Cationic antimicrobial peptides serve as activation signals for the *Salmonella* Typhimurium PhoPQ and PmrAB regulons *in vitro* and *in vivo*

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*Salmonella enterica* serovar Typhimurium uses two-component regulatory systems (TCRSs) to respond to environmental stimuli. Upon infection, the TCRSs PhoP-PhoQ (PhoPQ) and PmrA-PmrB (PmrAB) are activated by environmental signals detected in the lumen of the intestine and within host cells. TCRS-mediated gene expression leads to upregulation of genes involved in lipopolysaccharide (LPS) modification and cationic antimicrobial peptide (CAMP) resistance. This research expands on previous studies which have shown that CAMPs can activate *Salmonella* TCRSs *in vitro*. The focus of this work was to determine if CAMPs can act as environmental signals for PhoPQ- and PmrAB-mediated gene expression *in vitro*, during infection of macrophages and in a mouse model of infection. Monitoring of PhoPQ and PmrAB activation using recombinase-based *in vivo* expression technology (RIVET), alkaline phosphatase and β-galactosidase reporter fusion constructs demonstrated that *S. Typhimurium* PhoQ can sense CAMPs *in vitro*. In mouse macrophages, the cathelicidin CRAMP does not activate the PhoPQ regulon. Acidification of the *Salmonella*-containing vacuole activates PhoP- and PmrA-regulated loci but blocking acidification still does not reveal a role for CRAMP in TCRS activation in mouse macrophages. However, assays performed in susceptible wild type (WT), CRAMP knockout (KO), and matrilysin (a metalloproteinase necessary for activating murine α-defensins) KO mice suggest CRAMP, but not α-defensins, serve as a putative direct TCRS activation signal in the mouse intestine. These studies provide a better understanding of the *in vivo* environments that result in activation of these virulence-associated TCRSs.

**Keywords:** *Salmonella* Typhimurium, CAMPs, PhoPQ, PmrAB, lipopolysaccharide modification

INTRODUCTION

Upon infection, bacteria must evade the arsenal of host immune defenses, including cationic antimicrobial peptides (CAMPs), designed to eliminate invading pathogens. CAMPs are structurally diverse innate immune molecules providing protection against infection for all classes of life (Menendez and Brett Finlay, 2007). CAMPs are amphipathic peptides that are classified based on their secondary structure, and can be separated into categories including α- or β-defensins and cathelicidins. The antimicrobial activity of CAMPs comes from the ability of these molecules to insert into the microbial membrane, resulting in membrane destabilization and microbial lysis (Matsuizaki et al., 1997; Radek and Gallo, 2007; Bucki et al., 2010). These peptides are rapidly produced by macrophages and epithelial cells in response to infection or injury and can mediate inflammation and stimulate the immune system upon detection of pathogens (Menendez and Brett Finlay, 2007).

The Gram-negative enteric pathogen *Salmonella enterica* serovar Typhimurium (S. Typhimurium) can use two-component regulatory systems (TCRSs), such as PhoP-PhoQ (PhoPQ) and PmrA-PmrB (PmrAB), to detect environmental signals and to mediate changes in gene expression that promote survival and virulence (Miller et al., 1989; Roland et al., 1993; Gunn and Miller, 1996). Indirect activation of PmrAB occurs through the PhoPQ TCRS. PhoPQ activates the expression of PmrD, which can regulate PmrA activity at the post-transcriptional level by binding to and stabilizing PmrA in its phosphorylated form (Kox et al., 2000; Kato and Groisman, 2004). In addition to known *in vitro* signals such as low magnesium (Mg<sup>2+</sup>), acidic pH and high concentrations of iron (Fe<sup>3+</sup>), *S. Typhimurium* PhoPQ and PmrAB and their regulons are also activated by unknown environmental signals in macrophages, in other host cells and in the intestinal lumen (Foster and Hall, 1990; Alpuche Aranda et al., 1992; Garcia Vescovi et al., 1996; Pearson et al., 1998; Wosten et al., 2000; Merighi et al., 2005).

CAMPs also have been shown to activate PhoPQ *in vitro* and may be *in vivo* factors involved in *Salmonella* TCRS-mediated gene expression and LPS modification (Bader et al., 2003). Several CAMPs bind to acidic patches on the inner membrane-facing region of the PhoQ periplasmic domain, resulting in PhoQ conformational changes and activation of the PhoPQ regulon (Bader et al., 2005). Therefore, *Salmonella* may sense and respond to CAMPs in the environment and during infection through PhoQ to prevent killing by these host molecules and other...
immune defenses (Bader et al., 2003, 2005). PmrAB-regulated modification of LPS lipid A with positively-charged molecules such as aminoarabinose (Ara4N) and phosphoethanolamine promotes CAMP resistance by reducing the anionic charge of the bacterial outer membrane (Gunn et al., 2000; Tamayo et al., 2005). PhoPQ-regulated lipid A modification also promotes resistance to CAMP killing.

Merighi et al. used recombinase-based in vivo expression technology (RIVET) analysis to examine in vivo expression of PhoP-and PmrA-regulated genes (Merighi et al., 2005). These authors found that the PhoPQ-regulated gene pagP [palmitoyl transferase that mediates palmitate addition to Salmonella lipoid A (Belden and Miller, 1994; Guo et al., 1998)] and the PmrB-regulated gene pmrH [first gene in the seven gene operon involved in Ara4N addition to the lipid A (Gunn et al., 1998b, 2000)] are expressed early during Salmonella infection in response to unknown factors in the in vivo environment. Known in vitro activating signals, acidic pH and high iron concentrations, were not responsible for TCRS-mediated gene activation (Merighi et al., 2005). These authors also showed that TCRS-mediated gene activation in vivo requires the presence of active PmrA and PhoP, as pmrA and phoP mutants did not express pmrH in response to the in vivo environment (Merighi et al., 2005).

The complex interplay between host immune factors and bacterial defense systems during the early stages of Salmonella infection is still poorly understood. CAMPs are likely to be one of the earliest-encountered components of the immune system. They protect the host against infection both directly, through potent bactericidal activity, and indirectly, by inducing chemotaxis of monocytes and neutrophils to the site of infection. Bader et al. proposed a model in which CAMP detection by Salmonella leads to TCRS-induced signaling that could result in regulation of virulence genes, as well as increased resistance to these innate immune molecules and other host defenses through LPS modification (Bader et al., 2003, 2005). We further hypothesize that S. Typhimurium is able to detect and respond to sublethal levels of CAMPs through the PhoPQ/PmrAB TCRS in vivo, in the lumen of the intestine and within host cells.

MATERIALS AND METHODS

BACTERIAL STRAINS AND GROWTH CONDITIONS

S. Typhimurium strains used in this study are listed in Table 1. The pmrl::MudJ mutant strains listed in Table 1 were originally generated by random MudJ mutagenesis in a pmrlA background (as previously described by Tamayo et al.) and then were transduced into WT and PhoP strains of S. Typhimurium to create the pmrl::MudJ and pmrl::MudJ PhoP strains used in this study (Tamayo et al., 2002). Luria-Bertani (LB) broth and agar were used for strain maintenance and experimentation. When appropriate, antibiotics were added at the following concentrations: ampicillin, 50 μg/ml; chloramphenicol, 25 μg/ml; kanamycin, 50 μg/ml; tetracycline, 15 μg/ml. Two of the CAMPs tested were murine CRAMP and the human cathelicidin ortholog, LL-37 (Gudmundsson et al., 1996; Gallo et al., 1997). Other molecules of interest were the cationic lipopeptide polymyxin B (PMB), which targets Gram-negative bacteria in a manner similar to CAMPs, and a PMB derivative, polymyxin B nonapeptide (PMBN) (Vaara and Vaara, 1983). PMBN is used to examine gene activation by cationic molecules, but it cannot penetrate bacterial membranes or cause cell death due to the lack of a fatty acid tail (Vaara and Vaara, 1983). A range of peptide concentrations was used initially for all in vitro assays and concentrations of 0.5–5 μg/ml of PMBN, CRAMP and LL-37 showed similar results. Results from 5 μg/ml (5 μM PMBN, 1.3 μM CRAMP, and 1.1 μM LL-37) were reported in most of the assays as this represented the highest, yet non-lethal concentration in this range. For PMB, a range of 0.01–0.1 μg/ml was tested and all showed similar results. Results from 0.06 μg/ml (0.04 μM) were reported in the assays in this work.

ALKALINE PHOSPHATASE (AP) DETECTION OF PhoPQ ACTIVATION

S. Typhimurium AP reporter strains capable of displaying AP activity when TnphoA-containing gene products are translated [pagI::TnphoA (JSG162), pagI::TnphoA PhoP (JSG163), pagP::TnphoA (JSG174), and phoN::TnphoA (JSG2860)] were used to measure PhoPQ two-component system activation following CAMP exposure. S. Typhimurium strains were grown in a rotating drum at 37°C with aeration to early log phase (all cultures were normalized to an optical density at 600 nm [OD600] of 0.2), and incubated with 1.1 μM LL-37 (AnaSpec, Fremont, CA), 1.3 μM CRAMP (Anaspec), 0.04 μM PMB (Sigma-Aldrich, St. Louis, MO), or 5 μM PMBN (Sigma-Aldrich) for 90 min at 37°C with aeration to induce PhoPQ-mediated gene expression. These methods were based on the protocol described by Bader et al. (2005). After incubation with the CAMPs, a standard AP assay protocol was followed. Miller units, representing relative AP activity, were calculated according to the following equation: Miller units = (1000 × OD420)/(OD600 × Sample volume (0.1 mL) × Time lapsed until color change).

β-GALACTOSIDASE DETECTION OF PmrAB ACTIVITY

CAMP-induced activation of PmrAB-regulated gene expression was quantified using S. Typhimurium MudJ reporter fusion strains (JSG1051, JSG1070 and JSG1071). S. Typhimurium strains were grown as for the AP assays and incubated with 1.1 μM LL-37, 1.3 μM CRAMP, 0.04 μM PMB or 5 μM PMBN for 90 min at 37°C with aeration to induce PmrAB-mediated gene expression. β-galactosidase assays were carried out using a spectrophotometric method with ortho-nitrophenyl-β-galactoside (ONPG) as a substrate. Miller units, representing relative β-galactosidase activity, were calculated as for the AP assays.

RIVET in vitro ASSAYS

The S. Typhimurium RIVET strains (pagP [JSG2502], pmrl [JSG2428] or fepA [control, JSG2579]) were used to measure PhoPQ and PmrAB-mediated gene activation by the heritable loss of bacterial Tet resistance. The RIVET strains were grown as described for the AP assays and then were incubated for 4 h or 24 h in the presence or absence of sublethal concentrations of LL-37 (1.1 μM), CRAMP (1.3 μM) or PMB (0.04 μM), as well as low (10 μM) or high (10 mM) Mg2+, and enumeration plating was employed to determine bacterial survival. Resolution of antibiotic resistance cassettes from RIVET strains was quantified through patching individual colonies (n ≤ 100) onto LB or LBtet and calculating percent sensitivity.
RIVET assay in murine macrophages
All tissue culture experiments were incubated at 37°C plus 5% CO₂. Bone marrow-derived macrophages (BMDMs) were obtained from the femurs of WT (BALB/c) and background-matched CRAMP knockout (KO) mice and were cultured in Iscove’s Modified Dulbecco’s Medium ([IMDM], Gibco, Grand Island, NY) containing 10% heat-inactivated fetal bovine serum (FBS), 50% L929-conditioned IMDM, 0.6% non-essential amino acids (Gibco), 100 U/ml penicillin, and 100 µg/ml streptomycin (Gibco) at 37°C in a humidified incubator containing 5% CO₂ for 5 days to allow for monocytes to differentiate into macrophages (Swanson and Isberg, 1993; Stanley, 1997; Amer and Swanson, 2005; Abdelaziz et al., 2011). After 5 days, BMDMs were counted and 2 × 10⁵ cells were seeded into each well of a 24-well tissue culture plates (BD Falcon, Franklin Lakes, NJ), and incubated overnight to allow for BMDM adherence. BMDMs were then infected with S. Typhimurium grown in a rotating drum at 37°C with aeration to late log phase (cultures were adjusted to an OD₆₀₀ of 0.9) at an MOI of 10:1 (bacteria:macrophage). For experiments with bafilomycin, bafilomycin A1 from Streptomyces griseus (Sigma-Aldrich) was added to macrophage wells at a final concentration of 100 nM. For experiments with ammonium chloride [NH₄Cl (Sigma-Aldrich)], it was resuspended in IMDM and added to macrophage wells at a final concentration of 10 mM. Tissue culture wells were centrifuged briefly following the addition of bacteria to synchronize infection. BMDMs were incubated with bacteria at 37°C for 2 h to allow adequate time for phagocytosis of S. Typhimurium, and media was then removed from each well and replaced with IMDM plus 10% FBS containing either 50 µg/ml or 10 µg/ml gentamicin (Gibco) to kill extracellular S. Typhimurium. Wells that received 50 µg/ml gentamicin represent the initial infection timepoint and were incubated for an additional 30 min to allow for killing of extracellular bacteria, after which the BMDMs were processed for enumeration of intracellular S. Typhimurium. Wells that received 10 µg/ml of gentamicin were processed for enumeration of intracellular S. Typhimurium after 6, 10 and 24 h of S. Typhimurium exposure. BMDMs were lysed by 0.01% Triton-X-100. Intracellular survival was determined by enumeration of macrophage lysates on LB agar. Gene activation of RIVET strains was calculated as a percentage of colonies having resolved the Tet cassette as described previously. The percentage of activation in medium alone was subtracted from the percentage of Tet-sensitive colonies recovered from macrophage lysates at each time point.

MOUSE ASSAYS
Experiments with WT and CRAMP KO mice were performed in 6–10 week old female BALB/c (WT) mice from Harlan Sprague Dawley (Indianapolis, IN), as well as background-matched CRAMP KO mice obtained from Dr. Bradford McGwire at The Ohio State University. Female C57BL/6 mice and B6.129-Mmp7 (MMP7 KO) mice (6–10 weeks old) were purchased from The Jackson Laboratories (Bar Harbor, ME). Mice were used in accordance with guidelines established by The Ohio State University Institutional Animal Care and Use Committee (IACUC). Food and water were removed from the mouse cages 4 h prior to infection. Mice were orally infected (in triplicate) with 1 × 10⁸ colony forming units (CFU) S. Typhimurium in 100 µl PBS. After infection, food and water were returned to the cages. Mice were euthanized at 4, 12, 24 and 48 h post-infection (p.i.) For experiments in BALB/c and CRAMP KO mice, the intestine lumen contents, 2–4 Peyer’s patches near the distal ileum and the spleen were removed from each mouse and homogenized in 1 mL of cold PBS. For experiments in C57BL/6 and MMP7 KO mice, 2–3 inches of the small intestine closest to the caecum (distal ileum) and the spleen were removed from each mouse, homogenized in PBS diluted, plated, incubated and patched as above (see RIVET Assay in Murine Macrophages). The percentage of Tet-sensitive colonies obtained from each sample/condition was calculated to determine the level of in vivo promoter activation.

STATISTICS
For all experiments, each condition was analyzed in duplicate or triplicate in 2–5 independent experiments. The results obtained

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Table 1 | Salmonella enterica serovar Typhimurium strains.

| Strain | Relevant characteristics | Source or reference |
|--------|--------------------------|---------------------|
| JSG162 | CS019; pagP::TnphoA (Kan/Cam) | Belden and Miller, 1994 |
| JSG163 | 14028s; pagP::TnphoA phoP105::Tn10d (Kan/Tet) | Belden and Miller, 1994 |
| JSG174 | CS019; pagP::TnphoA (Kan/Cam) | Belden and Miller, 1994 |
| JSG2680 | phoP::TnphoA (Kan) | Miller et al., 1989 |
| JSG051 | pmrI::MudJ | This study |
| JSG1070 | pmrI::MudJ PhoP⁻ | Miller et al., 1989 |
| JSG1071 | pmrC::MudJ | This study |
| JSG2428 | pmrH::tnpR RIVET strain; JSG246 + pJSG2413 cointegrated at pmrH→ PhipmrH⁻-tnpRmut135-lacZ+J666 res1-tet-res1 = akaMM566 (Amp/Tet) | Merighi et al., 2005 |
| JSG2502 | pagP::tnpRRIVET strain; JSG246 + pJSG2483; fipagP⁻-tnpRmut135-lacZ+J608-6 res1-tet-res1 (Amp/Tet) | Merighi et al., 2005 |
| JSG2579 | fepA::tnpRRIVET strain; JSG246 + pJSG2535; Phi(fepA⁻-tnpRmut135-lacZ+J2535 res1-tet-res1 (Amp/Tet) | Merighi et al., 2005 |

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Richard et al. CAMPs activate Salmonella TCRS

July 2012 | Volume 2 | Article 102 | 3
were averaged and then analyzed for statistical significance using the two-tailed Student’s *t*-test.

**RESULTS**

**CATIONIC ANTIMICROBIAL PEPTIDES DIFFERENTIALLY ACTIVATE THE Salmonella PhoPQ AND PmrAB REGULONS in vitro**

Previous research has shown that certain CAMPs can activate the expression of PhoP-regulated genes (Bader et al., 2003, 2005; Shprung et al., 2012). To further examine Salmonella PhoPQ-mediated CAMP activation in vitro, we performed AP assays with S. Typhimurium strains containing a *phoA* (AP) reporter fusion to PhoP-regulated genes *phoN* and *pagJ* (Kier et al., 1979; Miller et al., 1989; Belden and Miller, 1994; Gunn et al., 1998a). The S. Typhimurium AP reporter strains were grown in the presence or absence of PMB, PMBN, CRAMP or LL-37 for 90 min. A significant increase in PhoPQ-regulated promoter expression was observed when S. Typhimurium was grown in the presence of all four peptides (*phoN* increased with LL-37 but not significantly), compared to growth in the absence of each CAMP (Figure 1). PMB and PMBN induced the highest levels of *pagJ* and *phoN* promoter-driven AP activity among the CAMPs examined. Interestingly, *pagJ* and *phoN* show different levels of induction in response to the various CAMPs used. To verify that the observed in vitro CAMP sensing by Salmonella was dependent on PhoP-mediated gene activation, AP assays were also performed with a *pagJ::TnphoA PhoP−* reporter fusion strain (Bader et al., 2003), where AP activity in the presence of peptides was greatly reduced in comparison to the WT *pagJ* reporter strain (Figure 1). These experiments confirm previous work published by Bader et al. demonstrating that specific CAMPs play a role in the expression of the PhoPQ regulon in vitro. In addition, PhoP is required for gene activation in the presence of CAMPs (Bader et al., 2003, 2005).

To measure the ability of CAMPs to activate genes in the PmrAB regulon, β-galactosidase assays were performed with the S. Typhimurium strains containing the *pmr1* or *pmrC* promoter fused to the *lacZ* reporter gene by the MudJ transposon (JSG1051, JSG1070 and JSG1071). PMB, PMBN, CRAMP and LL-37 all were able to induce statistically higher levels of β-galactosidase activity from the *pmr1* promoter compared to S. Typhimurium grown without peptide (Figure 2). The observed *pmr1*-regulated reporter gene activation in the presence of CAMPs was dramatically reduced in the absence of PhoP (Figure 2). The *pmrC::MudJ* reporter strain also showed increased β-galactosidase activity in the presence of PMBN and LL-37 but not the other CAMPs tested (Figure 2). These findings indicate that, as observed with the PhoPQ-regulated genes in the AP assays, PmrAB-regulated genes can be induced in response to CAMPs, but this induction is not uniform or observed with all CAMPs to which *Salmonella* was exposed.

To further confirm that PhoQ can induce TCRS-mediated gene activation in response to CAMPs in vitro, the S. Typhimurium *pagP* and *pmrH* RIVET strains were incubated for 4 h or 24 h (24 h data not shown due to similar trends) in the presence or absence of sublethal concentrations of LL-37, CRAMP and PMB (Gallo et al., 1997). Addition of high (10 mM) or low (10 μM) Mg²⁺ served as a control for appropriate regulation of PhoPQ and PmrAB by a known signal. Exposure to high Mg²⁺ resulted in the absence of Tet-resistance colonies suggesting that there is complete repression of the regulons under these environmental conditions. Results from in vitro experiments performed using the *fepA* control RIVET strain did not show changes in the heritable loss of bacterial Tet resistance (data not shown). In general, the RIVET strains demonstrated little PhoPQ or PmrAB activation in the presence

![Figure 1](image1.png)

**FIGURE 1** | PhoPQ-mediated activation of S. Typhimurium *phoN* and *pagJ* in response to CAMPs. AP assays demonstrate that different CAMPs induce varying levels of PhoP-regulated gene promoter expression. CAMP sensing by PhoQ and subsequent PhoP-mediated AP reporter gene activation is abolished in the absence of PhoP. Values shown represent the mean ± standard deviation from a representative experiment performed at least twice in triplicate. Statistical significance was measured against the samples to which no peptide was added. *Indicates *p* < 0.05. **Indicates *p* < 0.005.

![Figure 2](image2.png)

**FIGURE 2** | TCRS-mediated activation of S. Typhimurium *pmr1* and *pmrC* in response to CAMPs. β-galactosidase assays demonstrate that different CAMPs induce varying levels of PmrA-regulated gene promoter expression and CAMP sensing by PhoQ. Subsequent PmrA-mediated β-galactosidase reporter gene activation is abolished in the absence of PhoP. Values shown represent the mean ± standard deviation from a representative experiment performed at least twice in triplicate. Statistical significance was measured against the samples to which no peptide was added. *Indicates *p* < 0.05. **Indicates *p* < 0.005.
of sublethal concentrations of CAMPs in vitro. The pagP promoter was activated to a higher level than was the pmrH promoter; however, this greater activation was only seen with LL-37 under the conditions tested. LL-37 (1.1 μM) induced the highest level of S. Typhimurium TCRS-mediated signaling compared to the no peptide control, while CRAMP (1.3 μM) and PMB (0.04 μM) did not activate pagP or pmrH significantly in these experiments (Figures 3A–C). The observed greater pagP activation supports the idea that PhoQ is the direct sensor of CAMPs, since PhoQ activation of direct target genes would be stronger than downstream activation of pmrH that occurs through PhoP modulation of PmrA (Gunn et al., 1998b; Kato et al., 2003).

CRAMP IS NOT A SIGNIFICANT MACROPHAGE INTRACELLULAR SIGNAL FOR PhoPQ OR PmrAB ACTIVATION

CRAMP has been shown to be expressed in the intestines and has also been found to be produced by murine macrophages in response to Salmonella infection in vitro (Gallo et al., 1997; Rosenberger et al., 2004). Despite generally low levels of CAMP-mediated RIVET reporter gene activation in vitro, macrophages were infected with the RIVET strains to determine if CAMPs can induce pagP or pmrH expression in host environments that Salmonella is known to encounter during infection. To examine intracellular PhoP- and PmrA-regulated gene activation, BMDMs from WT (BALB/c) and background-matched CRAMP KO mice were infected with the S. Typhimurium RIVET strains at an MOI of 10:1 (bacteria: macrophage). As a control, patching of S. Typhimurium RIVET strains after overnight growth of the initial cultures used for infection confirmed that these strains are not activated in LB or IMDM. As expected, resolution dramatically increased for pagP and pmrH over the 24 h macrophage infection period (Figures 4A,B). Early time points (≤6 h p.i.) surprisingly showed slightly higher resolution in macrophages from CRAMP KO vs. WT macrophages, which was contrary to our hypothesis concerning the involvement of CRAMP in PhoPQ-mediated signaling. At later time points p.i., there were no significant differences in the percentage of Tet-sensitive colonies recovered. In addition, there were no differences in recovery of total bacteria at these time points from CRAMP KO vs. WT macrophages that would account for biases in the results (data not shown). Thus, CRAMP alone does not appear to be responsible for the observed activation of the PhoPQ or PmrAB TCRSs in mouse macrophages. In addition, the equal recovery of bacteria from CRAMP KO vs. WT macrophages demonstrate that CRAMP does not kill S. Typhimurium during macrophage infection.

pH REDUCTION ACTIVATES S. Typhimurium TCRSs INSIDE MURINE MACROPHAGES

Acidic pH is an activating signal for the PhoPQ or PmrAB TCRSs in vitro and should be encountered by S. Typhimurium while inside the macrophage phagosome (Alpuche Aranda et al., 1992). To examine the role of pH as a signal for intramacrophage TCRS activation, as well as its potential synergy with CRAMP, bafilomycin A1 (Baf) or ammonium chloride (NH4Cl) was added to the medium to inhibit the acidification p.i., in the murine macrophage phagosome (Gordon et al., 1980; Lukacs et al., 1990; Tapper and Sundler, 1995).

Again, resolution increased for pagP and pmrH over the 24 h infection period as expected, (Figures 5A,B) but not as dramatically as seen in the experiments in Figure 4. At all time points except 24 h p.i., the addition of Baf resulted in lower recovery
of Tet-sensitive colonies in the \( \text{pagP} \) RIVET fusion strain (i.e., reduced PhoPQ activation; Figure 5A). The addition of Baf to CRAMP KO macrophages showed no additional decrease in fusion activity, consistent with the data in Figure 4 suggesting the lack of a role for CRAMP in intramacrophage PhoPQ activation. Regarding \( \text{pmrH} \) fusion activity, there was no significant difference in the percentage of Tet-sensitive colonies recovered under any of the conditions examined (Figure 5B). The use of NH\(_4\)Cl instead of Baf to abrogate phagosome acidification showed similar trends to the results with Baf (Figure 6A), with the exception that NH\(_4\)Cl treatment dramatically decreased fusion activity for \( \text{pmrH} \) (Figure 6B). Thus, pH plays a role in the induction of \( \text{pagP} \) and \( \text{pmrH} \) gene expression over the course of infection in mouse macrophages, but CRAMP does not appear to play a role in PhoP or PmrA-mediated gene expression even in the absence of phagosome acidification.

**S. Typhimurium CAN SENSE MURINE CRAMP WITHIN THE MOUSE INTESTINE**

Previous work from our lab has demonstrated that *S. Typhimurium* isolated from the lumen of the distal ileum displayed increased levels of \( \text{pmrH} \) and \( \text{pagP} \) expression (Merighi et al., 2005). To expand upon these findings, CAMP-deficient mouse models were used to determine if antimicrobial peptides could serve as environmental factors promoting \( \text{pagP} \) and \( \text{pmrH} \) expression in the small intestine. To examine this, BALB/c (WT) and CRAMP-deficient mouse bone marrow-derived macrophages were pre-treated with bafilomycin in DMSO and infected with the *S. Typhimurium* (A) \( \text{pagP} \) or (B) \( \text{pmrH} \) RIVET strain. Gentamycin-treated macrophages were lysed at 2.5, 6, 12 and 24 h p.i., Lysates containing intracellular bacteria were plated on LB. Resulting colonies were patched onto LB and LB Tet to detect loss of Tet-resistance due to promoter activation. Graphed values represent the mean ± standard deviation of one representative experiment performed at least twice in triplicate. *Indicates \( p < 0.05 \).
at most time points (Figures 7C,D). Thus, while CRAMP did not appear to play a dramatic role in activation of the PhoPQ/PmrAB TCRS in macrophages in vitro, may promote TCRS activation in intestinal/Peyer’s patch tissues during infection.

As observed with the RIVET experiments performed in WT and CRAMP KO mice (Figure 7), the S. Typhimurium pagP promoter exhibited increased activation in both C57BL/6 and background-matched MMP7 KO (matrilysin-deficient) mice over time (Figure 8). The pmrH promoter again showed early activation (as observed in previous tissue/macrophage experiments in this work) that was sustained at later time points. However, these data indicate that Salmonella does not directly respond to active cryptdins in the intestine, as there is no significant difference in RIVET reporter resolution in these mice.

**DISCUSSION**

The role of CAMPs as environmental signals for the expression of *Salmonella* PhoPQ- and PmrAB-regulated genes was examined using reporter fusions in vitro, in macrophages and in mice. Taken together, the results confirm that sublethal concentrations of CAMPs can activate *Salmonella* TCRS-mediated gene expression, as reported by Bader et al. (2003, 2005). This activation

**FIGURE 6 | TCRS-mediated pagP and pmrH activation in NH4Cl-treated WT and CRAMP KO macrophages.** BALB/c and CRAMP-deficient mouse bone marrow-derived macrophages were pre-treated with NH4Cl in IMDM and infected with the *S. Typhimurium* (A) pagP or (B) pmrH RIVET strain. Gentamicin-treated macrophages were lysed at 2.5, 6, 10 and 24 h p.i. Lysates containing intracellular bacteria were plated on LB. Resulting colonies were patched onto LB and LB Tet to detect loss of Tet-resistance due to promoter activation. Graphed values represent the mean ± standard deviation of one representative experiment performed three times in triplicate.

**FIGURE 7 | TCRS-mediated activation of *S. Typhimurium* pagP and pmrH in the WT and CRAMP KO murine intestinal lumen and Peyer’s patches.** BALB/c and CRAMP KO mice were infected orally with 10^8 CFU of the *S. Typhimurium* pagP or pmrH RIVET strains. Mice were sacrificed at 4, 12, 24 and 48 h p.i. for removal of infected organs. Tissue samples were homogenized, diluted, plated on LB and grown overnight at 37°C to recover intracellular bacteria. One hundred colonies recovered from each sample were patched on LB and LB Tet to detect loss of Tet-resistance due to promoter activation in vivo. Graphed values represent the mean percentage ± standard deviation of Tet-sensitive colonies recovered from the (A) and (C) intestinal lumens and (B) and (D) Peyer’s patches of three BALB/c and three CRAMP KO mice infected with the pagP or pmrH RIVET strain for each time point in one representative experiment performed at least twice in triplicate. One asterisk (*) indicates *p* < 0.005 and two asterisks (**) indicate *p* < 0.001.
Richards et al. CAMPs activate Salmonella TCRS

Most likely occurs through the binding of CAMPs to the periplasmic domain of the inner membrane-bound protein kinase PhoQ, resulting in a signal transduction cascade that promotes expression of the PhoPQ and downstream PmrAB regulons (Bader et al., 2003, 2005).

Specific CAMPs were found to play a role in the activation of the PhoPQ and PmrAB regulons in vitro. The AP and β-galactosidase assays showed a significant increase in reporter expression of the PhoPQ and downstream PmrAB regulons (Bader et al., 2003, 2005).

Using the RIVET system, which measures TCRS-mediated gene activation by the heritable loss of bacterial Tet-resistance, was used in this work to expand on the findings of Merighi et al. and determine if PhoQ or PmrB can detect CAMPs in the murine intestine and inside murine macrophages (Merighi et al., 2005).

The RIVET reporters used in this work indicate that LL-37, but not CRAMP and PMB, significantly activates PhoPQ- or PmrAB-regulated virulence genes under the tested experimental conditions. This suggests that the RIVET system may be less sensitive than the other reporter constructs used in this work (AP and β-galactosidase). Despite high sensitivity seen with the in vitro assays, AP and β-galactosidase fusions are less well suited to in vivo experimentation for which the RIVET technique was specifically designed.

Changes in strain survival in CRAMP KO macrophages vs. WT macrophages. Furthermore, we saw no differences in strain survival in CRAMP KO macrophages vs. WT macrophages (data not shown). As acidification of the Salmonella containing vacuole occurs post-phagocytosis, inhibitors of acidification (bafilomycin and NH4Cl) were used to determine if low pH, a known in vitro activating signal of both PhoQ and PmrB, had an effect on gene activation (Perez and Groisman, 2007; Prost and Miller, 2008). The results show a general effect of low pH on activation of pagP and pmrH fusions in macrophages, with bafilomycin treatment having the greatest effect on pagP and NH4Cl had the greatest effect on pmrH. However, consistent with our previous data in this work, CRAMP did not play a role even in the absence of acidification of the vacuole.

While CRAMP is primarily localized in macrophages, it has also been reported to be found in the intestine and intestinal lumen (Gallo et al., 1997). Thus, we examined RIVET fusion induction in Peyers patch tissue and in the intestinal lumen after oral infection of mice. These data demonstrated a general effect of CRAMP in both locations, though not always reaching

**FIGURE 8 | TCRS-mediated activation of S. Typhimurium pagP and pmrH in the WT and Matrilysin-deficient murine intestine.** C57BL/6 and MMP7 KO mice were infected orally with 10^5 CFU of the S. Typhimurium pagP and pmrH RIVET strains. Mice were sacrificed at 6, 24 and 48 h p.i., for removal of two-three inches of the small intestine (measured from the distal ileum). Tissue samples were homogenized, diluted, plated on LB and grown overnight at 37°C to recover intracellular bacteria. One hundred colonies recovered from each sample were patched on LB and LB Tet to detect loss of Tet-resistance due to promoter activation in vivo. Graphed values represent the mean percentage ± standard deviation of Tet-sensitive colonies recovered from the intestine of three BALB/c and three CRAMP KO mice infected with the pagP or pmrH RIVET strain for each time point in one representative experiment.

CRAMP, the murine cathelecidin and human LL-37 ortholog, has previously been implicated in *Salmonella* killing in mouse macrophages, as a PhoP-null strain survived better in CRAMP KO macrophages than WT macrophages (Rosenberger et al., 2004). This data suggests that CRAMP interacts with *Salmonella* within macrophages. However, CRAMP was not found in our studies to be significantly or consistently involved in activation of PhoQ- or PmrB-regulated genes in macrophages. Furthermore, we saw no differences in strain survival in CRAMP KO macrophages vs. WT macrophages (data not shown). As acidification of the *Salmonella* containing vacuole occurs post-phagocytosis, inhibitors of acidification (bafilomycin and NH4Cl) were used to determine if low pH, a known in vitro activating signal of both PhoQ and PmrB, had an effect on gene activation (Perez and Groisman, 2007; Prost and Miller, 2008). The results show a general effect of low pH on activation of pagP and pmrH fusions in macrophages, with bafilomycin treatment having the greatest effect on pagP and NH4Cl had the greatest effect on pmrH. However, consistent with our previous data in this work, CRAMP did not play a role even in the absence of acidification of the vacuole.

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statistical significance or occurring at every time point examined. The variability seen here and with other experiments is often inherent in experiments relying on mouse infections. Thus, CRAMP appears to activate Salmonella pmrH and pagP loci in the intestine, likely prior to entrance into macrophages, and may be the luminal/intestinal signal observed in RIVET experiments reported by Meriggi et al. (2005). This is further supported by the fact that induction in vivo was dependent upon PhoPQ, which is the reported sensory system of CRAMP. It should be noted that the observed gene activation could result from the direct interaction of CRAMP with S. Typhimurium TCRSs and/or induction of other environmental factors or immune responses that are regulated by CRAMP. A complex indirect effect in vivo may explain why CRAMP demonstrated no role in PhoPQ/PmrA activation in vitro in cultured macrophages.

We also examined active intestinal α-defensins (cryptidins) as a potential luminal signal of pagP and pmrH induction. MMP7 KO mice were used in these studies, as this protease is required to cleave the mouse α-defensins from a pro-form to an active form. Similar induction of pagP and pmrH was observed in the intestines of both WT and MMP7 KO mice, suggesting a non-role for the cryptidins in TCRS induction. While it is possible that non-processed cryptidins could be released from Paneth cells and participate in PhoPQ induction, it is unlikely based on our current understanding of the secretion of these CAMPs (Wilson et al., 1999).

Taken together, the results of this project and related findings by other researchers highlight the intricacy, and often redundancy, of S. Typhimurium TCRS-mediated gene activation and LPS modification in response to environmental sensing. Tight control of these bacterial defense mechanisms allows Salmonella to respond quickly and efficiently to constant changes in local stressors and host immune factors, including CAMPs. The findings presented here also highlight the benefit of performing intramacrophage and in vitro experiments, despite weak results in vitro. Media conditions in a test tube cannot represent accurately the complex interplay between host and pathogen. Further investigation into bacterial virulence gene regulation, surface modification and evasion of host defenses will help to determine how pathogens evade killing and thrive in the various hostile environments encountered during infection.

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