¹ was used), E/D = 0.04, g∥ = 2.25, and g⊥ = 2.38, where D is the axial component of the zero-field splitting and E is the rhombic distortion of the axial zero-field splitting. These parameters and the broad lines are entirely consistent with a five-coordinate Co²⁺ ion with one or more water ligands, as observed in the crystal structure and deduced from UV/vis analysis. The
spectrum of 2-CoZnuA (Fig. 9, spectrum G) yielded a second, distinct Co$^{2+}$ signal (Fig. 9, spectrum D) upon subtraction of the signal from that for 1-CoZnuA. This second signal was also simulated (Fig. 9, spectrum E) as a single chemical species, with parameters consistent with five- or six-coordinate Co$^{2+}$ \[ S = 3/2, M_S = |\pm 1/2|, D \gg g/\beta BS \text{ (50 cm}^{-1} \text{ was used)}, E/D = 0.11, g_\perp = 2.25 \text{ and } g_\parallel = 2.36 \]. Although the coordination of the native Zn$^{2+}$ ion in the second metal binding site is not clear from the crystal structure, the broad lines of the EPR spectrum suggest that the coordination of Co$^{2+}$ in this metal binding site may be increased by one or more water molecules. Addition of the two simulations (Fig. 9, spectrum H) matches the experimental spectrum of 2-CoZnuA recorded under nonsaturating conditions. The signal from either 2-CoZnuA or 1-CoZnuA recorded at 5 K, 50 mW (Fig. 9, spectra A, F) provided no evidence for an \( M_S = |\pm 3/2| \) signal from tetrahedral Co$^{2+}$. The invariance of the signal attributed to the first Co$^{2+}$ ion upon binding a second Co$^{2+}$ ion and the lack of any features or intensity changes that could be assigned to spin-spin coupling indicate that the metal ions do not interact with one another, consistent with the Zn–Zn distances of 12.2–12.4 Å in ZnZnuA structures.

EXAFS studies of Zn- and ZnCo-substituted Eco-ZnuA

To better define the coordination environments of the two distinct metal sites, X-ray absorption spectroscopy experiments were performed. Examination of the main binding site (1-ZnZnuA, Figs. 10a, S4) shows a fairly symmetric first shell exclusively composed of Zn$^{2+}$–N/O scattering and significant outer-shell scattering. Fourier-filtered first-shell fits are consistent with a tetracoordinate metal ion, with four N/O ligands at an average distance of 1.98 Å (Table 3). Separate contributions from nitrogen and oxygen scatterers could not be resolved. Multiple-scattering fits further indicate the presence of three histidyl imidazole ligands, in agreement with the crystal structures.

The structure of the second metal binding site was probed by preparing a mixed-metal derivative, with 1 equiv of Zn$^{2+}$ added first, followed by addition of 1 equiv of Co$^{2+}$. Zinc K-edge EXAFS of the mixed-metal derivative (Fig. S4, Table 3) gave the same results as 1-ZnZnuA, indicating that Zn$^{2+}$ populates exclusively the main binding site. EXAFS of the Co$^{2+}$ in the auxiliary site shows an increase in Fourier transform magnitude, suggestive of higher
The best fits indicate asymmetric coordination (four O and two N donors) (Table 3). The distance difference of 0.13 Å is within the resolution of the data (0.12 Å). Multiple-scattering fits indicate the presence of two His ligands. Inclusion of an additional shell of two carbon scatterers at 2.66 Å, representative of carboxylate coordination, improves the Fourier-filtered fit by approximately 34% (while modestly increasing the number of variable parameters from 12 to 14), although it only minimally affects the unfiltered fit residual. Therefore, the predicted coordination of the second metal binding site based on EXAFS data is consistent with two His, two Asp/Glu, and two water molecules. These results are fully consistent with the crystallographic and EPR data.

Metal specificity of ZnuA monitored by ANS fluorescence

ANS is a useful hydrophobic fluorescent probe to study protein conformational changes, hydrophobic sites in native proteins, and protein/substrate interactions. The ANS fluorescence in aqueous solutions is low and increases substantially and blueshifts when ANS is associated with hydrophobic protein surfaces or nonpolar environments. The addition of 1 and 2 mol equiv of transition metals (Zn²⁺, Co²⁺, Mn²⁺, Ni²⁺, Cd²⁺, Cu⁺, and Cu²⁺) to a ZnuA/ANS mixture causes a decrease in the ANS fluorescence, indicating that each metal ion induces conformational changes that decrease exposed hydrophobic surfaces on ZnuA (Fig. 11a). However, selectivity for different metal ions is observed at higher metal concentrations (0.1–1.0 mM). At these concentrations, Zn²⁺, Cd²⁺, and Cu⁺ cause a substantial increase in ANS fluorescence (Fig. 11b). When Co²⁺ was added to the same mixture, the fluorescence intensity immediately decreased to the level of ANS fluorescence in the absence of protein. It is not clear why different results are obtained for ANS fluorescence in the presence of Co²⁺ as compared with Zn²⁺ given that both bind in the same site in the structures. A possible explanation lies in the different positions of the β6α6 loop in the Zn²⁺ and Co²⁺ structures. In three of four ZnZnuA molecules (A, B, and C), this loop is in the closed position, whereas in both CoZnuA molecules (E and F), the loop remains open (Fig. 2), suggesting Co²⁺ binding may not lead to the same conformational changes as Zn²⁺ binding. A similar decrease in ANS fluorescence upon initial metal addition was observed for Ni²⁺, Mn²⁺, and Cu²⁺. Addition of higher concentrations of these metals increased the observed fluorescence of the mixture, but not as much as with Zn²⁺, Cd²⁺, and Cu⁺. Minimal change in fluorescence was observed upon addition of Ca²⁺. In control experiments, the addition of buffer H containing 0.6% HNO₃ to the ZnuA/ANS mixture or the addition of metal solutions (Zn²⁺, Cd²⁺, Co²⁺, Ca²⁺, Mn²⁺, Ni²⁺, Cu²⁺, and Cu⁺) to ANS alone resulted in minimal if any change in fluorescence (Fig. S6). These data suggest that Zn²⁺, Cd²⁺, and Cu⁺ have the greatest effect on the Eco-ZnuA conformation, whereas Co²⁺, Mn²⁺, Ni²⁺, and Cu²⁺ either do not bind ZnuA as efficiently or do not induce substantial conformational changes.
Discussion

Metal specificity

According to the CD data, the addition of Zn$^{2+}$, Co$^{2+}$, and Cd$^{2+}$, but not Mn$^{2+}$, induces conformational changes in Eco-ZnuA (Figs. 5, S1). In agreement with this result, Zn$^{2+}$, Co$^{2+}$, Ni$^{2+}$, and Cu$^{2+}$, but not Mn$^{2+}$ or Cu$^{+}$, stabilize Eco-ZnuA against thermal denaturation, with Zn$^{2+}$ having the highest stabilization effect ($\Delta T_m = 12.5$ K). ANS fluorescence data using low metal concentrations (1–2 mol equiv) indicate that a broad range of metal ions affect the conformation of Eco-ZnuA. When excess metal ions are added, Zn$^{2+}$, Cd$^{2+}$, and Cu$^{+}$ have a much stronger effect (Fig. 11b). For Cu$^{+}$, the conformational changes may lead to decreased stability (Fig. 6). Although ZnuA interacts with various divalent transition metals, only Zn$^{2+}$ is capable of substantially improving its structural stability. This finding, combined with the high affinity of ZnuA for Zn$^{2+}$ ($K_d < 20$ nM) and the observation that Zn$^{2+}$ and Cd$^{2+}$ most significantly affect the conformation as monitored by ANS fluorescence, suggests that Zn$^{2+}$ is the physiological substrate.

A comparison of the primary metal binding site in all of the structurally characterized cluster 9 MBRs provides insight into how metal ion specificity is conferred by the
protein scaffold. Syn-ZnuA, Eco-ZnuA, Spn-PsaA, and Tpa-TroA were crystallized with Zn$^{2+}$ in the metal binding site, although Spn-PsaA is predicted to bind Mn$^{2+}$ [2]. The specificity of Tpa-TroA is more controversial, with Mn$^{2+}$, Zn$^{2+}$, and Fe$^{2+}$ as possible substrates [19, 47]. Syn-MntC is the only MBR crystallized with Mn$^{2+}$, its natural substrate [12]. In all the structures, the main metal binding site has a similar location and ligand composition. Two ligands, His60 and His143 (Eco-ZnuA numbering), are strictly conserved. Beyond these similarities, three structural features can be correlated with metal specificity. The first determinant is the identity of the third ligand at position 207 with His (Syn-ZnuA, Eco-ZnuA, Tpa-TroA) specifying Zn$^{2+}$ and Glu (Spn-PsaA and Syn-MntC) specifying Mn$^{2+}$ or Fe$^{2+}$, which prefer a higher coordination number [11]. However, the Glu residue in this position adopts monodentate coordination in all the reported structures. The second determinant is the role of the conserved Asp residue at position 279 [7]. In the Mn$^{2+}$-binding proteins, Asp279 coordinates the metal ion, whereas in the Zn$^{2+}$-binding proteins, it points away from the metal binding site. Asp279 is monodentate in Spn-PsaA and Syn-MntC specifying Mn$^{2+}$ or Fe$^{2+}$, which prefer a higher coordination number [11]. However, the Glu residue in this position adopts monodentate coordination in all the reported structures. The second determinant is the role of the conserved Asp residue at position 279 [7]. In the Mn$^{2+}$-binding proteins, Asp279 coordinates the metal ion, whereas in the Zn$^{2+}$-binding proteins, it points away from the metal binding site. Asp279 is monodentate in Spn-PsaA and Syn-MntC specifying Mn$^{2+}$ or Fe$^{2+}$, which prefer a higher coordination number [11]. However, the Glu residue in this position adopts monodentate coordination in all the reported structures.

Metal stoichiometry

Metal analysis, spectroscopic, and crystallographic data clearly indicate that Eco-ZnuA binds at least two metal ions, of which only one was detected in the previous ZnuA structures [13, 14]. This finding is consistent with the reported stoichiometry of 1.6–1.9 Zn$^{2+}$ ions per molecule of the ZnuA homolog Hin-Pzp1 [4], which is 49.2% identical to Eco-ZnuA. The present Eco-ZnuA structures, the second Zn$^{2+}$ ion is coordinated by His224 and at least three additional ligands, presumably provided by the His-rich loop. The His-rich loop was suggested to play a role in metal acquisition [7] or in periplasmic zinc sensing [8] in Syn-ZnuA, but only a single Zn$^{2+}$ ion was found to bind to this protein in the crystal structures. Of the other cluster 9 MBRs, His224 is only conserved in *Vibrio cholerae* ZnuA. The additional Zn$^{2+}$ ligands at this site have not been identified, but the structures of ZnZnuA and spectroscopic...
data for CoZnuA suggest the involvement of His119, two carboxylate residues, and perhaps two water molecules. Coordinating water molecules are likely since the second metal binding site is solvent-exposed. It may be that the presence of multiple His, Asp, and Glu residues in the His-rich loop provides multiple options for Zn$^{2+}$ binding, thus increasing the probability of metal acquisition.

Mechanistic implications

The structural and biochemical data presented here combined with previous reports [7, 10, 11, 15, 19] provide insight into the mechanism of metal acquisition and transport by Eco-ZnuA. The binding of Zn$^{2+}$ in the main metal binding site of apoZnuA probably results in local conformational changes in the loop that houses ligands Glu59 and His60 (Fig. 3e). The rest of Eco-ZnuA, including the long connecting helix, is unaltered. Therefore, Eco-ZnuA does not function by the "Venus-flytrap" mechanism proposed for PLBPs from other clusters. Similar conclusions were drawn from the structures of apo and Zn$^{2+}$-loaded Tpa-TroA [11, 15] and mutant Syn-ZnuA [8, 13] and are consistent with the idea that cluster 9 MBRs constitute an evolutionary distant group of ABC-type PLBPs [10]. The conservation of secondary-sphere interactions between apoZnuA and ZnZnuA suggests that the metal binding site is largely preformed prior to Zn$^{2+}$ binding. In the two apoZnuA molecules, the $\beta6z6$ loop occupies position 1, whereas in three of the four ZnZnuA molecules, the loop is closed (position 2). The binding of the first Zn$^{2+}$ is thus probably followed by a shift of the $\beta6z6$ loop from open position 1 to closed position 2 (Fig. 2b), shielding the metal site from solvent. The loop movement is accompanied by shifts in helices $z6$ and $x7$ (Fig. 2b) and subtle alterations in the main metal binding site such as the shift of His207 from position 1 to position 2 (Fig. 3b) and the movement of a water molecule away from the metal ion. These proposed conformational changes are consistent with the CD (Fig. 5) and ANS fluorescence (Fig. 11) data and might be necessary for interaction with ZnuB and subsequent metal release.

The role of the second metal binding site in Eco-ZnuA function is not clear. In the structures, the occupancy of this site is correlated with the closed position of the $\beta6z6$ loop. The orientation of $z6$ in the closed-loop position allows for the interaction between the side chain of Arg237 and the carbonyl oxygen of His224, the only visible ligand to the second Zn$^{2+}$ ion, and may stabilize the second site. This site may only be filled after Zn$^{2+}$ binds in the primary site, and its reduced occupancy in the structures implies a somewhat lower affinity for Zn$^{2+}$. It is not known whether the second site needs to be occupied for delivery to ZnuB, but it seems unlikely since $z6$ and the $\beta6z6$ loop in Syn-ZnuA, Spn-PsaA, and Tpa-TroA with a single Zn$^{2+}$ ion adopt conformations very similar to those in Eco-ZnuA with two bound Zn$^{2+}$ ions (Fig. S7). It may be that the second site functions as a scavenger or chaperone, sequestering Zn$^{2+}$ when periplasmic concentrations are limited. Biochemical and mutagenesis studies targeting the His-rich loop and its individual residues should further elucidate the functional importance of the second metal binding site.

In sum, the crystal structures of ZnZnuA reveal a second weakly occupied metal binding site not detected in the recent structures of Eco-ZnuA [13, 14]. This site, the existence of which is confirmed by metal analysis and spectroscopic data, likely involves the His-rich loop. Binding of Zn$^{2+}$ stabilizes ZnuA, and the Zn$^{2+}$ dissociation constant, $K_d$, is estimated to be less than 20 nM. Eco-ZnuA interacts with Co$^{2+}$, Ni$^{2+}$, Cu$^{2+}$, Cu$^+$, and Cd$^{2+}$, but not Mn$^{2+}$. Several structural changes in the C-terminal domain of Eco-ZnuA occur upon binding of the initial Zn$^{2+}$ ion. Replacement of Zn$^{2+}$ with Co$^{2+}$ results in almost identical coordination geometry at the primary site. Although the second site is not occupied in the CoZnuA structure, EPR and EXAFS data on samples with Co$^{2+}$ in the second site provide additional insight into possible ligands to the second site. Finally, fluorescence and CD data provide insight into metal ion specificity and conformational changes upon metal binding. Since homologs of Eco-ZnuA are found in numerous pathogens [1], a detailed understanding of its mechanism could impact the development of new antimicrobials.

Acknowledgements This work was supported by National Institutes of Health Grants GM079411 (to M.W.C.), AI056231 and EB001980 (to B. B.), P20RR-16480 (to D.L.T.), and GM58518 (to A.C.R.). Use of the Advanced Photon Source was supported by the US Department of Energy, Office of Science, Office of Basic Energy Sciences under contract no. DE-AC02-06CH11357. We thank the staff at the APS DND-CAT, SBC-CAT, and GM/CA-CAT beamlines for assistance with data collection, P. Focia for preliminary data collection, the Northwestern Keck facility for use of the fluorimeter and UV/vis spectrometer, and J. Arguëllo, E. Eren, B. Mitra, and M. Golynskiy for valuable discussions.

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