A Novel Zinc Exporter CtpG Enhances Resistance to Zinc Toxicity and Survival in *Mycobacterium bovis*

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**ABSTRACT**  Zinc is a microelement essential for the growth of almost all organisms, but it is toxic at high concentrations and represents an antimicrobial strategy for macrophages. *Mycobacterium tuberculosis* and *Mycobacterium bovis* are two well-known intracellular pathogens with strong environmental adaptability, including zinc toxicity. However, the signaling pathway and molecular mechanisms on sensing and resistance to zinc toxicity remains unclear in mycobacteria. Here, we first report that P1B-type ATPase CtpG acts as a zinc efflux transporter and characterize a novel CmtR-CtpG-Zn\(^{2+}\) regulatory pathway that enhances mycobacterial resistance to zinc toxicity. We found that zinc upregulates ctpG expression via transcription factor CmtR and stimulates the ATPase activity of CtpG. The APC residues in TM6 is essential for CtpG to export zinc and enhance *M. bovis* BCG resistance to zinc toxicity. During infection, CtpG inhibits zinc accumulation in the mycobacteria, and aids bacterial survival in THP-1 macrophage and mice with elevated inflammatory responses. Our findings revealed the existence of a novel regulatory pathway on mycobacteria responding to and adapting to host-mediated zinc toxicity.

**IMPORTANCE**  Tuberculosis is caused by the bacillus *Mycobacterium tuberculosis* and is one of the major sources of mortality. *M. tuberculosis* has developed unique mechanisms to adapt to host environments, including zinc deficiency and toxicity, during infection. However, the molecular mechanism by which mycobacteria promote detoxification of zinc, and the associated signaling pathways remains largely unclear. In this study, we first report that P\(_{1\beta}\)type ATPase CtpG acts as a zinc efflux transporter and characterize a novel CmtR-CtpG-Zn\(^{2+}\) regulatory pathway that enhances mycobacterial resistance to zinc toxicity in *M. bovis*. Our findings reveal the existence of a novel excess zinc-triggered signaling circuit, provide new insights into mycobacterial adaptation to the host environment during infection, and might be useful targets for the treatment of tuberculosis.

**KEYWORDS**  *Mycobacterium bovis*, CtpG, zinc toxicity, intracellular survival, P1B-type ATPase

Tuberculosis (TB) is caused by the bacillus *Mycobacterium tuberculosis* (Mtb) and is a leading cause of death and disability worldwide. Vaccination with the Bacillus Calmette-Guerin (BCG) and chemotherapy are the common strategies for TB control (1). Mtb is a facultative intracellular pathogen able to survive in an infected host for decades, with little or no replication (2). Thus, the pathogen must exist various strategies to stand with diverse hostile environment within the host, including a toxic environment and nutritional immunity by depriving bacteria from some metals (Fe, Mn) or poisoning the bacteria with others (Cu, Zn) (3–6).

Metals are essential components for nearly all forms of life. Indeed, approximately one-half of all enzymes in organism require a metal cofactor to fulfill their biological activity (7). However, it is toxic at high concentrations and represents a cell-autonomous...
immunity strategy (3, 8–11). During infection, the concentrations of Cu$^{2+}$, Fe$^{2+}$, and Zn$^{2+}$ inside phagosomes infected with intracellular pathogens increase between 1 and 24 h postinfection (3, 12), until antibacterial levels of Cu$^{2+}$ and Zn$^{2+}$ are reached (8). High concentrations of Cu$^{2+}$ generates a ROS response via oxidative stress, protein denaturation, inactivation of enzymes by substitution of other metal cofactors, and membrane destabilization (3, 13, 14). Zn$^{2+}$, unlike Cu$^{2+}$, is a redox-inactive nutrient metal. It can bind to non-homologous proteins at high concentrations, then prevents the protein from obtaining the required metal ion and cause protein dysfunction (15, 16). For example, zinc competitively inhibits manganese binding to an importer in S. pneumoniae and the glycolytic enzymes phosphofructokinase and glyceraldehyde-3-phosphate dehydrogenase in S. pyogenes (17–19). In response, bacteria have evolved elegant strategies, including extensive transporters, transcription factors, and ligands, to counteract the host defense (3, 20–22). Until now, three types of zinc export systems that protect cells from high concentrations of zinc have been identified: RND multidrug efflux transporters, P-type ATPases, and cation diffusion facilitators (CDF) (23, 24). In addition, zinc homeostasis is achieved by export systems and uptake systems which are regulated by zinc-responsive regulator ZntR and zinc uptake repressor Zur, respectively (23, 25). In the presence of zinc, the ion stimulates the DNA-binding activity of these metal regulators, then Zur represses the expression of zinc uptake systems genes and ZntR stimulates the export systems genes expression to lower the intracellular zinc concentration and contributes to the bacterial growth (23).

**RESULTS**

Zinc broadly affects gene expression of M. bovis BCG. Zinc is an essential microelement for almost all organisms, but it is toxic at high concentrations, and thus represents an antibacterial mechanism of macrophages. It is really true for intracellular pathogens such as M. tuberculosis (Mtbc) and M. bovis. Although these pathogens have developed several mechanisms to adapt to zinc toxicity, the specific zinc exporter and
the corresponding signaling pathway for zinc detoxification are yet to be elucidated. For this purpose, we performed RNA-seq and transcriptomic assays to compare the differences in gene expression of M. bovis BCG strains under 0.5 mM zinc treatment. As shown in Fig. 1A, a total of 240 genes were found to be significantly differentially expressed, of which 131 were upregulated and 109 were downregulated. The differentially expressed genes (DEGs) were analyzed using the Cluster of Orthologous Groups (COG) database to classify them and predict their functions. 182 of 240 DEGs were assigned to COG classifications and functionally classified into 19 protein families of which the top six groups in terms of abundance ratio are showed in Fig. 1B. The cluster predicted for energy production and conversion (11.68%) emerged as the largest group, followed by transcription (10.66%), general function prediction only (10.66%), and amino acid transport and metabolism (8.63%). Notably, 7.11% DEGs are involved in inorganic ion transport and metabolism. Interestingly, inorganic ion transport and metabolism & energy production and conversion are related to metal export, so we focused on the changes in transcript levels of 12 P-type ATPase genes. Indeed, the expression of ctpC and ctpG was induced under zinc treatment (Fig. 1C and D), which is consistent with previous reported data (30). Therefore, these results indicate that zinc broadly affects genes expression in M. bovis BCG.

**Zn\(^{2+}\) remarkably induces the expression of ctpG in M. bovis BCG.** Among DEGs mentioned above, we found the upregulation fold of transcription factor cmtR was the highest (~17 times), and the expression of cmtR operon genes (Fig. S1A) included ctpG which was significantly upregulated (Fig. 2A). And CtpG is a P\(_{1\beta}\)-type ATPase (Fig. S1B), which indicates that CtpG is implicated in zinc detoxification. To test this hypothesis, we first assayed ctpG expression profile of M. bovis BCG and M. smegmatis under 0.1 mM and/or 0.5 mM zinc condition by RT-qPCR, respectively. As shown in Fig. 2B, the induction of ctpG expression was enhanced with the increase of zinc concentration in M. bovis BCG, but the expression of ctpG was not induced by 0.5 mM zinc in M. smegmatis (Fig. S2A). Next, we compared ctp genes expression of M. bovis BCG under 0.5 mM zinc treatment by RT-qPCR assays. As shown in Fig. S2B, ctpG expression was strongly induced (up to ~50 times) upon exposure to 0.5 mM zinc and its induction fold change was much higher than that of the reported Zn\(^{2+}\)-transporter gene ctpC (~8 times), and the expression of ctpH and ctpV is inhibited by zinc while other ctp genes were not obviously affected under the same experiment conditions, which is consistent with the transcriptome data described above. These results indicate that CtpG may be a main zinc efflux protein in M. bovis BCG. To investigate whether the induction expression of ctpG by zinc has specificity, we treated M. bovis BCG strains with different kinds of divalent cations. As shown in Fig. 2C, ctpG expression was induced by Zn\(^{2+}\) and the control metal Cd\(^{2+}\), and the induction effect of Zn\(^{2+}\) was much better than that of Cd\(^{2+}\). Then, we further confirmed the zinc induction effect by β-gal activity assays. As shown in Fig. 2D, hsp60p significantly promoted the expression of lacZ in M. bovis BCG strains relative to the nonpromoter lacZ plasmid, which indicates that the reporting system functioned properly. Compared with no treatment, Zn\(^{2+}\) instead of Mg\(^{2+}\) significantly stimulated the expression of lacZ when the ctpG promoter, cmtRp, was used as a promoter. In contrast, no significant difference was observed in the expression of lacZ under different experiment conditions when a negative-control fbpBp was used as a promoter. Collectively, these results suggest that zinc remarkably stimulates ctpG expression in M. bovis BCG.

**CtpG enhances the resistance of M. bovis BCG to Zn\(^{2+}\) toxicity.** Next, we performed the phylogeny analysis of CtpG homologs. As shown in Fig. 3A, CtpG are phylogenetically related to Zn\(^{2+}\)/Cd\(^{2+}\) transporters (ZosA and ZntA) (data from P-TYPE ATPase DATABASE). Many conserved sites in these proteins such as phosphatase domain, conserved HP and GXGXXG/A as well as GDGXNDXP motif are highly similar, but the metal-coordinating transmembrane residues in TM6 of CtpG are the APC residues (Ala-Pro-Cys\(_{420}\)) other than the SPC and CPC residues in the P-type ATPases mentioned above (Fig. 3B). Thus, we speculate that CtpG is a novel protein involved in zinc transport and APC residues play an important role in this process. To confirm this, the
FIG 1 RNA-seq assays for studying the effect of zinc on gene expression in M. bovis BCG. (A) Volcano plot of the difference in gene expression of M. bovis BCG under 0.5 mM zinc treatment determined by RNA-Seq assays. The x axis and y axis indicate the log₂(Fold change) values and the log₁₀ (adjusted P value) values, respectively, of all genes. The Cuffdiff program was executed to perform differential expression tests using the edgeR package. The differential expression of a gene is identified as significant if the fold change is ≥ 2 and the false discovery rate-adjusted P value < 0.05. The significantly upregulated and downregulated genes are indicated by red and blue spots, respectively. Genes that did not undergo significant changes in expression are indicated by gray spots. The cmtR operon genes included ctpG, BCG_2010c, (Continued on next page)
cysteine of the APC motif in the sixth TM segment of CtpG was changed to Ala (the APA mutant) and the motif was deleted (ΔAPC) as well. Then, we via an integrative vector constructed wild-type strain (WT/pMindD), ctpG-deleted strain (ΔctpG/pMindD), and complementary strains (comp-ctpG, comp-ctpG [Mut], comp-ctpG [ΔAPC]) in M. bovis BCG (Fig. S3 and S4), and assayed their resistance to Zn²⁺/Cd²⁺ toxicity. As shown in Fig. 3D, the growth of ΔctpG/pMindD, comp-ctpG (Mut), comp-ctpG (ΔAPC) cultured in liquid medium under 0.5 mM zinc were much slower than WT/pMindD for 4 and 6 days. However, no obvious growth difference was observed among these strains in the absence of zinc stress (Fig. 3C). Interestingly, 10 nM Cd²⁺ result in a similar phenotype (Fig. S5A and S5B), which is consistent with the previous results (35). Furthermore, we observed the growth difference among these strains when they were diluted and spotted on 7H10 solid medium containing 0.5 mM zinc. As shown in Fig. S6 (left panel), no growth difference was observed among these strains in medium under normal conditions. In contrast, ΔctpG/pMindD and comp-ctpG (Mut) grew worse than that of the WT/pMindD under 0.5 mM zinc treatment, and comp-ctpG (ΔAPC) did not grow; no growth difference was observed between comp-ctpG and WT/pMindD under the same experiment condition (Fig. S6, right panel). These results indicate that ctpG expression enhances M. bovis BCG resistance to zinc toxicity and APC residues play an important role in this process.

The induction of ctpG by zinc depends on CmtR. The expression of cmtR operon where ctpG is located was significantly induced by zinc which can inhibit the DNA-binding ability of CmtR to its promoter (32). Therefore, we assume that zinc-induced ctpG expression is dependent on CmtR. To confirm this, we performed RNA-Seq and transcriptomic assays to compare the differential gene expression profile of cmtR-deleted strains in M. bovis BCG under 0.5 mM zinc treatment, and then analyzed the correlation of the differentially expressed genes in wild-type strains and cmtR-deleted strains. As shown in Fig. 4A (and 4B & Table S1), 38 genes, including ctpG, depended on CmtR to respond to zinc. Further, we verified the dependence between ctpG expression and CmtR under 0.5 mM zinc by RT-qPCR and β-gal activity assays. As shown in Fig. S7, the expression of ctpG and the control gene fbpB was remained unaffected in the cmtR-deleted strain under 0.5 mM zinc treatment compared with that under no treatment, but ctpG expression was significantly induced; no obvious difference was observed in the expression of lacZ when cmtRp as well as control promoter fbpBp used as the promoter in cmtR-deleted strain under 0.5 mM Zn²⁺ or 0.5 mM Mg²⁺ (Fig. 4C). Taken together, these data indicate that ctpG expression induced by zinc is dependent on CmtR.

Zn²⁺ stimulates the ATPase activity of CtpG. CtpG belongs to the family of P₁₈-ATPase which follows the classical E₁/E₂ Albers-Post transport of metals across the membrane. The central characteristic of the transport mechanism of all P-ATPases is coupling of transmembrane substrate transport to ATP hydrolysis (28). Therefore, we cloned and expressed CtpG, CtpG (Mut) and CtpG (ΔAPC) in E. coli (Fig. S8A), then assayed their ATPase activity. As shown in Fig. S8B, the ATPase activity increased parallel to the amount of these proteins that were added to the enzymatic reaction, and no significant activity difference among these proteins was observed. Then, we verified the effect of zinc on the ATPase activity of CtpG. As shown in Fig. 5A, the ATPase activity of CtpG was stimulated by addition zinc into the assay media, and the dependence of the ATPase ac-

FIG 1 Legend (Continued)
FIG 2 Analysis of the effect of zinc on ctpG expression in M. bovis BCG. (A) Heat map and transcriptional profile analysis of cmtR operon genes expression profile in the RNA-Seq assays mentioned in Fig. 1. Red-blue density display showing the expression levels. (B-C) RT-qPCR assays for ctpG expression of M. bovis BCG under zinc treatment (B) or following the exposure of the mycobacterial strain to various divalent metal cations (C). Bacteria were left untreated or were incubated with 0.5 mM Zn\(^{2+}\), 0.2 mM Cd\(^{2+}\), 0.5 mM Mg\(^{2+}\), 0.5 mM Ca\(^{2+}\), 0.2 mM Mn\(^{2+}\) or 0.2 mM Ni\(^{2+}\) in 7H9 medium for 24 h. (D) Assays for the cmtR operon promoter activities under Zn\(^{2+}\) or Mg\(^{2+}\) treatment. β-gal activity was evaluated in wild-type strain of M. bovis BCG. Left column: schematic representation of recombinant strain generation using reporter plasmids. Null promoter-lacZ, hsp60p-lacZ, and fbpBp-lacZ were used as controls. Error bars represent the (Continued on next page)
tivity on the zinc concentration showed an apparent $K_{1/2}$ of 0.24 ± 0.05 μM, a $V_{\text{max}}$ of 6 ± 0.4 U/mg. Furthermore, we evaluated metal ion specificity for the ATPase activity of CtpG. As shown in Fig. 5B, CtpG exhibited the maximum ATPase activity in the presence of zinc compared to other divalent metal cations when 1 μM metal ion was used, which indicates that zinc specifically stimulated the ATPase activity of CtpG. Based on this result, we compared the ATPase activity difference of CtpG, CtpG (Mut), and CtpG

FIG 3 Assays for studying the effect of CtpG on zinc detoxification in M. bovis BCG strains. (A) Phylogenetic relationships of CtpG paralogs. Phylogeny was constructed using the MEGA X software involving the Neighbor-Joining method (bootstrap: 10000 replicates, bootstrap values indicated by circle sizes). CtpG of Mycobacterium bovis BCG and Mycobacterium tuberculosis H37Rv are colored in red. Bacterial sequences shown are from P-type ATPase database (http://traplabs.dk/patbase/): CtpG from Mycobacterium bovis BCG (Mbb) and Mycobacterium tuberculosis H37Rv (Mtu); ZosA from Bacillus subtilis (strain 168) (Bsu); ZiaA from Synechocystis sp (Syn); ZntA from Escherichia coli (strain K-12) (Eco). (B) Amino acid sequence alignment (generated using ClustalW) of CtpG and Zn$^{2+}$-ATPases mentioned in (A) for the analysis of conservative amino acid sequences. (C-D) Assays for the effect of ctpG deletion on the growth of M. bovis BCG strain in 7H9 medium (C) or medium supplemented with 0.5 mM zinc(D). WT/pMindD represents the BCG/pMindD strain; ΔctpG/pMindD represents the BCG ctpG::hyg/pMindD strain; comp-ctpG represents the BCG ctpG::hyg/pMindD-ctpG strain; comp-ctpG (Mut) represents the BCG ctpG::hyg/pMindD-ctpG strain, and comp-ctpG (ΔAPC) represents the BCG ctpG::hyg/pMindD-ctpG (ΔAPC) strain. Error bars represent the S.D. from three biological experiments. The $P$-values of the data were calculated by unpaired two-tailed Student's t test using GraphPad Prism7. Asterisks denote significant difference (*, $P < 0.05$; ***, $P < 0.001$, two-tailed Student's t test) between two groups.

FIG 2 Legend (Continued)

S.D. from three biological experiments. The $P$ values of the data were calculated by unpaired two-tailed Student’s t test using GraphPad Prism7. Asterisks denote significant difference (***, $P < 0.001$, two-tailed Student’s t test) between two groups.
As shown in Fig. 5C, the ATPase activity of CtpG was significantly increased in the presence of 1 mM zinc, but the ATPase activity of CtpG (Mut) and CtpG (ΔAPC) were not affected. Collectively, these results indicate that zinc effectively promotes the ATPase activity of CtpG in M. bovis BCG and the APC residues play an important role in the process.

CtpG enhances mycobacterial resistance to zinc toxicity. Since zinc is a major intracellular cation that can severely affect mycobacterial growth (ΔAPC). As shown in Fig. 5C, the ATPase activity of CtpG was significantly increased in the presence of 1 μM zinc, but the ATPase activity of CtpG (Mut) and CtpG (ΔAPC) were not affected. Collectively, these results indicate that zinc effectively promotes the ATPase activity of CtpG in M. bovis BCG and the APC residues play an important role in the process.

CtpG promotes Zn$^{2+}$ efflux of M. bovis BCG. When P$_{1B}$-ATPase binds the transported metal, the enzyme undergoes phosphorylation, followed by a conformational change, which expels the metal from the cytoplasm (28). Here, we observed zinc promotes the ATPase activity of CtpG which contributes to mycobacterial resistance to zinc toxicity, implying that CtpG plays an important role in zinc export. To confirm this, we constructed a deletion strain of the reported zinc transporter CtpC in M. bovis BCG (Fig. S4), and treated wild-type strain, ctpC-deleted strain, ctpG-deleted strain, and

**FIG 4** Assays for the regulatory effects of transcription factor CmtR on ctpG expression under zinc treatment. (A) Intersectioning the differentially expressed genes in the wild-type and cmtR-deleted M. bovis BCG strains under 0.5 mM zinc treatment determined by RNA-Seq assays. (B) Comparative analysis of P-type ATPase genes expression profile of the wild-type and cmtR-deleted M. bovis BCG strains using the RNA-seq data. (C) Assays for the promoter activities of cmtRp in the presence or absence of zinc. β-gal activity was evaluated in cmtR-deleted strain under 0.5 mM Zn$^{2+}$ or 0.5 mM Mg$^{2+}$ treatment. Left column: schematic representation of recombinant strain generation using reporter plasmids. Null promoter-locZ, hsp60p-locZ, and fbpBp-locZ were used as controls. Error bars represent the S.D. from three biological experiments. The P-values of the data were calculated by unpaired two-tailed Student’s t test using GraphPad Prism7. Asterisks denote significant difference between two groups using two-tailed Student’s t test.
complementary strains mentioned above with 0.5 mM zinc for 24 h, then assayed their intracellular zinc accumulation. As shown in Fig. 6A, zinc assay kit analysis revealed that the intracellular zinc content in $D\text{ctp}G/p \text{MindD}$, comp-$D\text{ctp}G$ (Mut), comp-$D\text{ctp}G$ ($D\text{APC}$), and $D\text{ctp}C/p \text{MindD}$ was much higher than that in WT/pMindD under 0.5 mM zinc treatment; interestingly, the intracellular zinc content in $D\text{ctp}G/p \text{MindD}$ is also obviously higher than that in $D\text{ctp}C/p \text{MindD}$. These results indicate that CtpG may be the main zinc exporter in $M. \text{bovis}$ BCG and its APC residues are essential for zinc efflux.

To further confirm this, we assayed the intracellular zinc content in these strains under the same experiment conditions by ICP-OES. As shown in Fig. 6B, the intracellular zinc content in $\Delta\text{ctpG}/p \text{MindD}$, comp-ctpG (Mut), and comp-ctpG ($\Delta\text{APC}$) was much higher than that in WT/pMindD under 0.5 mM zinc treatment; interestingly, the intracellular zinc content in $\Delta\text{ctpG}/p \text{MindD}$ is also obviously higher than that in $\Delta\text{ctpC}/p \text{MindD}$. These results indicate that CtpG may be the main zinc exporter in $M. \text{bovis}$ BCG and its APC residues are essential for zinc efflux. To further confirm this, we assayed the intracellular zinc content in these strains under the same experiment conditions by ICP-OES. As shown in Fig. 6B, the intracellular zinc content in $\Delta\text{ctpG}/p \text{MindD}$, comp-ctpG (Mut), and comp-ctpG ($\Delta\text{APC}$) was significantly higher than that in WT/pMindD under 0.5 mM zinc treatment. In contrast, no obvious difference was observed among these strains under normal conditions. Taken together, these results indicate that CtpG facilitates excess zinc export and the APC residues play a pivotal role in the process.

CtpG enhances $M. \text{bovis}$ BCG resistance to Zn$^{2+}$ toxicity and enhances mycobacterial survival in THP-1 cells. Excess zinc represents an antimicrobial strategy for macrophages, and CtpG enhances $M. \text{bovis}$ BCG resistance to the stress, which implies that CtpG contributes to bacterial survival in the host cells. To confirm this, we utilized THP-1 cell model to evaluate the effect of CtpG on the intracellular survival of $M. \text{bovis}$ BCG.

**FIG 5** Assays for the effect of zinc on the ATPase activity of CtpG. (A) Kinetic parameters of CtpG. The Zn$^{2+}$-ATPase activity of CtpG follows a Michaelis-Menten kinetics. Zinc was added at different concentrations (0.1 to 2.0 $\mu$M) and the released phosphate (Pi) from the hydrolysis of ATP was quantified in the enzymatic reactions. The kinetic parameters for CtpG ($K_{1/2}$ 0.24 ± 0.05 nM and $V_{\text{max}}$ 6 ± 0.4 nmol/mg min) were calculated using GraphPad Prism 7. Error bars represent the S.D. from three biological experiments. (B) The ATPase activity of CtpG in the presence of 1 $\mu$M metals. The enzymatic ATPase activity corresponds to nmol of Pi released/mg of protein/min. (C) Assays for the effect of zinc on the ATPase activity of CtpG (Mut) and CtpG ($\Delta\text{APC}$). The purified proteins CtpG, CtpG (Mut) and CtpG ($\Delta\text{APC}$) were incubated with or without 1.0 $\mu$M zinc and measured the ATPase activity. Error bars represent the S.D. from three biological experiments. The $P$-values of the data were calculated by unpaired two-tailed Student’s t test using GraphPad Prism 7. Asterisks denote significant difference (**, $P < 0.01$; ***$, P < 0.001$, two-tailed Student’s t test) between two groups.
The results indicated that these strains invaded macrophages comparably at 2 h post-infection (hpi). However, the intracellular survival efficiency of ΔctpG/pMindD, comp-ctpG (Mut), and comp-ctpG (ΔAPC) in THP-1 macrophages was significantly lower than that of WT/pMindD at 24 and 36 hpi (Fig. 7A). Similarly, the survival efficiency of ΔctpG/pMV261 was significantly lower than that of WT/pMV261 in THP-1 macrophages at 20 hpi, whereas the survival efficiency of OE-ctpG was observably higher than that of WT/pMV261 (Fig. S9). Collectively, these results indicate that CtpG enhances mycobacterial survival in THP-1 macrophages.

Considering that some pathogens such as Mtb and M. bovis are usually exposed to zinc toxicity when infecting macrophages, and CtpG contributes to mycobacterial survival under zinc toxicity and in THP-1 macrophages, we speculate that CtpG enhances mycobacterial resistance to zinc toxicity in the cells. To test this hypothesis, we set up

![Graph A](image1)

**FIG 6** Assays for the effect of CtpG on zinc efflux in M. bovis BCG strains. (A) Zinc assay kit analysis of the metal accumulation in bacterial cells under 0.5 mM zinc treatment. ΔctpC/pMindD represents the BCG ctpC::hyg/pMindD strain. (B) ICP-OES assays for the effect of CtpG on zinc accumulation of M. bovis BCG strains under 0.5 mM zinc treatment. Recombinant strains (mentioned in Fig. 3) was left untreated or incubated with 0.5 mM zinc for 24 h at 37°C, washed twice in PBS, and the bacterial pellets processed for ICP-OES analysis. Intrabacterial zinc content expressed as nanograms zinc per grams of bacterial extract. Error bars represent the S.D. from three biological experiments. The P-values of the data were calculated by unpaired two-tailed Student’s t test using GraphPad Prism7. Asterisks denote significant difference (**, *P < 0.01; ***, *P < 0.001, two-tailed Student’s t test) between two groups.
FIG 7 Assays for the effects of CtpG on mycobacterial survival and zinc accumulation in THP-1 macrophages. (A) Intracellular survival of M. bovis BCG strains in macrophages. Recombinant strains (mentioned in Fig. 3) were used to infect THP-1 macrophages at a multiplicity of infection of 10 mycobacteria per cell. After 4 h, cells were washed and incubated in fresh medium, then lysed at the indicated postinfection time points. Serial dilutions of the supernatant were plated on 7H10 agar supplemented with 10% oleic acid-albumin-dextrose-catalase, and the number of CFU was counted 15–21 days later. (B) Composite images and quantification of DiI-labeled M. bovis BCG strains (in red), FluoZin-3 (FZ3)-labeled free zinc (in green) in the THP-1 macrophages after 20 h infection. Data points are the mean relative fluorescence of bacterial intracellular zinc from at least 35 FZ3-positive phagosomes in the graph left panel. (C) RT-qPCR analysis of the expression of the genes (lcn2, mt1, mt2, and mtf1) at 20 h after M. bovis BCG strains infection in THP-1 macrophages. Error bars represent the S.D. from three biological experiments. The P-values of the data were calculated by unpaired two-tailed Student’s t test using GraphPad Prism7. Asterisks denote significant difference (*, P < 0.05; **, P < 0.01; ***, P < 0.001, two-tailed Student’s t test) between two groups.
to visualize the intracellular free zinc in THP-1 macrophages with FluoZin-3 (FZ3), a fluorescent probe specific for free zinc, by confocal assays during *M. bovis* BCG strains infection (36, 37). As shown in Fig. 7B, significantly increased colocalization between ctpG-deleted strain and FZ3 in the infected THP-1 macrophages was observed compared with that of the wild-type strain, and the ctpG-deleted strain is more likely to induce the production of zinc in macrophages (Fig. S10A). Further, we assayed the expression of several metallothionein (MT)-encoding genes (*mt*1, *mt*2, and *mtf*1) which were induced when the THP-1 cell is exposed to potentially toxic concentrations of free zinc (38). As shown in Fig. S10B, the induction fold of these genes was the most obvious at 20 hpi when THP-1 macrophages were infected with *M. bovis* BCG. Subsequently, we compared the expression difference of these genes at 20 hpi when recombinant strains infected. As shown in Fig. 7C, ΔctpG/pMindD, comp-ctpG (Mut), and comp-ctpG (ΔAPC) were significantly induced higher expression of *mt*1, *mt*2, and *mtf*1 than that of WT/pMindD. Collectively, our data suggest that CtpG plays a crucial role in the ability of *M. bovis* BCG to resist zinc poisoning and enhance mycobacterial survival at least in the context of human macrophages.

**CtpG enhances *M. bovis* BCG survival in mice.** To investigate whether CtpG affects *M. bovis* BCG survival in mice, we determined the bacterial loads in the lungs of C57BL/6 mice infected with the wild-type, ctpG-deleted, and ctpG-overexpressing strains. As shown in Fig. 8A, the bacterial loads in the lung tissues of mice infected with all three strains were similar at 2 days postinfection (dpi); however, the bacterial load in the lungs of mice infected with the ctpG-overexpressing strain increased significantly compared with that of mice infected with the wild-type strain from 4 to 20 dpi. In contrast, compared with the mice infected with the wild-type strain, those infected with the ctpG-deleted strain exhibited an obvious decrease in bacterial load since 4 dpi, which indicates that CtpG enhances *M. bovis* BCG survival in the lungs of mice. Furthermore, histopathological assay of lungs showed that mice infected with the ctpG-deleted strain had less-severe lung pathology, marked by reduced total cellular and neutrophilic infiltration than those infected with the wild-type strain, probably because the high load of bacteria triggered regulatory immune responses in the host (Fig. 8B and Fig. S11). Consistently, levels of *tnf* mRNA and *il1b* mRNA in mice infected with the ctpG-deleted strain were much lower (Fig. 8C). Taken together, these data suggest that CtpG enhances *M. bovis* BCG survival in mice.

**DISCUSSION**

Although zinc is an essential micronutrient for the survival and proliferation of bacteria, it is toxic at high concentrations and presents an antibacterial mechanism for macrophages. Mtb is considered one of the most persistent intracellular pathogens and has developed unique mechanisms to adapt to host environments, including zinc deficiency and excess, during infection (10, 39). However, the molecular mechanism by which mycobacteria promote detoxification of zinc and the associated signaling pathways remains largely unclear. In this study, using *M. bovis* BCG as a model, we first characterized the P1B-type ATPase CtpG acts as a novel zinc efflux protein in which the APC domain plays an important role in zinc transport, and enhances mycobacterial resistance to zinc toxicity. We further observed that CtpG responding to zinc depends on transcription factor CmtR. Lastly, we provided evidence to demonstrate that CtpG affects mycobacterial interaction with host cells and contributes to bacterial survival during infection. Our findings revealed the existence of a novel regulatory pathway on mycobacteria responding to and adapting to host-mediated zinc toxicity.

An interesting finding from the present study is that CtpG is a novel and main zinc efflux protein in *M. bovis* BCG. It has a highly developmental homology to the zinc transport proteins that have been reported in other species and belongs to a metal transport P1B-ATPase with characteristic phosphorylation domains and membrane topology (28, 40). It is now well established that metal specificity is conferred by conserved residues in the TMs flanking the large cytoplasmic ATP binding and hydrolysis...
FIG 8 Assays for the effects of CtpG on the host innate immunity responses. (A) Bacterial titers in homogenates of lung from SPF C57BL/6 mice (n = 6 mice per group) infected intratracheally with 1.0 × 10^6 of M. bovis BCG strains for 0–20 days. WT/pMV261 represents the BCG/pMV261 strain; ΔctpG/pMV261 represents the BCG ΔctpG::hyg/pMV261 strain; OE-ctpG represents the BCG/pMV261-ctpG strain. (B) Histopathology of lungs of mice infected as in (A), assessed in sections stained with hematoxylin and eosin. (C) Quantitative PCR analysis of tnf mRNA and il1b mRNA in splenic cells from mice infected as in (A). Data are representative of one experiment with two independent biological replicates (mean and S.D. of n = 6 mice per group). The P-values were calculated by unpaired two-tailed Student’s t test using GraphPad Prism7. Asterisks denote significant difference (**, P < 0.01; ***, P < 0.001, two-tailed Student’s t test) between two groups.
The analysis of the eight transmembrane sequences of homologous CtpG proteins showed that CtpG has a novel residue APC signature in TM6, and has a unique HEFTE in TM8 (Fig. 3B and Fig. S1B). Furthermore, many experimental evidences indicate that conserved Cys in TM6 participate in metal binding and transport. It has been reported that various sequences in TM6 are related to ion transport and essential for enzyme function, such as CPC, CPH, SPC, TPC or CPS. In addition, the requirement of conserved transmembrane His(H), Glu(E), Asp(D), Ser(S) and Met(M) residues for the metal translocation by various transporters has been established (40). The presence of these unique residues is not trivial because their replacement leads to inactive proteins. Interestingly, these transformations make CtpG highly specific for zinc. Direct biochemical analysis of CtpG activity showed that maximum activation of the transporter occurs in the presence of zinc. What’s more, we found that mutating APC residues to APA residues or delete it can seriously affect the enzymatic activity and zinc efflux of CtpG as well as mycobacterial resistance to zinc toxicity. In addition, we found zinc significantly induces ctpG expression, and the induction fold change is higher than that of the previously characterized export transporter CtpC. And the zinc efflux effects of CtpG is significantly better than that of CtpC. Therefore, CtpG may be a main efflux protein for M. bovis BCG adaptation to the zinc poisoning environment.

Zinc homeostasis is crucial for bacterial cells, and too much or too little can result in growth defect. Therefore, intracellular zinc concentration is tightly regulated in the cells. Usually, the process involves pairs of zinc-responsive transcription factors, similar to Zur and ZntR, which separately regulate the zinc uptake and efflux (25, 43). For example, in B. abortus, the expression of the genes encoding the zinc uptake system ZnuABC is negatively regulated by Zur (44). And ZntR controls the expression of the gene encoding the zinc exporter ZntA by binding directly to its promoter to adaptation to zinc toxicity environment (45–47). However, only the zinc uptake repressor Zur have been characterized in mycobacterial species until now. And little is known about the mechanisms and regulatory pathway on zinc efflux in these bacteria. In the present study, we successfully characterized CtpG acts as a novel zinc export transporter in M. bovis BCG and significantly enhances mycobacterial resistance to zinc toxicity. Zinc induces the expression of ctpG and promotes the ATPase activity of CtpG. Furthermore, we found the expression of ctpG induced by zinc depends on its upstream repressor CmtR which has been reported to regulate zinc homeostasis by interaction with Zur in M. bovis BCG. Therefore, our findings revealed a novel regulatory model for mycobacterial resistance to zinc toxicity.

P-type ATPases make up a ubiquitous family of proteins which couple the hydrolysis of ATP to transport cations across plasma membranes and is pivotal for cell viability. For example, Mtb P1B-type ATPase, CtpA, is associated with Cu⁺ transport, and plays an important role in Cu⁺ detoxification (48). Similarly, a putative copper exporter CtpV contributes to maintain mycobacterial resistance to Cu⁺ toxicity and enhance intracellular survive during infection (49). In addition, CtpC has been reported to be a zinc efflux transporter and required for optimal growth of Mtb under excess zinc stress and in human macrophages. However, CtpC does not transport zinc under normal experiment condition and has no significant effect for bacterial survive in mouse. CtpG, like CtpC, belongs to be a P1B-type ATPase and transports Cd²⁺ to protect M. smegmatis against metal toxicity (35). In the present study, we found CtpG promotes zinc efflux and enhances mycobacterial resistance to zinc toxicity in M. bovis BCG. Zinc specifically induces ctpG expression and stimulates its ATPase activity. Moreover, during infection, CtpG lowers the intracellular zinc accumulation and contributes to mycobacterial survival in human macrophages and in mouse. Therefore, CtpG may represent a novel defense weapon for mycobacteria to survive in host.

In summary, we characterized CtpG acts as a novel zinc efflux transporter and significantly enhances M. bovis BCG survival under excess zinc stress and during infection. Our data combined with that of previous studies suggest that CmtR is able to sense zinc and derepresses ctpG expression, following which CtpG binds zinc and hydrolyzes
ATP to release energy for intracellular zinc efflux. Our findings first identified a regulation signaling pathway by transcription regulator CmtR on CtpG-mediated zinc detoxification in mycobacteria and provide insights into stress-induced mycobacterial adaptation to the host environment.

MATERIALS AND METHODS

Plasmids, enzymes, and reagents. The pET28a plasmid obtained from Novagen was used for over-expressing the mycobacterial protein in Escherichia coli BL21(DE3) pLyS S strain (Novagen, Darmstadt, Germany). The primers (listed in Table S2) for PCR were synthesized by Tsingke (China); DNA polymerase, restriction enzymes, T4 ligase, dNTPs, and antibodies were obtained from Takara Biotech (Shiga, Japan). The 7H9/7H10 medium and oleic acid-albumin-dextrose-catalase enrichment used for mycobacterial growth were purchased from BD Biosciences. Wild-type female SPF C57BL/6 mice were purchased from Chang-sheng Bio (Liaoning, China).

Expression and purification of recombinant proteins. Genes were amplified using PCR with specific primer pairs (Table S2). Mutant genes were obtained through site-directed mutagenesis by overlapping extension PCR. The amplified DNA fragments were digested using the corresponding restriction endonucleases and cloned into the pET-28a expression vector to produce recombinant plasmids (Table S3). The expression strains of E. coli BL21(DE3) pLyS S containing the recombinant plasmids were cultured, and the recombinant proteins were purified, as described in a previous study (50). All purification steps were carried out at 0–4°C. Cells were suspended in buffer A (25 mM Tris, pH 7.0, 100 mM sucrose, 1 mM phenylmethylsulfonyl fluoride [PMSF; Sigma]) and disrupted with a French press at 20,000 lb/in². Lysed cells were centrifuged at 8,000 × g for 30 min. The supernatant was then centrifuged at 229,000 × g for 1 h, and the pelleted membranes were resuspended in buffer A (10–15 mg/mL). For protein solubilization and purification, membranes were diluted to a final concentration of 3 mg/mL in buffer B (25 mM Tris, pH 8.0, 100 mM sucrose, 500 mM NaCl, 1 mM PMSF) and solubilized with 0.75% dodecyl-β- D-maltoside (DDM; Calbiochem). The preparation was incubated for 1 h at 4°C with mild agitation and centrifuged at 229,000 × g for 1 h. The supernatant was incubated overnight at 4°C with Ni²⁺-nitrilotriacetic acid resin (Qiagen) pre-equilibrated with buffer B, 0.05% DDM, and 5 mM imidazole. The resin was washed with buffer B, 0.05% DDM containing 10 mM and 20 mM imidazole, and the protein was eluted with buffer C, 0.05% DDM, 250 mM imidazole. Fractions were pooled and concentrated, and buffer was replaced by buffer C (25 mM Tris, pH 8.0, 100 mM sucrose, 50 mM NaCl, and 0.01% DDM) using 50 kDa cutoff centriacs (Millipore). The proteins were aliquoted and stored in 20% (vol/vol) glycerol at −80°C until use. All protein determinations were performed in accordance with the Bradford method (51). Purified CtpG protein was analyzed by 10% SDS-PAGE followed by Coomassie brilliant blue (CBB) staining.

Construction of recombinant mycobacterial strains. ctpG, ctpG (Mut), ctpG (ΔAPC) and ctpC genes were amplified by PCR using specific primer pairs (Table S2), and the amplicons were digested by the corresponding restriction endonucleases. The digested ctpG, ctpG (Mut), ctpG (ΔAPC) and ctpC genes were separately cloned into pMV261 overexpression vectors (52) or a pMindD vector (53) and transformed into the wild-type and knockout strains of M. bovis BCG to generate overexpression and complementary strains, respectively, in which the expression of the target genes was regulated using anhydrotetracycline hydrochloride. ctpG or ctpC knockdown was performed in M. bovis BCG strains, as described previously (54).

Transcriptomic analysis. M. bovis BCG strains (BCG/WT and BCG/cmrR::hyg) were cultured in 7H9 medium with shaking at 160 rpm at 37°C until the midlogarithmic phase. Then, the cells were treated with 0.5 mM zinc for 24 h and harvested from each sample (each strain in three biological replicates). Subsequent transcriptomic analysis was performed, as described previously (54). In brief, total RNA was isolated using RNAprep Pure Cell/Bacteria kit (Tiangen, China). Strand-specific libraries were prepared using the NEBNext Ultra RNA Library Prep kit for Illumina (Illumina, USA) according to the manufacturer’s instructions. Library construction and sequencing were performed at Novogene Corporation.

The volcano plot diagrams were constructed using the log10(P value) and log10(fold change) values of the genes of M. bovis BCG strains under zinc treatment conditions using the ggplot2 package. Briefly, the cuffdiff program (55) was performed to conduct differential expression tests for the wild-type or cmrR-deleted strains under zinc treatment using the edgeR package (56). A transcript will be reported as differential expression significant if the test gives that the false discovery rate-adjusted P-value after Benjamin-Hochberg correction for multiple-testing represents statistical significance (P value < 0.05) (57) and the fold change is ≥ 2. The changes in gene expression are indicated on the x axis, and the P values are indicated on the y axis. The red and blue spots represent the upregulated and downregulated genes, respectively, whereas the gray spots represent genes with insignificant changes in expression.

The heat map was constructed using the Hemi (Heatmap Illustrator, version 1.0) software. Briefly, the fragments per kilobase of exon per million fragments mapped values of target genes in each sample were normalized and imported to the Hemi software to construct the heat map diagram. Red represents high expression, whereas blue represents low expression of the target genes in different samples. The color scale beside the heat map indicates the color threshold.

Assay for β-gal activity. Assays for measuring β-gal activity was separately performed using the wild-type and cmrR-deleted M. bovis BCG strains by constructing operon-lacZ fusions based on the expression vector pMV261. The target and control promoters were amplified by PCR using the respective pairs, which are listed in (Table S2), after which the amplicons were digested using the corresponding restriction endonucleases and cloned into the pMV261 backbone. The reporter gene lacZ was cloned downstream of the promoters. The plasmids were separately transformed into the cmrR-deleted and wild-
type M. bovis BCG strains to obtain the corresponding recombinant reporter strains. The recombinant strains were cultured until the midlogarithmic phase, then treated with 0.5 mM Zn\(^{2+}\) and 0.5 mM Mg\(^{2+}\) for 24 h, respectively. The bacterial cells were harvested and washed using phosphate-buffered saline (PBS). The levels of galactosidase were measured as described previously (32).

**Evaluation of mycobacterial growth.** The growth patterns of M. bovis BCG strains were evaluated using modified versions of procedures described earlier (32). The recombinant strains were cultured in 7H9 medium supplemented with 30 μg/mL Kan and 50 ng/mL anhydrotetracycline hydrochloride, and the cultures were incubated under shaking conditions at 160 rpm at 37°C. When the culture reached the midlogarithmic phase, each culture was diluted (1:100) in 100 mL of fresh 7H9 broth supplemented with 0.5 mM zinc, and the sensitivity of the recombinant strains was determined every 2 days.

**Determination of metal ion content under zinc treatment.** Total zinc contents in the dried bacterial pellets was determined by ICP-OES (Varian, USA) according to a previously published procedure with several modifications (34). Briefly, the ctpG recombinant strains were cultured in 300 mL of 7H9 medium under shaking conditions at 160 rpm at 37°C until the midlogarithmic phase, after which the strains were treated with 0.5 mM zinc for 24 h. The harvested samples were washed twice in PBS containing 1 mM EDTA and 0.05% Tween 80, and were then washed only with PBS. The pellets were stored overnight at -80°C and dried using the Freeze Dryer machine (Thermo Fisher). The dried samples were weighed and acid digested with HNO\(_3\) (trace metal grade) for 4 h at 80°C and overnight at 65°C. The supernatants were filtered by passing through 0.22-μm filters (Corning Inc.). The metal content in the digested samples was measured by ICP-OES. The concentration of zinc in the ctpG recombinant strains was further determined using a Zinc assay kit (MAK032; Sigma-Aldrich). Briefly, M. bovis BCG strains were grown to midlogarithmic phase, were pelleted, washed, and suspended in PBS, and were ultrasonic crushing treatment 10 min (200 W). The extract was deproteinized by adding an equal volume of trichloroacetic acid (7%). The mixture was then centrifuged at 12000 rpm for 5 min to obtain the cell extract as the supernatant. Then, 50 μL of the cell extract was mixed with 200 μL of Zn reagent mix and the reaction mixture was incubated at room temperature for 10 min. Absorbance of the reaction mixture was read at 560 nm.

**ATPase assays.** The enzymatic reactions (50 μL) were performed in the reaction buffer (40 mM MOPS-TRIS, 3 mM MgCl\(_2\), 5 mM Na\(_2\)ATP, 0.25 mM Na\(_2\)MoO\(_4\), and 0.5 mM Cys, pH 7.0) using 4 μg of CtpG protein, and supplemented with 1 μM (final concentration) of the tested heavy metal cation (Cd\(^{2+}\), Zn\(^{2+}\), Mg\(^{2+}\), Ca\(^{2+}\), Mn\(^{2+}\), and Ni\(^{2+}\)). The enzymatic reactions were initiated by the addition of 3 mM Na\(_2\)ATP, followed by incubation at 37°C for 30 min. The ATPase activity was stopped by adding 100 μL of stop solution (3% ascorbic acid, 0.5% ammonium molybdate, and 3% SDS in 1.0 M HCl) and 150 μL of 3.5% bismuth citrate and 3.5% sodium citrate in 2.0 M HCl. Subsequently, the released inorganic phosphate (Pi) was quantified (58). The OD\(_{690}\) of the samples was measured using an iMark Microplate Absorbance Reader (Bio-Rad, Philadelphia, PA, United States) and interpolated in a calibration curve from 0 to 100 mM NaH\(_2\)PO\(_4\). The total enzymatic activity was reported as nmol of Pi released/mg of protein. min; the reactions were assessed in triplicate from three independent experiments.

**Kinetic parameters of CtpG.** The Zn\(^{2+}\)-ATPase activity of CtpG was assayed as described above (50). To estimate the dependence of CtpG on zinc concentration, enzymatic kinetics were performed under the optimal reaction conditions and varying the zinc concentration from 0.1 to 2 μM and the enzymatic reactions were performed for 30 min. The enzymatic activity was reported as nM of Pi released/mg of protein. min and assessed in triplicate for three independent experiments. The values of K\(_{1/2}\) and V\(_{max}\) were calculated using the nonlinear least-squares regression for Michaelis-Menten enzyme kinetics using Prism 7 version, GraphPad Software, La Jolla California USA.

**Intracellular survival assays.** Monocytic THP-1 cells were seeded in 24-well plates (5 × 10\(^3\) cells per well in RPMI 1640 supplemented with 10% heat-inactivated fetal bovine serum (FBS) (Gibco) in a volume of 1 mL) and differentiation was stimulated by the addition of 100 nM/mL phorbol 12-myristate 13-acetate (PMA) (Sigma-Aldrich). THP-1 cells were incubated for 48 h at 37°C and 5% CO\(_2\) prior to infection. M. bovis BCG strains cultured in 7H9 to midlogarithmic phase, were centrifuged and washed with infection medium (RPMI 1640, PAA) containing 1% heat-inactivated FBS. THP-1 cells were infected with M. bovis BCG strains using a multiplicity of infection (MOI) of 10 bacteria per phagocyte at 37°C and 5% CO\(_2\) in infection medium. Bacteria were slightly centrifuged (2 min, 200 × g) onto the cells to initiate a simultaneous contact with phagocytes. Post infection, phagocytes were washed with infection medium and subsequently incubated with Penicillin G (100 units/mL, Sigma-Aldrich) and Gentamicin (0.1 mg/mL, Sigma-Aldrich) for 1 h at 37°C and 5% CO\(_2\). After washing, the phagocytes were lysed using 0.025% SDS at the indicated time points. The CFU (CFU of released intracellular M. bovis BCG strains was determined by plating the bacteria in appropriate dilutions on 7H10 agar plates.

**Confocal microscopy.** For visualization of free zinc, the THP-1 cells infected with Dil-labeled M. bovis BCG strains were incubated for 1 h with the cell permeant reagent F23 (Invitrogen) at a final concentration of 1 mM in PBS. The confocal images were acquired using the Olympus FV1000 confocal microscope. An UPlanSapo100x/1.40 NA oil objective was used to obtain an over sampled 1024 × 1024 image with 49 nm pixel and a z-stack with a step size of 120 nm, recorded with Zen software (Carl Zeiss, Inc.), and analyzed using Zen 2010 with Olympus FV1000 confocal microscope.

**Mouse infection.** WT female SPF C57BL/6 mice were purchased from Chang-sheng Bio (Liaoning, China). The mice weighed 16–18 g and were 6–8 weeks old and housed in a specific pathogen-free facility using standard humane animal husbandry protocols that were approved by the Research Ethics Committee of the College of Veterinary Medicine, Huazhong Agricultural University, Hubei, Wuhan, China (HZAUMO-2019-013). M. bovis BCG strains were cultured until the midlogarithmic phase and were washed three times with PBS containing 0.05% Tween 80. Thirty-six mice were randomly divided into six groups.
(n = 6) and were intratracheally infected with 1 × 10^6 CFU of WT/pMV261, ΔctpG/pMV261, and OE-ctpG strains, respectively. Data were analyzed as described previously [19].

**Statistical analysis.** Statistical analyses of data were performed using unpaired two-tailed Student’s t test with GraphPad Prism 7. Data are expressed in terms of mean ± S.D. Asterisks represent significant difference, *P < 0.05; **P < 0.01; ***P < 0.001; and ns, not significant (P ≥ 0.05).

**Data availability.** All data described are presented either within the article or in the supporting information. The transcriptomic data of M. bovis BCG strains under zinc treatment conditions were deposited to the Gene Expression Omnibus (GEO) for the accession No: GSE179966.

**SUPPLEMENTAL MATERIAL**

Supplemental material is available online only.

**SUPPLEMENTAL FILE 1**, PDF file, 1 MB. **SUPPLEMENTAL FILE 2**, XLSX file, 0.02 MB. **SUPPLEMENTAL FILE 3**, XLSX file, 0.03 MB. **SUPPLEMENTAL FILE 4**, XLSX file, 0.03 MB. **SUPPLEMENTAL FILE 5**, XLSX file, 0.1 MB. **SUPPLEMENTAL FILE 6**, XLSX file, 0.1 MB.

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CtpG Enhances Mycobacterial Resistance to Zinc Toxicity

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