Metal Binding Properties of Escherichia coli YjiA, a Member of the Metal Homeostasis-Associated COG0523 Family of GTPases

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**ABSTRACT:** GTPases are critical molecular switches involved in a wide range of biological functions. Recent phylogenetic and genomic analyses of the large, mostly uncharacterized COG0523 subfamily of GTPases revealed a link between some COG0523 proteins and metal homeostasis pathways. In this report, we detail the bioinorganic characterization of YjiA, a representative member of COG0523 subgroup 9 and the only COG0523 protein to date with high-resolution structural information. We find that YjiA is capable of binding several types of transition metals with dissociation constants in the low micromolar range and that metal binding affects both the oligomeric structure and GTPase activity of the enzyme. Using a combination of X-ray crystallography and site-directed mutagenesis, we identify, among others, a metal-binding site adjacent to the nucleotide-binding site in the GTPase domain that involves a conserved cysteine and several glutamate residues. Mutations of the coordinating residues decrease the impact of metal, suggesting that metal binding to this site is responsible for modulating the GTPase activity of the protein. These findings point toward a regulatory function for these COG0523 GTPases that is responsive to their metal-bound state.

GTPases are molecular switches that contribute to a wide variety of critical cellular processes ranging from ribosomal protein synthesis to the cell cycle.1−3 The majority of GTPases are members of the phosphate-binding loop (P-loop) NTPase class that share a mononucleotide-binding fold composed of a central, mostly parallel, β-sheet surrounded by α-helices.4,5 All P-loop NTPases contain conserved structural elements, including the P-loop (Walker A motif), the Walker B motif, and the switch I region, which are essential for binding and hydrolysis of GTP as well as for the ensuing conformational changes that trigger the downstream effects.2,4 The P-loop NTPases can be phylogenetically sorted into several families based on shared structural and sequence features.4 One such family is composed of the G3E NTPases, which possess a glutamate residue in place of the conserved aspartate in the Walker B motif, responsible for coordinating the catalytically essential magnesium ion.4 The G3E family features four main subfamilies, three of which (HypB, UreG, and MeaB/ArgK) are based on prototypical proteins with established roles in the assembly of metalloenzyme active sites.6 The fourth subfamily, COG0523, is a more diverse and ubiquitous group of proteins with mostly unknown functions and a distribution across all three domains of life.4,6

A recent phylogenetic study, which suggested that the COG0523 subfamily of P-loop GTPases can be separated into at least 15 subgroups, predicted that some members of this subfamily function in metal metabolism.5 For example, several subgroups were linked to zinc homeostasis because of promoters containing the DNA recognition sequence of Zur, a zinc-responsive transcription factor, consistent with the zinc-dependent regulation observed for COG0523 members from several organisms.6−8 In addition, there are a few reports about individual COG0523 proteins that support specific roles in transition metal pathways. The first member of this subfamily identified was *Pseudomonas denitrificans* CobW, a protein essential for cobalamin biosynthesis.10 The function of CobW is ambiguous, but it has been suggested that it is responsible for the delivery of cobalt to the cobaltochelatase during assembly of the metallocofactor.11−13 Another COG0523 member is the nitrile hydratase activator protein Nha3, which is involved in the biosynthesis of the iron-dependent nitrile hydratase in *Rhodococcus* sp. N-771 and is proposed to be responsible for trafficking the proper metal to the enzyme precursor.14 A third described COG0523 factor is *Bacillus subtilis* YciC. Originally designated as a low-affinity zinc transporter,7 YciC has since been proposed to serve as a metallochaperone for delivery of the metal cofactor into YciA, a backup enzyme involved in folate biosynthesis.15 Both YciC in *B. subtilis*7 and the more recently reported YeiR from *Escherichia coli*16 help their host...
bacteria survive under zinc-limited growth conditions. Furthermore, in vitro analysis of YeIR revealed a link between metal binding and the protein’s oligomerization state and GTPase activity.\textsuperscript{16}

Despite the large size of this subfamily and its connection with metal homeostasis, very little is known about the biochemical properties of the COG0523 constituents. The only high-resolution structural information about this subfamily is provided by the crystal structure of apo-YjiA from \textit{E. coli},\textsuperscript{17} making this protein an attractive target for structure–activity analysis. To garner further information about the metal binding activities of the COG0523 proteins and to better understand the possible connection with metal homeostasis, we conducted a bioinorganic characterization of YjiA. Metal binding studies demonstrate that YjiA can bind stoichiometric cobalt, 2 equiv of nickel, or four zinc ions in solution, and loading the protein with metal inhibits the GTPase activity. Using X-ray crystallography and spectroscopy, we identified a unique metal-binding site that, unlike those found in the UreG or HhpB subfamilies,\textsuperscript{18,19} is located in a solvent-accessible pocket formed by a cysteine and two glutamate residues. This metal-binding site is adjacent to the nucleotide-binding site and is likely responsible for the observed effects of metal on GTP hydrolysis by YjiA. This report improves our understanding of this large class of proteins, supporting a role for metal-dependent regulation, and will be useful in dissecting the function of the uncharacterized COG0523 proteins.

\section*{Materials and Methods}

\textbf{Materials.} Restriction endonucleases and T4 DNA ligase were obtained from New England Biolabs. Primers (Table S1 of the Supporting Information) were purchased from Sigma Genosys. All chromatography media were from GE Healthcare. Kanamycin, tris(2-carboxyethyl)phosphine (TCEP), phenylmethylsulfonyl fluoride (PMSF), and isopropyl \(\beta\)-d-thiogalactopyranoside (IPTG) were purchased from BioShop (Toronto, ON). Nickel chloride, zinc sulfate, and cobalt sulfate salts (as a minimum, 99.9\% pure) were purchased from Aldrich. Nickel chloride, zinc sulfate, and cobalt chloride were obtained from New England Biolabs. Primers (Table S1 of the Supporting Information) were purchased from Sigma (St. Louis, MO).

\textbf{WT and Mutant YjiA Expression Vector Construction.} The \textit{yjiA} gene was amplified from genomic \textit{E. coli} (DH5\(\alpha\)) DNA using primers (Table S1 of the Supporting Information) designed with restriction sites for NdeI (YjiA forward) and XhoI (YjiA reverse). The digested polymerase chain reaction (PCR) product was ligated with T4 DNA ligase into the pET24b vector (Novagen) digested with NdeI and XhoI and dephosphorylated with calf intestinal phosphatase (New England Biolabs). The mutations were introduced into the YjiA-pET24b vector using QuikChange PCR mutagenesis (Stratagene) with \textit{Pfu} Turbo DNA polymerase and the primers listed in Table S1 of the Supporting Information. The template was subsequently digested with DpnI. The E42A/C66A/C67A and E37A/C66A/C67A triple mutants were constructed using the C66A/C67A YjiA-pET24b vector as a template and the E42A and E37A primers, respectively. For production of large amounts of the WT and mutant plasmids, the plasmids were transformed into XL-2 Blue \textit{E. coli} (Stratagene) and isolated using the Fermentas GeneJET plasmid miniprep kit. All plasmids were sequenced (ACGT, Toronto, ON) in the forward and reverse directions.

\textbf{Protein Expression and Purification.} For expression of WT and mutant YjiA, the plasmids were transformed into BL21 Star(DE3) \textit{E. coli} (Novagen). Overnight cultures were grown, and 25 mL was used to inoculate 1.5 L of LB medium supplemented with 50 \(\mu\)g/mL kanamycin. The cells were grown aerobically at 37 \(^\circ\)C until the OD\_600 reached 0.6, at which point they were induced with 1 mM IPTG. After incubation at 37 \(^\circ\)C for 5 h, the cells were harvested by centrifugation and resuspended in 20 mM Tris (pH 7.5) and 100 mM NaCl supplemented with 4 mM TCEP and 200 \(\mu\)M PMSF. For each preparation of purified protein, a total of 9 L of cell culture was used. All subsequent steps were performed at 4 \(^\circ\)C or on ice. The resuspended cells were sonicated and centrifuged at 25000g for 40 min. The supernatant was passed through a 0.45 \(\mu\)m syringe filter and then loaded onto a DEAE-Sepharose anion-exchange column (GE Healthcare) equilibrated with buffer A [20 mM Tris (pH 7.5) and 1 mM TCEP]. Fractions from a NaCl gradient were screened by sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS–PAGE) using 12.5\% gels. WT and mutant YjiA eluted at approximately 250 mM NaCl. Fractions containing the protein of interest were pooled and dialyzed against buffer A. The sample was then loaded onto a HiTrapQ anion-exchange column (GE Healthcare) equilibrated with buffer A. Once again, fractions from a NaCl gradient were screened by SDS–PAGE, and fractions containing the protein of interest (eluted at approximately 150 mM NaCl) were pooled. Following concentration of the pooled fractions to 1 mL using Amicon Ultra 3K molecular weight cutoff centrifuge concentrators (Millipore), the protein was loaded onto a Superdex 200 gel filtration column (GE Healthcare) equilibrated with 25 mM HEPES (pH 7.6), 200 mM NaCl, and 1 mM TCEP. Fractions containing the protein of interest were pooled, concentrated such that the final concentration was in the range of 250–500 \(\mu\)M, and stored at \(-80 \^\circ\)C. The protein concentrations were calculated by using an extinction coefficient of 26930 M\(^{-1}\) cm\(^{-1}\) at 280 nm for both YjiA and the YjiA mutants in 25 mM EDTA and 4 M guanidinium hydrochloride (GuHCl).\textsuperscript{20} Unless otherwise noted, electronic absorption measurements were taken on an Agilent 8453 spectrophotometer with a 1 cm path length cuvette. Each protein was analyzed by electrospray ionization mass spectrometry (ESI-MS) (Department of Chemistry, University of Toronto), and the observed molecular masses of the proteins are listed in Table S2 of the Supporting Information. All proteins were >90\% pure as estimated by Coomassie-stained SDS–PAGE quantified by using ImageJ [National Institutes of Health, Bethesda, MD (http://rsb.info.nih.gov/ij/)].

\textbf{Preparation of Proteins.} The reduced apoprotein was produced by incubating the protein with 10 mM EDTA and 20 mM TCEP in a Coy anaerobic glovebox at 4 \(^\circ\)C for 48 h. The TCEP and EDTA were removed by exhaustive dialysis into protein buffer [25 mM HEPES and 100 mM NaCl (pH 7.6)]. The absence of any metal bound to the protein was confirmed by a \(\text{[4-(2-pyridylazo)resorcinol (PAR) assay,}^{21}\) in which the protein was denatured with 4 M GuHCl and 50 \(\mu\)M PAR was added to the sample. The absorbance at 500 nm, due to the formation of the (PAR)\(_2\)Me(II) complex, was monitored and compared to a standard curve produced with 50 \(\mu\)M PAR in 4 M GuHCl and known metal concentrations. The free thiol content of the proteins was quantified via reaction with 5,5'-

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dithiobis(2-nitrobenzoic acid) (DTNB) in the presence of 6 M GuHCl and 1 mM EDTA. β-Mercaptoethanol was used as a standard, and the absorbance of the 5-mercapto-2-nitrobenzoic acid product was measured at 412 nm. Protein samples were >95% reduced after they had been treated with TCEP.

**Metal Binding and Stoichiometry.** For the difference spectra, individual samples containing 20 μM apo-YjiA in protein buffer and either 20 μM NiCl₂ or 20 μM CoSO₄ were prepared in the glovebox and incubated overnight at 4 °C. The electronic absorption spectrum was monitored between 250 and 500 nm and corrected by background subtraction at 600 nm. The difference spectrum for each metal was generated by subtracting the apo-YjiA spectrum from that of the metal-loaded protein.

Metal stoichiometry experiments were conducted by incubating 120 μM WT or mutant apo-YjiA with either 480 μM NiCl₂, 360 μM CoSO₄, or 600 μM ZnSO₄ overnight at 4 °C in the glovebox. Samples containing mixtures of metal were similarly prepared with 120 μM WT apo-YjiA and either 360 μM NiCl₂ and 360 μM CoSO₄, 600 μM ZnSO₄, or 480 μM NiCl₂, or 600 μM ZnSO₄, and 360 μM CoSO₄. Samples with GDP contained 1.5 mM GDP and 5 mM MgCl₂. Excess metal was removed by passing the protein through a PD10 gel filtration column (GE Healthcare) equilibrated with protein buffer in the glovebox. The protein concentration was subsequently measured in GuHCl and EDTA, as described above, and the metal content was determined via a PAR assay. In the case of the YjiA samples containing a mixture of metals, a potential cation of metal ions in solution was utilized. For HPLC analysis, at least 50 μg of protein was dried by centrifugation under vacuum, reconstituted with metal-free concentrated HCl (SeaStar Chemicals), and incubated overnight at 95 °C for protein hydrolysis. The sample was once again dried to remove HCl and reconstituted in 80 μL of Milli-Q water. This sample was injected onto a Dionex IonPak CSSA column, equilibrated with 7 mM pyridine-2,6-dicarboxylic acid, 66 mM KOH, 5.6 mM K₂SO₄, and 74 mM HCOOH, attached to a metal-free Dionex BioLC HPLC system. The metals were detected at 500 nm following postcolumn mixing with PAR.

**ESI-MS Measurements.** Before mass spectrometry experiments, 120 μM WT apo-YjiA was incubated with 600 μM ZnSO₄ overnight at 4 °C in the glovebox and then buffer exchanged into 10 mM ammonium acetate (pH 7.5) using two consecutive PD10 columns. Mass spectra were recorded on an AB/Sciex QStarXL mass spectrometer equipped with an ion spray source in the positive mode and a hot source-induced desolvation interface. Ions were scanned in the m/z 1000–2000 range with accumulation times of 1 s per spectrum, with no interscan time delay. The instrument parameters were as follows: ion source temperature, 200 °C; ion source gas, 50.0 psi; curtain gas, 50.0 psi; ion spray voltage, 5000 V; declustering potential, 60.0 V; focusing potential, 60.0 V; MCP (detector), 2200 V. The spectra were deconvoluted using the Bayesian protein reconstruction program included with Analyst QS (AB/Sciex) over a mass range of 30000–40000 Da, with a mass step of 1 Da.

**YjiA Metal Affinities.** To define the affinity of YjiA for Ni(II), 250 nM YjiA in protein buffer was titrated with a nickel atomic absorption standard solution and the absorbance at 340 nm was monitored using a 10 cm path length cuvette and a GBC Cintra 404 spectrophotometer. The apparent $K_d$ was calculated by determining the fractional saturation, $r$, and free nickel concentration, $[\text{Ni(II)}]_{\text{free}}$ by using eqs 1 and 2, respectively:

$$ r = \frac{[\text{YjiA} - \text{Ni(II)}]}{2[\text{YjiA}]_{\text{total}}} = \frac{A_{440} - A_{\text{min}}}{A_{\text{max}} - A_{\text{min}}} $$

$$ [\text{Ni(II)}]_{\text{free}} = \frac{[\text{Ni(II)}]_{\text{total}} - 2r[\text{YjiA}]_{\text{total}}}{2} $$

where $[\text{YjiA} - \text{Ni(II)}]$ is the concentration of the protein bound to Ni(II), $[\text{YjiA}]_{\text{total}}$ is the total protein concentration, $A_{440}$ is the absorbance at 340 nm for apo-YjiA, $A_{\text{max}}$ is the absorbance at 340 nm upon saturation, and $[\text{Ni(II)}]_{\text{total}}$ is the total Ni(II) concentration added to the sample. The resulting values were plotted as $r$ versus $[\text{Ni(II)}]_{\text{free}}$ and the data were fit to the Hill equation (eq 3):

$$ r = \frac{[\text{Ni(II)}]_{\text{free}}^n}{K_d^n + [\text{Ni(II)}]_{\text{free}}^n} $$

where $n$ is the Hill coefficient.

To determine the affinity of YjiA for Co(II), 5 μM apo-YjiA was titrated with CoSO₄ and the absorbance at 350 nm was monitored. The Co(II) $K_d$ was calculated in the same manner as for Ni(II), except that eqs 1 and 2 were adjusted to solve for a single metal site instead of two.

**YjiA Zincon Zn(II) Competitions.** To estimate the Zn(II) $K_d$ of YjiA, the competitor zincon was selected because of its ability to form a 1:1 complex with Zn(II) with a reported $K_d$ of 12.7 μM. Stocks of zincon were prepared in Milli-Q water. The affinity of zincon for Zn(II) was verified under our experimental conditions by titrating 400 nM zincon in protein buffer with increasing amounts of ZnSO₄. The absorbance at 620 nm was monitored using a 10 cm path length cuvette and a GBC Cintra 404 spectrophotometer. The $K_d$ was calculated by using the same method that was used for the Ni(II)—and Co(II)—YjiA titrations, where $n = 1$. Competition experiments were conducted by incubating 10 μM apo-WT or E37A/C66A/C67A YjiA with 140 μM zincon and various amounts of Zn(II). The samples were incubated overnight at 4 °C in the glovebox. The absorbance of zincon at 620 nm was monitored using a 2 mm path length cuvette.

**Analytical Gel Filtration Chromatography.** Samples containing 60 μM YjiA were incubated with the desired metal, GDP, or GTP at the indicated concentrations (Tables S2 and S3 of the Supporting Information) overnight at 4 °C in the glovebox. For samples containing GTP, the protein was preincubated with GTP for at least 2 h at 4 °C in the glovebox prior to injection onto the gel filtration column. All samples contained 25 mM HEPES (pH 7.6), 100 mM NaCl, and 5 mM MgCl₂. Apoprotein- and metal-containing samples were loaded onto a Superdex 200 10/300 analytical gel filtration column (GE Healthcare), pre-equilibrated with chelexed and filtered 25 mM HEPES (pH 7.6), 200 mM NaCl, and 5 mM MgCl₂. The column was calibrated with thyroglobulin (670 kDa), γ-globulin (158 kDa), ovalbumin (44 kDa), myoglobin (17 kDa), and vitamin B₁₂ (1.4 kDa) from Bio-Rad. The inclusion of metal in the standards did not affect their elution profiles (data not shown). Molecular masses were determined by plotting the log molecular masses of the standards versus the partition coefficient ($K_{av}$), where $K_{av} = (V_v - V_s)/(V_e - V_v)$ ($V_e$ represents the elution volume, $V_v$ the void volume, and $V_t$ the total column volume).
**Circular Dichroism (CD) Spectroscopy.** WT and mutant YjiA samples were prepared for CD spectroscopy by diluting the protein in Milli-Q water to a final concentration of approximately 10–20 μM in the glovebox. For metal titrations, ZnSO₄, CoSO₄, or NiCl₂ was added to the diluted samples and allowed to equilibrate overnight at 4 °C in the glovebox. All samples were analyzed on an Olis RSM 1000 spectropolarimeter with a capped 1 mm path length cuvette to minimize exposure to the air. Spectra were collected at 1 nm intervals over a spectral range of 200–260 nm with an integration time of 2 s and 2400 grating lines/nm. The final spectra obtained are averages of three scans. The observed ellipticity was converted into mean residue ellipticity (θ mre, in degrees square centimeters per decimole) using the following formula:26

$$\theta_{\text{mre}} = \frac{100 \theta}{\text{MW} \\ N \\ \text{protein}}$$

where MW is the molecular mass of the protein in daltons, N is the number of amino acids, θ is the observed ellipticity in degrees, [protein] is the protein concentration in grams per milliliter, and l is the path length. The secondary structure content was quantitated in the 200–240 nm range using K2D3.27 The CD spectra of the mutants were similar to that of the WT protein, demonstrating that these mutations do not affect the secondary structure of the protein (data not shown).

**GTPase Assay.** GTPase activity was determined by the Malachite Green assay for free phosphate adapted from Lanzetta et al.27 A series of 400 μL samples containing 0.5–2 μM WT or E37A/C66A/C67A YjiA in protein buffer supplemented with 5 mM Mg(II) and concentrations of GTP between 5 and 950 μM were incubated at 37 °C in the glovebox for 2.5–5 h. Controls without protein were prepared alongside the protein samples and received the same treatment. After incubation, 100 μL of the phosphate detection reagent (2.6 mM Malachite Green, 1.5% ammonium molybdate, and 0.2% Tween 20) was added to each sample. The samples were then mixed by being vortexed and incubated at room temperature for 3 min, after which sodium citrate was added to a final concentration of 3.5%. The samples were vortexed again and incubated at room temperature for 30 min prior to being plated on a 96-well plate, and the absorbance at 630 nm was measured with a Tecan Safire2 microplate reader. The amount of phosphate released was determined via a standard curve based on a phosphate standard (Molecular Probes). The data were analyzed by fitting to the Michaelis–Menten equation using OriginPro version 7.5. Samples containing metal were preincubated with the metal overnight prior to the assay [prepared as a stock of 10–40 μM protein with 100 μM Zn(II), 100 μM NiCl₂, or 250 μM CoSO₄ for WT or 225 μM Zn(II), 400 μM NiCl₂, or 800 μM CoSO₄ for the E37A/C66A/C67A triple mutant]. These stocks were then diluted to the final protein concentration of 0.5–2 μM for the assay in a buffer that contained 100 μM Zn(II), 100 μM NiCl₂, or 250 μM CoSO₄ for WT or 225 μM Zn(II), 400 μM NiCl₂, or 800 μM CoSO₄ for the E37A/C66A/C67A triple mutant.

**Crystallization and Zn(II) Soaks.** WT YjiA and E37A/C66A/C67A YjiA were crystallized at 25 °C by the hanging drop vapor diffusion technique. For WT YjiA, 1.5 μL of a protein solution [13.2 mg/mL in 100 mM NaCl and 25 mM HEPES (pH 7.6)] was mixed with 0.3 μL of 100 mM CaCl₂ and 1.5 μL of a precipitant solution [1.55 M (NH₄)₂SO₄ and 0.1 M HEPES (pH 6.9)] on a glass coverslip. The coverslip was sealed with grease over a reservoir containing 500 μL of the precipitant solution. Crystals appeared overnight and grew to maximal dimensions of ~300 μm x 50 μm x 50 μm after 3 days. Crystals of E37A/C66A/C67A YjiA were grown in the same fashion with the following modifications: the precipitant solution consisted of 1.3–1.35 M (NH₄)₂SO₄ and 0.1 M HEPES (pH 7.5–7.7), the protein concentration was 12.4 mg/mL, and the protein solution contained 1 mM TCEP in addition to the other components.

Because cocrystallization of WT YjiA with Zn(II) was unsuccessful, Zn(II)-soaked WT YjiA crystals were generated by incubating apo-WT crystals in the precipitant solution supplemented with 3 mM ZnSO₄ and 1 mM TCEP at 25 °C for 16 h. The crystals were then directly transferred to a cryogenic solution [1.5 M (NH₄)₂SO₄, 0.1 M HEPES (pH 6.9), 3 mM ZnSO₄, 1 mM TCEP, and 20% (v/v) glycerol], incubated for 5 s, and flash-frozen in liquid nitrogen. Zn(II)-soaked E37A/C66A/C67A YjiA crystals were generated in a similar fashion with the following modifications: the soaking solution contained an additional 20 mM CaCl₂ to better stabilize the crystals, the cryogenic solution consisted of 1.35 M (NH₄)₂SO₄, 0.1 M HEPES (pH 7.7), 20 mM CaCl₂, 3 mM ZnSO₄, 1 mM TCEP, and 20% (v/v) glycerol, and the crystals were transferred into the cryogenic solution in three steps at increasing glycerol concentrations and incubated in that solution for 20 s.

**Data Collection.** All Zn(II)-soaked YjiA crystals belong to space group P2₁ (a = 56 Å, b = 69 Å, c = 78 Å, and β = 104°) and contain two protomers in the asymmetric unit. All diffraction data were collected at 100 K.

Diffraction data for Zn(II)-soaked WT YjiA were collected at Stanford Synchrotron Radiation Laboratory beamline 9-2 in Portola Valley, CA, using a MARmosaic 325 CCD detector. An initial Zn(II)-soaked WT YjiA crystal was used for an X-ray fluorescence scan to verify the presence of bound Zn(II). On the basis of the fluorescence scan, a peak wavelength of 1.2827 Å (9665.5 eV) was determined using CHOOCB.28 A wavelength of 1.1808 Å (10500.0 eV) was chosen for the remote data set. Remote and peak data sets were collected on the same crystal in wedges of 30° in 1° oscillation steps. For each wedge, the corresponding images related by a 180° crystal rotation were collected immediately after completion of the wedge. An inflection data set was also collected on the same crystal, but not used for this study. Bijvoet mates were treated as separate reflections for both the remote and the peak data set.

Diffraction data for Zn(II)-soaked E37A/C66A/C67A YjiA were collected at Advanced Photon Source beamline 24ID-C in Argonne, IL, using a Pilatus 6M pixel detector. The crystal was annealed for 5 s prior to data collection.29 Then, a single data set was collected at a wavelength of 0.9795 Å (12658.0 eV) in 1° oscillation steps. For structure solution, Bijvoet mates were treated as symmetry-related reflections. For calculation of Zn anomalous difference Fourier maps of Zn(II)-soaked E37A/C66A/C67A YjiA, Bijvoet mates were treated as separate reflections.

All data were integrated using HKL2000 and scaled using Scalepack.30 The same reflections were marked for the free set of reflections for Zn(II)-soaked WT YjiA and for Zn(II)-soaked E37A/C66A/C67A YjiA. In addition, 5% of the reflections in the resolution ranges of 75–30 and 2.57–2.05 Å were included in the free set for the data of the triple mutant. All data...
collection statistics are summarized in Table S5 of the Supporting Information.

**Structure Building and Refinement.** The crystal structure of Zn(II)-soaked WT YjiA was determined at 2.57 Å resolution by molecular replacement with the crystal structure of native WT YjiA without water molecules [Protein Data Bank (PDB) entry I11] in PHASER, yielding a single solution. The output model was subjected to 20 cycles of simulated annealing in PHENIX to reduce model bias. The model was adjusted for any changes that had occurred through a cycle of model building in COOT and refinement in PHENIX. Initial difference electron density maps revealed the presence of five Zn(II) ions in the asymmetric unit. Zn(II) ions and water molecules were inserted in subsequent rounds of refinement.

The intermediate model of Zn(II)-soaked WT YjiA without water molecules or Zn(II) was used to determine the crystal structure of Zn(II)-soaked E37A/C66A/C67A YjiA to 2.05 Å resolution. The structure was determined by rigid body refinement in PHENIX followed by automated rebuilding of the structure using phenox.autobuild to reduce model bias. Initial difference electron density maps revealed the presence of all three mutations as well as the presence of three Zn(II) ions in the asymmetric unit. Subsequently, Zn(II) ions and water molecules were inserted, and the model was adjusted to account for the mutations as well as other changes through iterative cycles of model building in COOT and refinement in PHENIX.

Because of the superior quality of the Zn(II)-soaked WT YjiA data set (Table S5 of the Supporting Information), the model derived from this data set was then used to finalize the structure of Zn(II)-soaked WT YjiA. Following rigid body refinement of the mutant structure into the WT YjiA data in PHENIX, additional Zn(II) ions were inserted and the structure was completed through iterative cycles of model building in COOT and refinement in PHENIX.

For both structures, the two protomers in the asymmetric unit were initially restrained by strict non-crystallographic symmetry (NCS) parameters. In advanced stages of refinement, NCS restraints were loosened for all residues and removed for residues involved in crystal contacts or Zn(II) binding. Side chains with limited electron density were truncated at the last atom with visible electron density. Zn(II) coordination was loosely restrained to average bond lengths observed in previously determined structures of proteins with Zn(II) ions bound. The final stages of refinement for both structures involved TLS parametrization using two TLS groups per protomer, as determined by the TLSMD server. The locations of Zn(II) ions were verified by Zn anomalous difference Fourier maps calculated from the peak data set (WT YjiA) or the remote data set (E37A/C66A/C67A YjiA) using FFT, which is part of the CCP4 program suite.

Crystallographic refinement of both structures yielded models that possess excellent stereochemistry and small root-mean-square deviations from ideal values for bond lengths and bond angles. Because of the presence of pseudo-translational symmetry in the data, the crystallographic R factors are relatively high, but within an acceptable range. Refinement statistics are summarized in Table S5 of the Supporting Information. The final model of WT YjiA contains all 318 residues of YjiA in protomer A and residues 2–318 in protomer B. The final model of E37A/C66A/C67A YjiA contains all 318 residues in both protomers. The models were validated using simulated annealing composite omit maps calculated in CNS. The geometries of the final models were analyzed with Molprobity. Figures were generated using Pymol.

**RESULTS**

**YjiA Binds Zn(II), Ni(II), and Co(II) with Micromolar Affinities.** The presence of a CXXC motif in YjiA, conserved in all COG0523 proteins, led to the proposal that the protein can bind metals, but experimental evidence was lacking. Because of the homology of YjiA to G3E NTPases known to be involved in Zn(II) (YeiR and YciC), Ni(II) (UreG and HypB), or Co(II) (CobW) homeostasis, these three metals were the focus of this investigation. To determine whether YjiA can bind these metals and, if so, how much, 120 μM apo-YjiA was incubated with either 480 μM NiCl₂, 360 μM CoSO₄, or 600 μM ZnSO₄, and excess metal was removed by gel filtration chromatography. Subsequent metal analysis via a PAR assay revealed roughly stoichiometric cobalt, two Ni(II) ions, or four Zn(II) ions bound per monomer (Table 1).

| Table 1. Stoichiometry of Binding of Metal to WT and Mutant YjiA |
|-----------------|-----------------|-----------------|
| WT with GDP     | 1.3 ± 0.1       | 0.6 ± 0.2       |
| E37A/C66A/C67A  | 0.5 ± 0.2       | 0.1 ± 0.1       |
| E42A/C66A/C67A  | 1.2 ± 0.2       | 0.2 ± 0.1       |

*480 μM CoSO₄ was incubated with either 480 μM NiCl₂, 360 μM CoSO₄, or 600 μM ZnSO₄ overnight at 4 °C in an anaerobic glovebox. Excess metal was removed by passing the proteins over a PD10 gel filtration column, and the bound metal was detected via a PAR assay. In samples containing GDP, 1.5 mM GDP and 5 mM MgCl₂ were included. The data listed are average values and standard deviations of the number of metal ions bound per protein monomer from at least three independent experiments.*

After incubation of YjiA with 1 equiv of nickel or cobalt, broad charge-transfer bands appear in the electronic absorption spectrum region of 250–400 nm (Figure 1A). The difference spectra, obtained by subtracting the signal of apo-YjiA from that of the protein loaded with 1 equiv of Ni(II) or Co(II) (Figure 1A), reveal a peak maximum at 280 nm and a broad shoulder around 340 or 350 nm for nickel and cobalt, respectively. The 280 nm and shoulder absorptions for both metals can be attributed to Cyg-S⁻ → Ni(II)/Co(II) ligand-to-metal charge transfer (LMCT) indicating the presence of at least one thiolate in the coordination sphere of both metals.

The LMCTs for Ni(II) and Co(II) can be used to monitor metal binding by YjiA. A titration of 5 μM apo-YjiA with CoSO₄ yields an apparent Kₐ of (2.0 ± 0.1) × 10⁻⁵ M and a Hill coefficient n of 1.7 ± 0.2 (Figure 1B). This Hill coefficient suggests cooperativity in cobalt binding to YjiA, a surprising result given that the protein binds only 1 equiv of cobalt. This discrepancy may be explained by the ability of cobalt to induce oligomerization in the protein (see below). The titration of YjiA with Ni(II) is more complicated because of the 2:1 Ni(II):YjiA stoichiometry as the titration may be monitoring either a single site or both Ni(II) sites simultaneously. The electronic absorption spectrum demonstrates that at least one site is spectrascopically active, but the mutants described below suggest that the second Ni(II) site does not yield a discernible
electronic absorption signal. The appearance of a LMCT upon the addition of substoichiometric amounts of nickel to YjiA indicates that the second site is not significantly tighter than the spectroscopically active site. Thus, the second nickel site has either a similar or a lower affinity than the spectroscopically active site. If this site is much weaker, then the nickel titration in Figure 1B represents only the spectroscopically active site and yields an apparent $K_d$ of $(3.9 \pm 0.3) \times 10^{-6} \text{M}$ with a Hill coefficient $n$ of $1.2 \pm 0.2$ for this one site. If the two sites have similar affinities, an apparent average $K_d$ for the two sites can be calculated from the Ni(II) titration to be $(3.7 \pm 0.3) \times 10^{-6} \text{M}$ ($n = 1.2 \pm 0.3$).

Because of the spectroscopically quiet nature of Zn(II), a metallochromic indicator, zincon, was used to estimate the affinity of Zn(II) for YjiA. The $K_d$ of zincon for Zn(II) was determined to be $10 \pm 1 \mu\text{M}$ under our buffer conditions (data not shown), in agreement with the reported $K_d$ of 12.7 $\mu\text{M}$. In the absence of protein, metal loading of 140 $\mu\text{M}$ zincon is complete upon addition of 140 $\mu\text{M}$ Zn(II) (Figure 2). Upon the inclusion of 10 $\mu\text{M}$ YjiA, 180 $\mu\text{M}$ Zn(II) is required to observe saturation of the zincon (Figure 2), demonstrating competition by YjiA for Zn(II) and consistent with the measured Zn(II) stoichiometry of four ions per monomer. Furthermore, binding of Zn(II) to zincon is not observed until more than 20 $\mu\text{M}$ metal has been added, suggesting that YjiA has two Zn(II) sites with a $K_d$ significantly tighter than that of zincon ($K_d < 10 \mu\text{M}$), and that the remaining two Zn(II) sites have affinities comparable to that of zincon ($K_d \sim 10 \mu\text{M}$). Finally, to confirm the Zn(II) stoichiometry, ESI-MS of YjiA incubated with excess Zn(II) was performed (Figure S1 of the Supporting Information). The mass spectrum revealed up to four Zn(II) ions bound to the protein, with two sites completely filled and two partially filled, consistent with the affinities of the sites estimated in the competition experiment.

**Ni(II), Co(II), and Zn(II) Share a Common Site.** To determine if the binding sites of the different transition metals overlap, apo-YjiA was incubated simultaneously with two types of metal, followed by HPLC metal analysis to identify which metals are bound. When YjiA was incubated with both CoSO$_4$ and NiCl$_2$, 1.9 $\pm$ 0.3 Ni(II) ions and 0.15 $\pm$ 0.01 Co(II) ions were detected bound to the protein, suggesting that Co(II) shares a site with Ni(II) and that Ni(II) can outcompete Co(II) for this site. Similar experiments conducted with ZnSO$_4$ and either NiCl$_2$ or CoSO$_4$ demonstrate a selectivity for Zn(II) as 3.6 $\pm$ 0.3 Zn(II) ions and 4.0 $\pm$ 0.4 Zn(II) ions per YjiA were detected, respectively, with no detectable Ni(II) or Co(II).

**The GTPase Domain Contains the Common Metal-Binding Site.** A sequence alignment of YjiA with other G3E GTPases demonstrates the presence of conserved putative metal-binding residues between the Walker A and Walker B motifs (Figure 3A). In the COG0523 subfamily, this conserved sequence corresponds to C$^{\text{Glu}}$XXC$^{\text{Cys}}_G$ of YjiA. Furthermore, examination of the published crystal structure of apo-YjiA$^{17}$
revealed two nearby glutamates, Glu37 and Glu42, which could also serve as metal-binding residues (Figure 3B). To investigate the role of these glutamates as well as the CXCC motif in metal binding, we generated the E37A/C66A/C67A and E42A/C66A/C67A triple mutants of YjiA and characterized their metal-binding capabilities. Upon addition of Ni(II) or Co(II) to either triple mutant, the absorbance spectra lacked the LMCTs at 340 or 350 nm, respectively, suggesting that Cys66 and/or Cys67 are the source of these signals (data not shown). Furthermore, both triple mutants exhibited weakened Co(II) and Zn(II) binding, but the Zn(II) stoichiometry was unchanged (Table 1). However, the previous observation that Zn(II) competes with the other metals (see above), suggesting overlapping sites, led us to investigate further the Zn(II) coordination of E37A/C66A/C67A YjiA by using a zincon titration. This result is consistent with the maintained Zn(II) stoichiometry of the mutant protein but indicates that the mutations have significantly weakened the affinities of two of the metal ions and suggests that there is at least one Zn(II)-binding site involving some combination of Glu37 and Glu42, which could also serve as metal-binding residues (Figure 3B). To investigate the role of these glutamates as well as the CXCC motif in metal binding, we generated the E37A/C66A/C67A and E42A/C66A/C67A triple mutants of YjiA and characterized their metal-binding capabilities. Upon addition of Ni(II) or Co(II) to either triple mutant, the absorbance spectra lacked the LMCTs at 340 or 350 nm, respectively, suggesting that Cys66 and/or Cys67 are the source of these signals (data not shown). Furthermore, both triple mutants exhibited weakened Co(II) and Zn(II) binding, but the Zn(II) stoichiometry was unchanged (Table 1). However, the previous observation that Zn(II) competes with the other metals (see above), suggesting overlapping sites, led us to investigate further the Zn(II) coordination of E37A/C66A/C67A YjiA by using a zincon titration. Unlike the WT protein, no initial plateau region upon adding metal to the E37A/C66A/C67A and E42A/C66A/C67A mutants, although the impact on quaternary structure was diminished compared to the impact in the WT protein (Table S4 of the Supporting Information). No Major Secondary Structure Changes. The CD spectra of apo-WT YjiA (Figure S2 of the Supporting Information) is indicative of a mixed α/β protein, in close agreement with the published crystal structure. Upon addition of Co(II), Zn(II), or Ni(II), minor changes in the CD spectrum were detected, but the calculated percentage of secondary structure content was unchanged (Figure S2 of the Supporting Information), indicating that YjiA does not undergo any dramatic secondary structure changes upon binding metal. Analytical gel filtration chromatography was used to probe the oligomeric state of apo-YjiA, as well as the metal- and nucleotide-bound species. In the apo form, the protein elutes from the column as a monomer (Figure 4 and Table S3 of the Supporting Information). The calculated molecular mass of the monomeric apo-YjiA species from the gel filtration experiments (30.0 ± 1.5 kDa) is smaller than the predicted molecular mass of 35.7 kDa, suggesting a compact protein structure. Upon the addition of 2 equiv of Ni(II), Co(II), or Zn(II), the protein oligomerizes, forming a small amount of dimer as well as a larger oligomeric species (Figure 4). The protein also dimerizes in the presence of GDP or GTP, but to a lesser degree than with metal (Figure 4). Some oligomerization was also observed upon adding metal to the E37A/C66A/C67A and E42A/C66A/C67A mutants, although the impact on quaternary structure is diminished compared to the impact in the WT protein (Table S4 of the Supporting Information). Metal Binding Affects YjiA GTPase Activity. YjiA was previously shown to bind GTP and, on the basis of its GTPase motifs, is predicted to possess GTP hydrolysis (GTPase) activity. In agreement with this prediction, the apo-WT protein has low GTPase activity, with a $k_{cat}/K_m$ of $14 ± 9 M^{-1} s^{-1}$ (Table 2). To determine whether metal binding modulates the GTPase activity, we incubated apo-YjiA with Ni(II), Co(II), or Zn(II) prior to analysis. The addition of Co(II) slightly
disrupts GTPase activity, reducing the $k_{cat}/K_m$ to $2.3 \pm 0.8 \text{ M}^{-1} \text{s}^{-1}$. The presence of Ni(II) significantly diminishes the GTPase activity of YjiA, such that substrate saturation was not observed with up to 950 μM GTP, indicating a significantly weaker $K_m$ versus that of the apoprotein. Attempts to use higher GTP concentrations were unsuccessful because of the departure from the thermal response region of the assay (data not shown). Zn(II) also inhibits the enzyme, and in this case, no activity was detectable in the presence of 100 μM ZnSO₄ (Table 2). The results of these GTPase assays prompted experiments that aimed to examine if there was a connection between GDP results of these GTPase assays prompted experiments that even though it was added at lower concentrations than Co(II) and Ni(II). While the E37A/C66A/C67A triple mutant has reduced affinity for all three metals (see above), it appears that Zn(II) binds to at least some sites in YjiA with higher affinity than Co(II) and Ni(II) do, and that this affinity remains sufficiently high to cause inhibition at the concentrations used. To probe whether this inhibition of GTPase activity occurs by direct Zn(II) binding to the active site or in an allosteric fashion by Zn(II) binding to other metal sites on YjiA, we conducted a structural analysis of Zn(II)-bound YjiA.

Zn(II) Binding Induces a Space Group Transition in YjiA Crystals. We determined the crystal structure of Zn(II)-bound WT YjiA to 2.57 Å resolution after soaking apo-WT YjiA crystals with ZnSO₄ (Figure 5). Surprisingly, the crystals underwent a space group transition upon soaking. While uns soaked (apo) WT YjiA crystals belong to space group C2 with one protomer in the asymmetric unit, the space group is P2₁, after soaking and features two protomers in the asymmetric unit (for a more detailed discussion of space group assignment, see the Supporting Information). The arrangement of molecules in the crystal lattices of Zn(II)-soaked WT YjiA and apo-WT YjiA, however, is nearly identical despite the different space groups (Figure S3 of the Supporting Information). It seems that Zn(II) soaking induced a small shift in the crystal lattice, causing a reduction in symmetry from C2 to P2₁ while leaving the overall crystal lattice intact.

Unfortunately, we have not been able to generate nucleotide-bound structures of YjiA itself. While YjiA exhibits the canonical GTPase motifs, these motifs are not well-ordered in the structure and the phosphate-binding Walker A motif in particular seems to exhibit multiple conformations (Figure S4 of the Supporting Information). Likely, nucleotide binding will result in a conformational change and depend on the presence of an effector protein, consistent with our unsuccessful attempts to generate nucleotide-bound structures.

YjiA Binds Zn(II) in Four Distinct Sites. Analysis of the electron density and of Zn anomalous difference Fourier maps revealed the presence of Zn(II) ions in four distinct types of sites: a “bridging site” that is located at the interface of the two YjiA protomers in the asymmetric unit (site B in Figure SA), an “internal site” (site C in Figure SA), and two types of “surface

![Image](https://example.com/image.png)

Figure 4. Effect of metal on the quaternary structure of YjiA. In the absence of metal, 60 μM YjiA elutes at a volume corresponding to a monomer (solid black line). The addition of 2 equiv of NiCl₂ (solid red line), CoSO₄ (dashed black line), or ZnSO₄ (solid blue line) results in the formation of dimeric and oligomeric species. Incubation of YjiA with 2 equiv of GDP (dashed red line) or GTP (dashed blue line) followed by chromatography with 400 μM nucleotide in the running buffer resulted in only a small portion of dimeric protein. The chromatographic traces (monitored at 280 nm) are representative data sets from experiments with a Superdex 200 10/300 analytical column equilibrated with 25 mM HEPES (pH 7.6), 200 mM NaCl, and 5 mM MgCl₂. The ticks at the top of the graph denote the elution volumes of the protein standards. From left to right, the identities of the standards (and their molecular masses) are thyroglobulin (670 kDa), γ-globulin (158 kDa), ovalbumin (44 kDa), myoglobin (17 kDa), and vitamin B₁₂ (1.4 kDa), respectively.

Table 2. Kinetics of GTP Hydrolysis by WT and E37A/C66A/C67A YjiA

<table>
<thead>
<tr>
<th></th>
<th>$k_{cat}$ (s⁻¹)</th>
<th>$K_m$ (M)</th>
<th>$k_{cat}/K_m$ (M⁻¹ s⁻¹)</th>
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<tbody>
<tr>
<td>WT</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>apo</td>
<td>$(6 \pm 2) \times 10^{-1}$</td>
<td>$(5 \pm 3) \times 10^{-1}$</td>
<td>14 ± 9</td>
</tr>
<tr>
<td>Co(II)</td>
<td>$(5 \pm 2) \times 10^{-4}$</td>
<td>$(2.3 \pm 0.7) \times 10^{-4}$</td>
<td>2.3 ± 0.8</td>
</tr>
<tr>
<td>Ni(II)</td>
<td>$(1.1 \pm 0.1) \times 10^{-3}$</td>
<td>$(1.0 \pm 0.7) \times 10^{-4}$</td>
<td>6 ± 3</td>
</tr>
<tr>
<td>Zn(II)</td>
<td>$(6 \pm 1) \times 10^{-4}$</td>
<td>$(1.7 \pm 0.1) \times 10^{-5}$</td>
<td>78 ± 9</td>
</tr>
<tr>
<td>E37A/C66A/C67A</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>apo</td>
<td>$(5 \pm 1) \times 10^{-4}$</td>
<td>$(1.0 \pm 0.7) \times 10^{-4}$</td>
<td>6 ± 3</td>
</tr>
<tr>
<td>Co(II)</td>
<td>$(1.1 \pm 0.1) \times 10^{-3}$</td>
<td>$(1.7 \pm 0.1) \times 10^{-5}$</td>
<td>78 ± 9</td>
</tr>
<tr>
<td>Ni(II)</td>
<td>$(6 \pm 1) \times 10^{-4}$</td>
<td>$(3 \pm 1) \times 10^{-4}$</td>
<td>2 ± 1</td>
</tr>
<tr>
<td>Zn(II)</td>
<td></td>
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All GTPase assays were conducted with 0.5–2 μM WT or E37A/C66A/C67A YjiA in protein buffer supplemented with 5 mM MgCl₂. Samples containing metal were preincubated with either zinc, cobalt, or nickel overnight at 4 °C in an anaerobic glovebox. The amount of released phosphate was detected using a modified Malachite Green assay. The data listed are average values from at least three independent experiments. aWeak hydrolysis (see the text for details). bNo measurable hydrolysis (see the text for details).
sites” (sites D and E in Figure 5A). In total, the two YjiA protomers in the asymmetric unit could bind seven Zn(II) ions: two in the internal site, one in the bridging site, and four in surface sites (Figure 5A). The asymmetric unit, however, contains only five Zn(II) ions because all four surface sites are involved in crystal lattice contacts and each of those bound Zn(II) ions is shared with a neighboring asymmetric unit molecule.

**The Internal Site Is Located near the GTPase Active Site.** The internal site is located in the N-terminal region of a YjiA protomer, with the side chains of E37, E42, and C66 coordinating Zn(II) (Figure 5C). Although hard to resolve in the electron density at this resolution, a fourth coordination site on Zn(II) could be occupied by a water molecule, which would give rise to a tetrahedral coordination geometry. E37 and E42 are located in a loop region that undergoes a conformational change upon Zn(II) binding to bring the side chain of E42 into the proximity of the bound Zn(II) (Figure 5C). Furthermore, the region around C66 undergoes a slight shift, bringing C66 closer to the Zn(II) ion and rearranging the subsequent loop region significantly (Figure 5C).

It is clear that the internal Zn(II)-binding site is in the neighborhood of the nucleotide-binding site (Figure 5C), as demonstrated by a structural alignment with the GTP analogue-bound structure of *Methanocaldovoccus jannaschii* HypB (PDB entry 2HF8,18 GTPγS shown with cyan carbons and Mg(II) shown as an orange sphere). (D and E) Close-up views of the two types of surface sites, located at crystal contacts between YjiA protomers in the asymmetric unit and crystallographically related molecules. Zn(II) is coordinated by the side chains of E167 and/or H170 from one molecule and H187 from the other molecule.
The Bridging Site and the Surface Site Connect YjiA Protomers. As mentioned briefly above, the bridging site is located between two YjiA protomers on a noncrystallographic 2-fold symmetry axis, with Zn(II) coordinated by the side chains of E74 and H114 from both protomers in an approximate tetrahedral geometry (Figure 5B). Binding to this site likely disrupted the C2 symmetry of apo-WT YjiA crystals, as the bridging site would be located on a crystallographic axis in the C2 lattice. Analysis of the resulting dimer interface with the PISA server50 indicates that this interface alone is not as extensive as is typically observed in physiologically relevant dimers, with an average buried surface area of 1244 Å² and a low complex formation significance score of 0.132 (on a scale of 0–1 with increasing relevance to complex formation in solution), not including the contributions from Zn(II). Thus, it seems unlikely that apo-YjiA would be a dimer in solution, in agreement with our analytical gel filtration chromatography results. Zn(II) binding, however, could strengthen this interface and thereby induce dimerization and oligomerization of YjiA.

All four surface Zn(II) sites are located at crystal contacts at the interfaces of YjiA protomers (Figure 5D,E). The Zn(II) ions in these sites are coordinated by H170 and/or E167 from one protomer and H187 from a protomer in the neighboring asymmetric unit, or vice versa. These sites exhibit a significant amount of disorder and high crystallographic B factors, indicating that the Zn(II) ions are not bound at high occupancy within the crystal lattice. There are also some differences in the exact coordination spheres of the surface sites, providing a further rationale for the asymmetry that led to the space group transition upon Zn(II) binding. Most likely, Zn(II) binding affects the individual protomers in a slightly different manner, thus causing breakdown of the crystallographic symmetry. It is unclear whether these sites would be occupied by Zn(II) ions in solution or whether binding to these sites occurred adventitiously, after the protein ligands were brought into proximity by crystallization. These sites, however, provide an additional explanation for the observed metal-induced oligomerization of YjiA, as they could stabilize higher-order oligomers.

**DISCUSSION**

The mechanisms through which cells ensure the proper allocation and trafficking of metal ions are not well understood. The COG0523 subfamily of GTPases represents a diverse group of proteins, and on the basis of homology to known metallochaperones in the other G3E subfamilies and comparative genomics analysis, it was suggested that they play a role in metal trafficking or some other metal homeostasis process.6,51 This study of YjiA lends credence to this proposal. In particular, we demonstrate that YjiA can bind transition metals and provide biochemical and structural details about this interaction. Furthermore, the impact of metal binding on the GTPase activity of YjiA suggests that metal ions play a regulatory function.

YjiA belongs to subgroup 9 of the COG0523 subfamily and was predicted to bind metals via a conserved CXCC motif.17 As with the other members of this subgroup, YjiA does not have an assigned function, but there is a growing body of evidence that suggests a role for YjiA in the cellular response to carbon starvation.6,52–54 Within the genomic context, yjiA is found downstream of yjiY, a homologue of carbon starvation protein CstA, and yjiX, a small cytosolic protein of unknown function.6

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**Figure 6.** E37 or E39 of YjiA could coordinate nucleotide-associated Mg(II). GTP analogue-bound HypB (cyan carbons, PDB entry 2HF818) is superimposed onto the structure of Zn(II)-bound WT YjiA (green carbons). Bound GTPγS is shown with cyan carbons, and GTPγS-associated Mg(II) is shown as an orange sphere. D75 of HypB is involved in Mg(II) coordination. Both E37 and E39 of YjiA are in the proximity of the modeled Mg(II), as indicated by the purple dashed lines, raising the possibility that these residues are involved in Mg(II) coordination in YjiA. Notably, E37 is also involved in binding Zn(II) (purple sphere). E39 was truncated to the Cγ atom because of a lack of electron density for the side chain.
These three genes are transcribed as a single transcript, which is believed to be under the control of one of the central regulators in the carbon catabolism pathway, cyclic AMP receptor protein (CRP). Furthermore, yjiA is downregulated by the noncoding RNA molecule Spot 42 that participates in a feed-forward loop to help enact catabolite repression in E. coli. Further studies will be needed to establish what role YjiA plays in the carbon starvation response and the function of metals in this pathway. Here, we present a characterization of the metal binding capabilities of YjiA. In particular, our data identify a Zn(II)-binding site created by the side chains of E37, E42, and C66, the latter from the predicted CxxCxxC metal-binding motif. Solution studies of WT and mutant proteins suggest that Ni(II) and Co(II) also bind to at least a subset of the same ligands, although the coordination is likely to be different. Whether the other conserved cysteines in this motif have a functional role in binding metals or performing downstream effects remains to be determined. The relative metal affinities mirror the Irving-Williams series of small-molecule chelators for divalent metal ions, suggesting that the protein structure is not enforcing strong metal selectivity. Furthermore, at this time, it is not clear which metals are physiologically relevant for YjiA function. Many of the COG0523 subfamilies were linked to zinc pathways because of the presence of zinc-regulated genetic elements, and deletions of B. subtilis yeIR or E. coli yjiA sensitize the bacteria to zinc-deficient conditions. However, a similar phenotype was not observed in a ΔyjiA strain of E. coli, and binding sites for zinc metalloregulators were not detected in the genetic neighborhoods of subgroup 9.7,16 On the other hand, although nickel is required in E. coli as a cofactor for several [NiFe]-hydrogenase isofoms,6 the uptake of this metal is restricted to anaerobic growth and there is no evidence that Ni(II) suppresses GTPase activity to different rates determined in vitro. In vitro, the isolated GTPases lack additional partner proteins and cofactors that may stimulate GTPase activity in vivo. Furthermore, the addition of SlyD, a protein involved in the nickel insertion stage of [NiFe]-hydrogenase biosynthesis, enhances the GTPase activity of E. coli HypB. By analogy, it is possible that YjiA requires an additional factor to maximally activate GTP hydrolysis. A possible candidate is the small cytosolic protein YjIX that is co-transcribed with YjiA.

In addition to the internal metal site, YjiA also binds multiple Zn(II) ions at interface sites between separate YjiA molecules, which may provide an explanation for the metal-induced oligomerization observed by analytical gel filtration chromatography. It is also possible, however, that rearrangements around the internal metal site could create interfaces for oligomerization. Large-scale metal-induced rearrangements would be masked in our crystal soaking experiments because the interfaces between YjiA molecules are already preformed in the crystal lattice. This scenario would explain how Co(II) can mediate changes in quaternary structure, given that the solution analysis indicates that only a single Co(II) ion binds to YjiA at the internal site. Furthermore, this model is consistent with the observation that mutation of two of the internal site ligands diminishes oligomerization induced by Ni(II) and Zn(II). Finally, there may be other metal sites on YjiA not identified by crystallography. A recent bioinformatics analysis of the apo-YjiA crystal structure suggested two surface-exposed sites with potential for metal binding, composed of H23, E27, and H29 as well as D52, D79, and D82, respectively. We do not observe bound metal in either of these two sites, and E27 and D52 do not appear to be in the correct positions for metal binding. We note, however, that binding to these sites could rely on structural rearrangements that are prohibited by the crystal lattice. Further studies will be necessary to probe the contribution of these residues to metal binding as well as the functional relevance of metal ions bound on the surface of YjiA.

Although characterization of the E37A/C66A/C67A mutant provides information about the structure and impact of the internal Zn(II) site, several observations require additional explanation. The ability of this triple mutant to bind four Zn(II) ions and to oligomerize in the presence of Zn(II), albeit to a lesser degree than WT YjiA, suggests that mutation of these residues is not sufficient to completely disrupt Zn(II) binding. This conclusion, supported by the zinc competition illustrating a decreased affinity for Zn(II), is surprising because the crystal structure depicts a Zn(II) coordination sphere comprised of E37, E42, and C66, such that mutation of two of these three protein ligands should be sufficient to stop metal binding. Structural rearrangement of this site to incorporate other residues as ligands, such as the nearby E39, could serve as...
a possible explanation. Such a rearrangement may not be observed in the E37A/C66A/C67A YjiA crystal structure because of the aforementioned interface differences in the crystal prior to Zn(II) binding. Possible binding to a rearranged site in the E37A/C66A/C67A mutant may also explain the GTPase inhibition in the presence of Zn(II). An alternative explanation for the inhibition of the GTPase activity by Zn(II) is the involvement of additional allosteric sites, but evidence to support this latter explanation is needed.

Altogether, the results presented here provide a detailed biochemical and structural picture of the metal binding properties of a COG0523 GTPase. A connection between members of this subfamily of GTPases and intracellular metal pathways is emerging. This study supports the model in which metals play a role in regulating the proteins’ enzymatic activities and presents a glimpse into how this metal-mediated regulation may occur. Furthermore, as noted earlier, subgroup 9 of the COG0523 GTPases is hypothesized to be involved in carbon starvation, and the metal binding activities of YjiA offer a tantalizing hint about the interplay of metal homeostasis with other cellular processes.

■ ASSOCIATED CONTENT

* Supporting Information
A discussion of the space group transition in the YjiA crystals upon Zn(II) binding, tables with primer sequences, representative reconstructive ESI-MS results, summary of gel filtration chromatography results, crystallographic data collection and refinement statistics, circular dichroism spectra, and figures of the crystal structure of WT and mutant YjiA with Zn(II). This material is available free of charge via the Internet at http://pubs.acs.org.

Accession Codes
The atomic coordinates for Zn(II)-soaked WT YjiA and Zn(II)-soaked E37A/C66A/C67A YjiA have been deposited in the PDB as entries 4IXM and 4IXN, respectively.

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Notes
The authors declare no competing financial interest.

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■ ABBREVIATIONS

CD, circular dichroism; DTNB, 5,5′-dithiobis(2-nitrobenzoic acid); ESI-MS, electrospray ionization mass spectrometry; EDTA, ethylenediaminetetraacetic acid; GuHCl, guanidinium hydrochloride; HPLC, high-pressure liquid chromatography; IPTG, isopropyl β-D-thiogalactopyranoside; LMCt, ligand-to-metal charge transfer; P-loop, phosphate-binding loop; PAR, 4-(2-pyridyldazo)resorcinol; PMSF, phenylmethanesulfonyl fluoride; rmsd, root-mean-square deviation; TCEP, tris(2-carboxyethyl)phosphine; WT, wild type.

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Biochemistry


