

# Alkyl Hydroperoxide Reductase Repair by *Helicobacter pylori* Methionine Sulfoxide Reductase

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Protein exposure to oxidants such as HOCl leads to formation of methionine sulfoxide (MetSO) residues, which can be repaired by methionine sulfoxide reductase (Msr). A *Helicobacter pylori* *msr* strain was more sensitive to HOCl-mediated killing than the parent. Because of its abundance in *H. pylori* and its high methionine content, alkyl hydroperoxide reductase C (AhpC) was hypothesized to be prone to methionine oxidation. AhpC was expressed as a recombinant protein in *Escherichia coli*. AhpC activity was abolished by HOCl, while all six methionine residues of the enzyme were fully to partially oxidized. Upon incubation with a Msr repair mixture, AhpC activity was restored to nonoxidized levels and the MetSO residues were repaired to methionine, albeit to different degrees. The two most highly oxidized and then Msr-repaired methionine residues in AhpC, Met<sub>101</sub> and Met<sub>133</sub>, were replaced with isoleucine residues by site-directed mutagenesis, either individually or together. *E. coli* cells expressing variant versions were more sensitive to *t*-butyl hydroperoxide than cells expressing native protein, and purified AhpC variant proteins had 5% to 39% of the native enzyme activity. Variant proteins were still able to oligomerize like the native version, and circular dichroism (CD) spectra of variant proteins revealed no significant change in AhpC conformation, indicating that the loss of activity in these variants was not related to major structural alterations. Our results suggest that both Met<sub>101</sub> and Met<sub>133</sub> residues are important for AhpC catalytic activity and that their integrity relies on the presence of a functional Msr.

*Helicobacter pylori* is a spiral, Gram-negative bacterium that colonizes the gastric epithelium (1) in about 50% of the world's population (2). *H. pylori* has been shown to be the etiological agent of peptic ulcer disease or chronic atrophic gastritis (3), which can subsequently develop into gastric cancer (4). Colonization of the gastric mucosa by *H. pylori* induces a strong inflammatory response in the host, who produces harmful reactive nitrogen species (RNS) or oxygen species (ROS) (5); the latter include superoxide anion (O<sub>2</sub><sup>-</sup>), hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), hydroxyl radical (·HO), and hypochlorous acid (HOCl). Increased production of ROS by gastric cells (6) and phagocytes (7) induced by *H. pylori* has been shown *in vitro*, and increased levels of ROS in the gastric mucosa have been found in *H. pylori*-infected patients (8). ROS have the ability to react with and damage bacterial proteins, as well as lipids and DNA. Protein covalent modifications include amino acid modifications or carbonyl group formation, usually leading to partial or total loss of catalytic activity. Among the amino acid modifications, oxidation of the sulfur-containing methionine (Met) into R- and S-diastereoisomers of Met sulfoxide (Met-SO) has been well documented (9–11).

To counter oxidative stress, *H. pylori* cells have evolved different defense strategies (reviewed in references 12 and 13), including (i) avoidance, (ii) use of antioxidant enzymes such as superoxide dismutase (SOD), catalase (kat), and alkyl hydroperoxide C (AhpC), and (iii) use of repair enzymes such as MutS or methionine sulfoxide reductase (Msr). The latter catalyzes the conversion of Met-SO back to Met (14). While most organisms have two distinct Msr proteins, MsrA and MsrB, shown to reduce the two isomers Met(S)O and Met(R)O of methionine sulfoxide, respectively (15), *H. pylori* possesses only one Msr protein that consists of MsrA-like and MsrB-like domains (16). *H. pylori* *msr* mutants show increased sensitivity to oxidants such as paraquat and H<sub>2</sub>O<sub>2</sub> as well as to nitrosative (GSNO) stress and exhibit attenuated colonization of mice compared to wild-type cells (17). Upon exposure of Msr-deficient mutants to oxidative

stress, several methionine-containing proteins were isolated, including GroEL, thioredoxin 1 (Trx1), and catalase (14). Cross-linking interaction studies using purified recombinant proteins revealed that Msr can bind to each of those proteins (14). In addition, oxidation of purified catalase by HOCl generated Met-SO residues within the H<sub>2</sub>O<sub>2</sub>-scavenging enzyme and these Met-SO residues were shown to be subsequently repaired upon incubation with an enzyme repair mix that required Msr (10). Recently, the UreG urease accessory protein was also identified in our laboratory as a primary target for Met oxidation (18). Evidence was provided for an Msr-UreG transient interaction as well as for Msr-mediated repair of Met-SO residues within UreG (18). While these findings shed new light on the role of Msr in repairing oxidative stress-damaged proteins, as well as on its interaction with methionine-rich proteins in *H. pylori*, we sought to find new Msr-repaired targets.

AhpC is a key enzyme to combat oxidative stress: it detoxifies harmful lipid hydroperoxides. In the current study, we hypothesized that this enzyme would be a target for methionine oxidation because (i) other proteins known to combat oxidative stress, such as catalase and Trx1, have been shown to be primary targets for Met-SO formation (14), (ii) AhpC is one of the most abundant proteins in *H. pylori*, and (iii) its Met content (3% of the total

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amino-acid content) is higher than average. To get a better assessment of the oxidative stress-induced protein damage and to determine which Met residues would be preferentially oxidized and repaired, we exposed recombinant AhpC to HOCl and measured the level of oxidation of each individual methionine residue before and after repair by Msr. Finally, the role of the two most oxidized methionine residues of AhpC was investigated by replacing these two residues with isoleucine residues and studying the effect of both substitutions on the *in vitro* and *in vivo* AhpC activity.

## MATERIALS AND METHODS

**Bacterial strains and plasmids.** *Escherichia coli* TOP10 (Invitrogen) was used for all cloning experiments, and *E. coli* BL21(DE3)RIL (Novagen) was used to overexpress recombinant *H. pylori* AhpC (*HpAhpC*) proteins. *H. pylori* wild-type strain 26695 (16) was used as a source of DNA for PCR. *H. pylori* SS1 (19) and *H. pylori* SS1  $\Delta$ *msr* (17) were used for all other experiments. Plasmid pET21b (Invitrogen) was used to clone and express native and variant AhpC proteins. All plasmids and PCR products used were sequenced at the Georgia Genomics Facility of the University of Georgia, Athens, GA, and compared to DNA sequences from strain 26695 (16) to ensure that no error had been introduced following PCR amplification.

**Growth conditions.** *E. coli* cells were grown aerobically in Luria-Bertani (LB) medium or plates at 37°C. Ampicillin (100  $\mu$ g/ml), chloramphenicol (30  $\mu$ g/ml), and isopropyl- $\beta$ -D-thiogalactopyranoside (IPTG) (0.1 to 0.2 mM) were added as needed. *H. pylori* was routinely grown on Brucella agar plates supplemented with 10% defibrinated sheep blood (BA) at 37°C under microaerophilic conditions with 1% to 5% O<sub>2</sub>, 5% CO<sub>2</sub>, and 90% to 94% N<sub>2</sub>, as indicated.

**Sensitivity of nongrowing cells to HOCl.** The *H. pylori* wild type (SS1) and the SS1 $\Delta$ *msr* mutant were grown for 36 h on BA plates under 1% O<sub>2</sub> conditions, harvested, and resuspended in phosphate-buffered saline (PBS) to an optical density at 600 nm (OD<sub>600</sub>) of 1. Cells (0.9 ml) were mixed with 0.1 ml freshly prepared HOCl (available as NaClO [Sigma]; final concentrations ranging from 31.25 to 500  $\mu$ M) and incubated for 30 min at room temperature, with gentle shaking (40 rpm). HOCl-exposed cells were then serially diluted, and 10  $\mu$ l of each dilution ( $n = 4$ ) was plated on BA plates. CFU were counted after 3 to 5 days. Data are shown as the averages and standard deviations of the results of quadruplicate assays from one experiment. Another experiment (also done in quadruplicate) confirmed that the mutants were significantly more sensitive to HOCl than the parental cells.

**Cloning, expression, and purification of AhpC.** Recombinant native and variant AhpC proteins were expressed as hexahistidine-tagged proteins in *E. coli* BL21(DE3)RIL. Briefly, primers AhpC1 (5'-CCGGATCCATATGTTAGTTACAAAACCTTGCC-3') and AhpC2 (5'-CACGGTC TCGAGAAGCTTAATGGAATTTCTTTG-3') were used to amplify a 600-bp-long DNA sequence containing the whole *hp1563* (*ahpC*) open reading frame (ORF) (without its stop codon), as well as to incorporate a 5' NdeI restriction site and a 3' XhoI restriction site, respectively. The PCR product was digested with NdeI and XhoI, gel purified, and cloned into similarly digested pET21b plasmid, generating pET-AhpC. Plasmid pET-AhpC.M<sub>101</sub>I was generated by overlapping PCR using genomic DNA and primers AhpC1 and AhpC2, along with primers AhpC3 (5'-CC TATCGTGGCTGATATTACCA-3') and AhpC4 (5'-CCACGATAGGGA AAGTACTTG-3') (new codons are shown in bold). Plasmid pET-AhpC.M<sub>133</sub>I was generated by overlapping PCR using genomic DNA and primers AhpC1 and AhpC2, along with primers AhpC5 (5'-AACATCAA AGTAAAGGCATGCGG-3') and AhpC6 (5'-CTTTGATGTTTTGTCAA TCAAAAAC-3'). Plasmid pET-AhpC.M<sub>101</sub>I.M<sub>133</sub>I (double substitution) was generated using pET-AhpC.M<sub>101</sub>I as the template and primers AhpC1, AhpC2, AhpC5, and AhpC6. BL21(DE3)RIL cells transformed with the appropriate pET-AhpC plasmid were grown at 37°C in 500 ml of LB supplemented with ampicillin and chloramphenicol until an OD<sub>600</sub> of 0.5 was reached. Gene expression was induced by the addition of 0.2 mM

IPTG followed by incubation at 22°C for 3 to 5 h. Cells were harvested by centrifugation (10,000  $\times$  g, 15 min, 4°C), and His-tagged proteins were purified following the manufacturer's instructions (Qiagen, Valencia, CA) and finally dialyzed three times against 1 liter of 50 mM NaH<sub>2</sub>PO<sub>4</sub> buffer (pH 7.5). Purified AhpC wild-type or mutant proteins were purified to a level close to homogeneity (as estimated by SDS-PAGE).

***In vitro* HOCl oxidation of purified AhpC.** The *in vitro* oxidation of AhpC was carried out as previously described by Mahawar et al. (10). Briefly, 10  $\mu$ M purified AhpC was incubated with a 60-fold-molar-excess ratio of HOCl in oxidation buffer (50 mM sodium phosphate [pH 7.5], 50 mM KCl, 10 mM MgCl<sub>2</sub>) for 15 min at room temperature in the dark. Residual oxidants were quenched by excess free methionine (5 mM) for 10 min on ice and finally dialyzed twice against 1 liter of 50 mM sodium phosphate buffer (pH 7.5) to remove HOCl and free methionine.

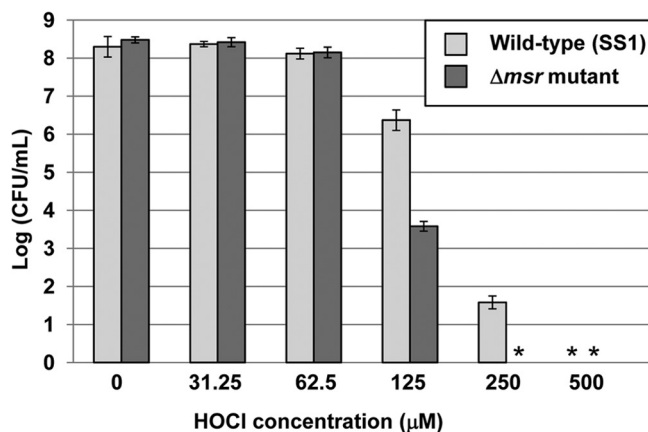
**Repair of AhpC Met-SO residues by Msr.** Equimolar amounts of HOCl-oxidized AhpC and Msr were incubated for 2 h at 37°C in the presence of 100  $\mu$ M dithiothreitol (DTT) and a thioredoxin system containing 5  $\mu$ M purified *H. pylori* thioredoxin 1 (Trx1), 0.1  $\mu$ M purified *H. pylori* thioredoxin reductase (TrxR), and 400  $\mu$ M NADPH in 50 mM sodium phosphate buffer (pH 7.4). These components plus Msr are referred to here as the repair mixture. Purification of Msr, Trx1, and TrxR was previously described (14).

**Peroxide reductase activity of recombinant AhpC.** AhpC activity was measured in a coupled assay by monitoring thioredoxin reductase-dependent oxidation of NADPH at 340 nm, as previously described (20, 21). Briefly, each assay was carried out with 1  $\mu$ M AhpC (previously unoxidized or HOCl oxidized or HOCl oxidized and Msr repaired) in a final volume of 100  $\mu$ l containing 50 mM potassium phosphate buffer (pH 7.0), 100 mM ammonium sulfate, and 0.5 mM EDTA in the presence of 0.5  $\mu$ M Trx1, 0.5  $\mu$ M TrxR, and 150  $\mu$ M NADPH; 1 mM H<sub>2</sub>O<sub>2</sub> was added to start the reaction. To account for non-AhpC-dependent NADPH oxidation, our control included all the above except AhpC. To rule out any Msr-dependent NADPH oxidation, a Msr-only control included Trx1, TrxR, and Msr. The decrease of NADPH absorbance (at room temperature) was followed for 2 min using an ELX800 BioTek microplate reader (Winooski, VT). One unit of specific activity is defined as 1  $\mu$ mol of NADPH oxidized per min per mg of AhpC protein. Data shown are means and standard deviations of the results from 4 to 15 replicate assays.

**Detection and quantification of oxidized methionine residues by LC-MS/MS.** For detection and quantification of oxidized Met residues within AhpC, the protein was digested with sequencing-grade GluC, following the recommendations of the manufacturer (Promega, Madison, WI). GluC digests of (i) unoxidized AhpC, (ii) unoxidized AhpC with Msr, (iii) oxidized AhpC, or (iv) oxidized and Msr-repaired AhpC were separated by reversed-phase liquid chromatography (LC) using trapping cartridges coupled to a 50-mm-long, 200- $\mu$ m-inside-diameter (ID) Halo C<sub>18</sub> column (Advanced Materials Technology, Wilmington, DE), and the eluent was analyzed by electrospray ionization-tandem mass spectrometry (ESI-MS/MS) using a linear trap quadrupole-Fourier transform (LTQ-FT) mass spectrometer (Thermo Scientific, West Palm Beach, FL) as described before (10, 18).

**Disk inhibition assays.** Disk inhibition assays were done following the method of Lundström and Bölin (22). Briefly, *E. coli* BL21(DE3)RIL strains with either pET21b vector (control) or one of the pET-AhpC plasmids were grown to an OD<sub>600</sub> of 2, and 0.2 ml of cells was mixed with 3.5 ml of top agarose (0.8%) before being spread on top of LB plates supplemented with ampicillin and chloramphenicol and with 0.1 mM IPTG when indicated. Sterile paper disks (7.5-mm diameter) were placed in the center of each plate, and 10  $\mu$ l of 5% *t*-butyl hydroperoxide was added to the disk. Cells were allowed to grow for 24 h, and the diameter of the inhibition zone was recorded. Data are shown as means and standard deviations of the results from measurements done on 3 to 5 plates.

**CD analysis.** Circular dichroism (CD) spectra of AhpC were obtained on an Aviv 400 spectropolarimeter (Lakewood, NJ) at the Robert Wood Johnson Medical School CD Facility, Piscataway, NJ. Purified recombi-



**FIG 1** Sensitivity of nongrowing *H. pylori* wild-type and  $\Delta msr$  mutant cells to HOCl. *H. pylori* SS1 (wild-type) and SS1  $\Delta msr$  mutant cells were grown on BA plates, harvested, and resuspended in PBS containing the indicated HOCl concentration for 30 min at room temperature. Cells were serially diluted and plated on BA plates, and CFU were counted after 3 to 5 days. Data shown represent the averages and standard deviations of the results of quadruplicate assays from one experiment, and the mutant is significantly more sensitive than the parent at 125 and 250  $\mu\text{M}$  HOCl ( $P < 0.05$ , as determined by Student's *t* test). \*, no CFU could be detected (detection limit, 100 CFU/ml).

nant wild-type, Met<sub>101</sub>Ile, or Met<sub>133</sub>Ile AhpC proteins dissolved in 10 mM NaH<sub>2</sub>PO<sub>4</sub> buffer (pH 7.5) to a final concentration of ~0.15 mg/ml were used for the Far-UV CD analysis at from 185 to 260 nm at 20°C in a 0.1-cm-gap cuvette. The molar ellipticity values ( $\theta$ ) were calculated using a molar concentration of 6.4  $\mu\text{M}$  and 206 amino acids.

## RESULTS

***H. pylori*  $\Delta msr$  mutants are more sensitive to HOCl.** Previous studies in our laboratory showed that nongrowing *H. pylori*  $\Delta msr$  mutants were more sensitive to a variety of oxidants (such as O<sub>2</sub>, H<sub>2</sub>O<sub>2</sub>, and GSNO) than the wild type (17). In addition, those mutants were also more sensitive to neutrophil-mediated killing (10). Neutrophils produce HOCl via their robust myeloperoxidase, so the increased sensitivity of the  $\Delta msr$  mutants was possibly due to an inability to repair HOCl-induced Met-SO residues. In the current study, we aimed at complementing our previous findings by comparing the sensitivities of nongrowing *H. pylori* wild-type and  $\Delta msr$  mutant cells to HOCl. Wild-type and  $\Delta msr$  mutant cells were grown on plates, resuspended in PBS, and mixed with increasing HOCl concentrations before being serially diluted and plated. CFU counts indicated that  $\Delta msr$  mutants were significantly more sensitive to HOCl than the parental strain (Fig. 1).

**AhpC inactivation by HOCl can be repaired by Msr.** Among the methionine-rich proteins known to be affected by HOCl treatment in the  $\Delta msr$  mutant strain are two proteins involved in oxidative stress combat, catalase and Trx1 (14). This finding led us to consider a third oxidative-stress-combating protein, AhpC, as a likely target for methionine oxidation. Indeed, AhpC has a high methionine content (more than 3% of its residues). It is also an abundant protein, making up approximately 3% of the total protein content, according to our own densitometry analysis of resolved peptide bands after gel electrophoresis resolution (data not shown). Therefore, we sought to study further a possible link between oxidative stress and Met oxidation in AhpC. To investigate the functionality of AhpC before and after oxidative repair, we analyzed the peroxide reductase activities of various purified

AhpC preparations by following TrxR-dependent NADPH oxidation (Table 1). AhpC was expressed as a hexahistidine-tagged recombinant protein in *E. coli* and purified to a level close to homogeneity (data not shown). Purified AhpC was divided in three pools: (i) as purified, (ii) oxidized with HOCl, and (iii) HOCl oxidized and subsequently Msr repaired. AhpC-dependent reduction of H<sub>2</sub>O<sub>2</sub> was observed with purified (unoxidized) AhpC; however, treatment of pure AhpC enzyme with excess HOCl abolished the reductase's activity (Table 1). Upon incubation with an enzyme repair mixture requiring Msr, oxidized AhpC fully recovered peroxide reductase activity (at a level similar to that seen with the nonoxidized sample). As a control, the presence of Msr did not affect the peroxidase enzyme activity of the nonoxidized AhpC (Table 1). Also, a Msr repair mixture control had negligible peroxidase activity (approximately 2 nmol NADPH/min/mg of Msr). Taken together, our results strongly suggest that HOCl-oxidized AhpC can be fully repaired by the Msr methionine sulfoxide repair enzyme.

**Identification of oxidized and Msr-repaired Met residues in *H. pylori* AhpC.** The primary sequence of the AhpC protein (HP1563 in *H. pylori* strain 26695) contains 198 amino acids, including 6 Met residues (Met<sub>1</sub>, Met<sub>101</sub>, Met<sub>133</sub>, Met<sub>152</sub>, Met<sub>155</sub>, and Met<sub>180</sub>). Each of the three AhpC pools defined above (unoxidized, oxidized, and oxidized and then repaired) and our control (Msr-incubated unoxidized AhpC) were digested with GluC (protease), with peptide identification confirmed by MS/MS (Fig. 2). Oxidation of residues Met<sub>152</sub> and Met<sub>155</sub>, which were located on the same peptide, was determined by LC-MS/MS of the oxidized peptide to determine the fraction of oxidation of that peptide measured by LC-MS to be assigned to each methionine on the peptide. All the other identified Met residues were distributed on different peptides. We identified residue Met<sub>101</sub> as the Met residue with the highest oxidation rate as well as the highest repair rate by Msr, followed by Met<sub>133</sub> (Fig. 2). Only partial Msr-mediated Met repair was observed for Met<sub>155</sub> or Met<sub>180</sub>.

**Met<sub>101</sub> and Met<sub>133</sub> residues are critical for AhpC activity both *in vivo* and *in vitro*.** Since Met<sub>101</sub> and Met<sub>133</sub> residues were found to be readily oxidized and repaired by Msr (Fig. 2), we hypothesized that both residues might be important for the stability and/or activity of AhpC. Therefore, both Met residues were independently or concomitantly replaced with an isoleucine (I) residue by site-directed mutagenesis. Each construct was cloned into pET21b and introduced into *E. coli* BL21(DE3)RIL host cells to

**TABLE 1** AhpC activity of native and variant AhpC proteins<sup>a</sup>

AhpC protein	AhpC activity (nmol NADPH oxidized/min/mg protein)
Native†	216 ± 22
Native, oxidized†	5 ± 5
Native, oxidized, Msr added†	242 ± 5
Native, Msr added†	244 ± 22
Native‡	144 ± 22
Variant M101I‡	23 ± 7
Variant M133I‡	57 ± 15
Variants M101I and M133I‡	8 ± 7

<sup>a</sup> AhpC activity was measured in a coupled assay by monitoring thioredoxin reductase-dependent oxidation of NADPH at 340 nm. "native" refers to the protein as purified from *E. coli*. † and ‡ indicate 2 independent sets of experiments done with different batches of native AhpC, Trx1, and TrxR. Data represent means and standard deviations from 4 replicates (†) or 10 to 15 replicates (‡).

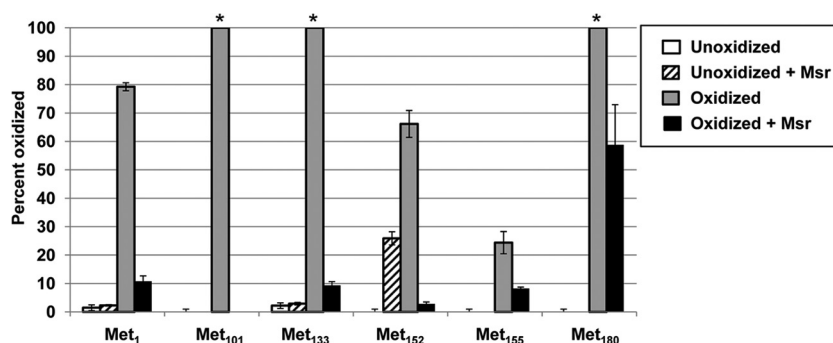


FIG 2 Identification and quantification of HOCl-oxidized and Msr-repaired methionine residues in *H. pylori* AhpC by LC-MS/MS. AhpC was oxidized with a 60-fold molar excess of HOCl and subsequently repaired (or not) by Msr. Repaired and unrepaired AhpC samples were subjected to in-solution GluC digestion, and Met residues were analyzed and quantified by LC-MS/MS. The values corresponding to the percentages of oxidation (y axis) correspond to the ratios of MetO residues to the total number of times the peptide was found (Met coverage). Error bars represent 1 standard deviation (from three measurements). \*, the unoxidized peptide was below our limit of detection.

allow T7-based expression in the presence of the IPTG inducer. First, cells expressing each AhpC variant were characterized for their sensitivity to *t*-butyl hydroperoxide (*t*-BH) (Table 2). Although *E. coli* host cells also have their own AhpC, we hypothesized that expression of active recombinant *H. pylori* AhpC should confer to them increased resistance to the chemical. While there was no difference between the inhibition zones in the absence of IPTG, upon addition of 0.1 mM IPTG in the medium, *E. coli* cells expressing native AhpC were indeed more resistant to *t*-BH (smaller inhibition zone) than *E. coli* cells with pET21b vector only (Table 2). Interestingly, *E. coli* cells expressing M<sub>101</sub>I, M<sub>133</sub>I, or M<sub>101</sub>I-M<sub>133</sub>I AhpC variant proteins were more sensitive to *t*-BH than cells expressing native AhpC. This phenotype was not due to differences in protein expression, as wild-type and mutant AhpC proteins were expressed to similar levels in the presence of 0.1 mM IPTG, as seen by SDS-PAGE (data not shown). Rather, these results indicate that both Met<sub>101</sub> and Met<sub>133</sub> residues are important for alkyl hydroperoxide detoxification *in vivo*.

The importance of both Met residues was confirmed *in vitro*, by measuring the AhpC activity of each purified AhpC variant protein. M<sub>101</sub>I, M<sub>133</sub>I, and M<sub>101</sub>I-M<sub>133</sub>I AhpC proteins were independently purified to a level close to homogeneity on a nickel-nitrilotriacetic acid (Ni-NTA) column (data not shown) and assayed by monitoring the TrxR-dependent oxidation of NADPH at

340 nm, following the exact same protocol as for native AhpC (Table 1). The levels of peroxide reductase activity of M<sub>101</sub>I, M<sub>133</sub>I, and M<sub>101</sub>I-M<sub>133</sub>I were 16%, 40%, and 6% of wild-type activity, respectively, confirming that the presence of both residues—individually and together—was critical (Table 1). To determine whether the AhpC protein structure was affected by the methionine substitutions, nonreducing SDS-PAGE and CD analyses were conducted. Both the M<sub>101</sub>I and M<sub>133</sub>I single variants as well as the double variant were able to form dimers as well as higher-molecular-mass complexes (like native AhpC), as seen by nonreducing SDS-PAGE (Fig. 3). CD analysis (Fig. 4) indicated a slight increase in helicity and decrease in total  $\beta$ -structure for M<sub>101</sub>I and a slight decrease in helicity and increase in total  $\beta$ -structure for M<sub>133</sub>I. Overall, these changes are minor and are highly unlikely to indicate a gross destabilization of the structure of these variants. Rather, it appears that these Met residues are important for maintaining catalytic activity.

## DISCUSSION

As part of the host defense system against invading microorganisms, neutrophils produce HOCl via the enzyme myeloperoxidase (23) and Met residues are considered primary amino acid targets

TABLE 2 Diameter of zone of inhibition of *E. coli* cells expressing *H. pylori* native or variant AhpC proteins<sup>a</sup>

Plasmid	Diam of zone of inhibition (mm)	
	No IPTG	0.1 mM IPTG
pET21b (vector control)	50 ± 4	52 ± 4
pET-AhpC	48 ± 1	41 ± 2
pET-AhpC.M <sub>101</sub> I	49 ± 1	53 ± 3
pET-AhpC.M <sub>133</sub> I	51 ± 1	58 ± 5
pET-AhpC.M <sub>101</sub> I.M <sub>133</sub> I	49 ± 1	55 ± 2

<sup>a</sup> *E. coli* BL21 RIL cells transformed with the appropriate plasmid were mixed with top agarose and spread on top of LB plates (ampicillin [Amp], chloramphenicol [Cm]) supplemented with 0.1 mM IPTG when indicated. Sterile paper disks (7.5-mm diameter) with *t*-BH were placed in the center of each plate, and the diameter of the inhibition zone was recorded after 24 h of growth. Results shown represent means and standard deviations of the results of measurements done on 3 to 5 plates, and the inhibition zone for pET-AhpC with IPTG was statistically significantly smaller than for all the other strains ( $P < 0.05$ , Student's *t* test).

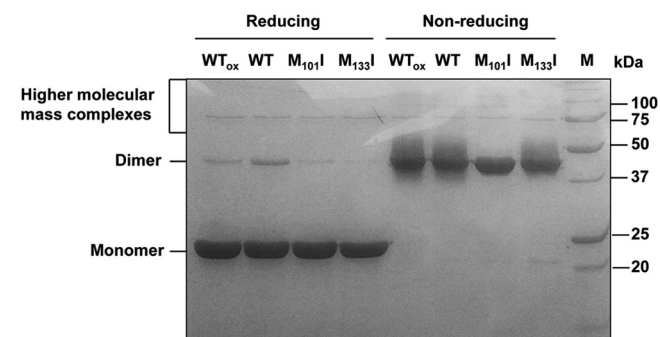


FIG 3 SDS-PAGE of native and variant AhpC proteins. Oxidized native (wild-type) AhpC (WT<sub>ox</sub>), nonoxidized native AhpC (WT), or variant AhpC (M<sub>101</sub>I, M<sub>133</sub>I) (2.5  $\mu$ g per lane) proteins were mixed with reducing buffer and heated at 95°C for 5 min or mixed with nonreducing loading buffer (and not heated) before being separated on an SDS-12.5% polyacrylamide gel. A molecular mass standard (M) is shown on the right, with sizes indicated on the right. Calculated (theoretical) molecular mass of AhpC-His<sub>6</sub>, 23.3 kDa.

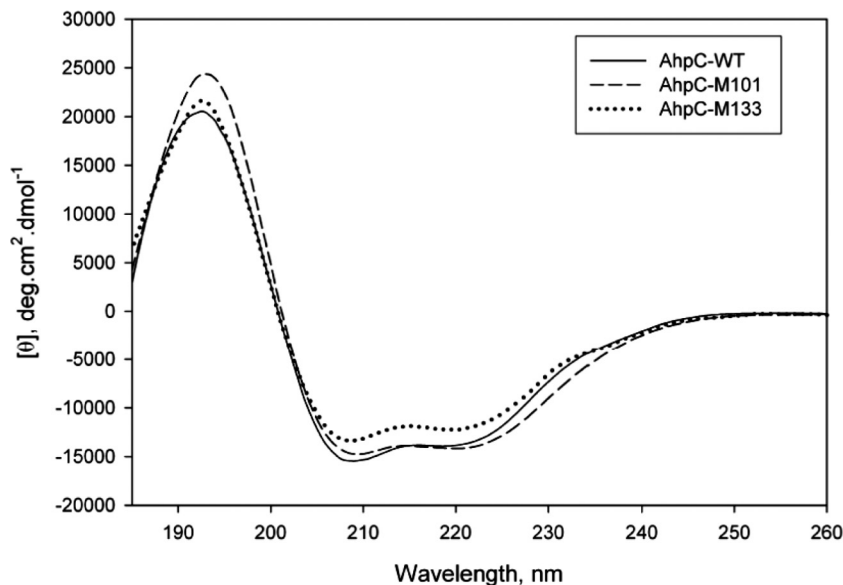


FIG 4 CD spectra of purified native,  $M_{101}I$  variant, or  $M_{133}I$  variant AhpC proteins. CD spectra of each purified protein were measured using an Aviv 400 spectropolarimeter in a 0.1-cm-gap cuvette from 185 to 260 nm at 20°C. Mean residue molar ellipticities ( $\theta$ ) were calculated using a molar concentration of 6.4  $\mu$ M and 206 amino acids.

of HOCl damage via conversion to Met sulfoxide (Met-SO) (24). These Met-SO residues can be repaired by Msr. In the current report, we show that nongrowing *H. pylori*  $\Delta$ msr mutants are more sensitive to HOCl than the parental strain. This phenotype reflects the inability of  $\Delta$ msr mutants to repair Met-SO residues in key enzymes, especially enzymes needed to fight against HOCl-generated oxidative stress. Indeed, previous results from our laboratory revealed that catalase and Trx1 were among the proteins most oxidized in  $\Delta$ msr mutants subjected to oxidative stress (14). Furthermore, *H. pylori* catalase is inactivated by HOCl treatment but can be repaired upon incubation with purified Msr and MS/MS tandem analysis revealed those Met residues specifically oxidized by HOCl and repaired by Msr (10). Repair of catalase by Msr suggests that a physical interaction occurs between the two proteins: Msr and catalase can indeed interact, as shown by cross-linking (14).

In light of these findings, we sought to investigate whether AhpC (HP1563 in *H. pylori* strain 26695 [16]), another key enzyme in the fight against oxidative stress, would be oxidized and then subsequently repaired by Msr. Indeed, its high methionine content as well as its overall abundance in *H. pylori* makes AhpC a prime candidate for methionine oxidation. AhpC is a member of the 2-Cys peroxiredoxin (Prx) family (25) whose catalytic activity involves two conserved cysteine residues. It is commonly accepted that reduced AhpC detoxifies  $H_2O_2$  or organic peroxides (such as lipid hydroperoxides) by converting them into their corresponding alcohols (26). However, new findings suggest that the role of AhpC might not be solely limited to peroxide detoxification: for instance, AhpC was recently shown to be also required for ferric ion transport and the optimal production of enterobactin in *E. coli* (27). Interestingly, *H. pylori* AhpC has been shown to have dual functions: depending on the duration of the oxidative stress, it can switch from a low-molecular-weight (LMW) complex with peroxide reductase activity (short-term oxidative stress) to a high-molecular-weight (HMW) complex with chaperone-like activity (short-term oxidative stress) (28). In addition, *HpAhpC* differs

significantly from other prokaryotes' AhpCs: according to Chuang and coworkers, the sequence of *HpAhpC* is more homologous to those of mammalian Prxs than to those of other eubacterial AhpC (21). *HpAhpC* has also been shown to have higher catalytic activity toward  $H_2O_2$  than other AhpC homologs do (20). Nevertheless, *HpAhpC* contains two highly conserved cysteine residues (Cys<sub>49</sub> and Cys<sub>169</sub>) which are the hallmarks of the Cys-2 Prx protein family. AhpC is one of the most abundant proteins synthesized by the gastric pathogen. The proteome analysis by Jungblut et al. (29) showed that AhpC (also called TsaA) is the third-most-abundant protein in *H. pylori* (29); according to our own densitometry analysis using crude extracts as well as known amounts of purified AhpC protein, we estimated that the protein makes up approximately 3% of the total protein content. Its abundance is reflected by the fact that *H. pylori* mutants with one-third-lower *ahpC* transcript levels (as controlled by antisense RNA technology) can still colonize mouse stomachs (30), even though *H. pylori* *ahpC* mutants fail to colonize, as shown by a different study (31).

Treatment of purified AhpC by HOCl led to full inactivation of the enzyme, which could be restored upon incubation with purified Msr, as previously shown for catalase (10) and UreG (17). When the two most oxidized and successfully Msr-repaired Met residues, Met<sub>101</sub> and Met<sub>133</sub>, were independently or concomitantly replaced with isoleucine, the AhpC enzyme activity of the variant proteins was significantly decreased, and those recombinant AhpC variant enzymes confer lower *t*-BH resistance to *E. coli* cells. While these mutations are not in the vicinity of the essential Cys<sub>49</sub> and Cys<sub>169</sub> cysteine residues, there was a possibility that the overall conformation of AhpC would be disturbed. However, when run on nonreducing SDS-polyacrylamide gels, both single variants as well as the  $M_{101}I$ - $M_{133}I$  double variant formed dimers and higher-molecular-mass complexes, similar to native AhpC (Fig. 3), suggesting that the oligomerization capacity is not affected in these variants. In addition, CD analysis confirmed there

was only a slight modification of the secondary structure in these variant proteins (Fig. 4). Therefore, one has to conclude that the loss of activity in these variants is not related to major structural alterations. The involvement of these methionine residues in AhpC activity is yet to be understood.

So far, only a few proteins have been shown to be repaired by Msr in bacteria. In *H. pylori*, the list now includes GroEL, SSR (site-specific recombinase), catalase, UreG, and AhpC (10, 14, 18). In *E. coli*, Msr was found to protect GroEL and Ffh from oxidative damage (32, 33). As the list expands, so does our knowledge on Msr target specificity; this will eventually lead to a better understanding of Msr-mediated repair mechanisms. Finally, methionine oxidation of a protein is usually considered detrimental to the protein's activity; however, a recent study from Drazic and coworkers actually suggested otherwise: those authors showed that methionine oxidation activates a hypochloride-responsive transcription factor (HypT) in *E. coli* (34). In this case, Msr repair of the oxidized HypT protein leads to an inactive protein. This finding suggests that Msr can almost be considered a master regulator at the posttranslational level, able to reactivate but also deactivate key enzyme activities in the cell.

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