**Burkholderia pseudomallei** Evades Nramp1 (Slc11a1)- and NADPH Oxidase-Mediated Killing in Macrophages and Exhibits Nramp1-Dependent Virulence Gene Expression

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Bacterial survival in macrophages can be affected by the natural resistance-associated macrophage protein 1 (Nramp1; also known as solute carrier family 11 member a1 or Slc11a1) which localizes to phagosome membranes and transports divalent cations, including iron. Little is known about the role of Nramp1 in *Burkholderia* infection, in particular whether this differs for pathogenic species like *Burkholderia pseudomallei* causing melioidosis or non-pathogenic species like *Burkholderia thailandensis*. Here we show that transfected macrophages stably expressing wild-type Nramp1 (Nramp1⁺) control the net replication of *B. thailandensis*, but not *B. pseudomallei*. Control of *B. thailandensis* was associated with increased cytokine responses, and could be abrogated by blocking NADPH oxidase-mediated production of reactive oxygen species but not by blocking generation of reactive nitrogen species. The inability of Nramp1⁺ macrophages to control *B. pseudomallei* was associated with rapid escape of bacteria from phagosomes, as indicated by decreased co-localization with LAMP1 compared to *B. thailandensis*. A *B. pseudomallei* bipB mutant impaired in escape from phagosomes was controlled to a greater extent than the parent strain in Nramp1⁺ macrophages, but was also attenuated in Nramp1⁻ cells. Consistent with reduced escape from phagosomes, *B. thailandensis* formed fewer multinucleated giant cells in Nramp1⁺ macrophages at later time points compared to *B. pseudomallei*. *B. pseudomallei* exhibited elevated transcription of virulence-associated genes of Type VI Secretion System cluster 1 (T6SS-1), the Bsa Type III Secretion System (T3SS-3) and the *bimA* gene required for actin-based motility in Nramp1⁺ macrophages. Nramp1⁺ macrophages were found to contain decreased iron levels that may impact on expression of such genes. Our data show that *B. pseudomallei* is able to evade Nramp1- and NADPH
INTRODUCTION

The Natural resistance-associated macrophage protein 1 (Nramp1), also known as solute carrier family 11 member a1 (Slc11a1), acts as a protein-coupled metal transporter for delivering divalent cations (i.e., Fe²⁺, Zn²⁺, and Mn²⁺) across the membrane of late endosomes/lysosomes (Blackwell et al., 2000). One explanation for the ability of Nramp1 to control intracellular bacterial infections is by limiting availability of divalent cations within phagosomes and thus restricting bacterial growth (Cheryayil, 2011). For example, in Mycobacterium bovis-infected macrophages, Nramp1 and the iron exporter ferroportin 1 (Fpn1; Slc40a1) are recruited to Mycobacterium-containing phagosomes, leading to transportation of iron outside macrophages and suppression of M. bovis growth (Van Zandt et al., 2008). In addition, Nramp1 can promote host immune responses via regulation of iron homeostasis. Nramp1-mediated reduction of cellular iron pools results in suppression of interleukin-10 (IL-10), an anti-inflammatory cytokine, leading to high TNF-α production and enhanced killing activity of S. Typhimurium-infected macrophage via iNOS-mediated NO release (Fritsche et al., 2008). Despite numerous studies on bacterial survival in macrophages, the influence of Nramp1 on expression of bacterial virulence factors, either directly or through its pleiotropic effects on macrophage function, has not been addressed.

Burkholderia pseudomallei is a Gram-negative intracellular bacterium that causes melioidosis, a severe invasive disease of humans that is endemic in subtropical areas. It is an environmental saprophyte and can survive in water and soil for long periods. It is closely related to Burkholderia thailandensis, which has been found in a similar habitat to B. pseudomallei, but is rarely reported to cause disease in humans, though at high doses in a murine model it could cause pneumonia-derived sepsis (Glass et al., 2006). B. pseudomallei is able to survive and replicate in macrophages, and is able to invade non-phagocytic cells, spread from cell-to-cell via actin-based motility and induce cell fusion. B. thailandensis is less efficient than B. pseudomallei in its ability to adhere to and invade human lung epithelial cells, resulting in lower host cell damage (Kespichayawattana et al., 2004). Although a number of studies reported the roles of Nramp1 expression in macrophages against bacterial infection (Nairz et al., 2014; Wessling-Resnick, 2015), the influence of Nramp1 in response to infection by Burkholderia species has not been reported.

Here we examine whether different Nramp1-controlled macrophage defense mechanisms act differently on Burkholderia pseudomallei compared to B. thailandensis. We examine the survival of the two Burkholderia spp. in stably transfected macrophage cell lines expressing wild-type (Nramp1⁺) and mutant (Nramp1⁻) Nramp1. We provide evidence that Nramp1 mediates killing of B. thailandensis via the production of reactive oxygen species. In contrast, intracellular net replication of B. pseudomallei was higher than observed for B. thailandensis in Nramp1⁺ macrophages. Our data indicate that evasion of Nramp1- and NADPH oxidase-mediated killing of B. pseudomallei in macrophages is associated with escape from phagosomes. Interestingly, even though B. pseudomallei escaped phagosomes to a comparable extent in macrophages with or without functional Nramp1, transcription of virulence genes associated with Type III and Type VI secretion systems and actin-based motility was elevated in Nramp1⁺ cells.

MATERIALS AND METHODS

Bacterial Strains, Macrophages Cell Lines, and Growth Conditions

Non-pathogenic B. thailandensis E264 and pathogenic B. pseudomallei K96243 strains were used throughout the study. A B. pseudomallei mutant lacking the Bsa T3SS translocon component BipB was constructed by insertion of the pKNOCK-Tet suicide vector and has been described (Suparak et al., 2005). Salmonella enterica serovar Typhimurium 13311 (hereinafter referred to as S. Typhimurium) was obtained from the American Type Culture Collection (ATCC). Bacteria were routinely cultured on Luria–Bertani (LB; TM MEDIA, India) agar or in broth. The bipB mutant was maintained on agar plates or in broth containing 60 µg/mL tetracycline (Sigma-Aldrich, USA). Growth of Burkholderia strains under iron-limiting conditions was performed by culturing bacteria in M9 minimal salts supplemented with 0.1% (w/v) casamino acids and 0.4% glucose (M9CG) as previously described (Burtnick and Brett, 2013). All experiments involving viable B. pseudomallei were conducted under biosafety level 3 containment.

Stably transfected RAW 264.7 macrophage cell lines expressing wild-type (Nramp1⁺) and G169D mutant (Nramp1⁻) Nramp1 were constructed as previously described (White et al., 2004), and obtained from the laboratory of Prof. Jenefer M. Blackwell. In transfected macrophages expressing wild-type Nramp1 the protein localizes to membranes of late endosomes/lysosomes, whereas the G169D mutant protein is retained in the endoplasmic reticulum and degraded (White et al., 2004). RAW 264.7 macrophages carry a mutated oxidase-mediated killing in macrophages and that expression of virulence-associated genes by pathogenic B. pseudomallei is enhanced in macrophages expressing wild-type compared to non-functional Nramp1. B. thailandensis has been proposed as surrogate for B. pseudomallei in the study of melioidosis however our study highlights important differences in the interaction of these bacteria with macrophages.

Keywords: B. pseudomallei, B. thailandensis, iron, macrophages, Nramp1, Slc11a1, secretion system
endogenous copy of the Nramp1 gene (White et al., 2004). The cell lines were routinely maintained in RPMI 1640 medium (Gibco, USA) supplemented with 10% (v/v) heat-inactivated fetal bovine serum (FBS; HyClone, USA) at 37°C under an atmosphere of 5% CO₂, and periodically grown under 400 µg/mL geneticin (G418 sulfate; Sigma-Aldrich, USA) selection to ensure that macrophage clones were stably expressing the Nramp1 transgene.

**Macrophage Activation and Inhibitor Treatments**

Sixteen hours prior to infection experiments, macrophages were routinely activated using 50 U/mL of recombinant mouse IFN-γ (Sigma-Aldrich, USA). A dose-response experiment for IFN-γ treated cells was also performed for bacterial survival assays. Activated macrophages (50 U/mL IFN-γ) were incubated with the NADPH oxidase inhibitor diphenyleneiodonium chloride (DPI; Sigma-Aldrich, USA) at 50 µg/mL, or the specific iNOS inhibitor amino guanidine (AG; Sigma-Aldrich, USA) at 250 µg/mL, at 37°C, 5% CO₂ for 1 h prior to bacterial infection. At these inhibitor concentrations, cell viability was more than 95% as determined by MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; Sigma-Aldrich, USA) assay.

**Macrophage Infection Assay**

Net intracellular survival and replication of bacteria in macrophages was quantified by means of net survival where bacterial numbers decline over time or net replication where numbers increase over time to reflect the balance of intracellular killing or intracellular replication processes as previously described (Muangsombut et al., 2008). Briefly, IFN-γ stimulated (50 U/mL, or dose response as indicated) Nramp1⁺ or Nramp1⁻ macrophages were seeded at a density of 4 × 10⁵ cells into 24-well tissue culture plates and cultured for 16 h. Activated macrophages were incubated at a multiplicity of infection (MOI) of 1 and incubated at 37°C, 5% CO₂. After 1 h of infection, extracellular *Burkholderia* spp. or *S. Typhimurium* were killed by incubation with culture medium containing 250 µg/mL kanamycin (for *Burkholderia* spp.) or 100 µg/mL gentamicin (for *S. Typhimurium*) for 2 h at 37°C under 5% CO₂ atmosphere. At this point, in some macrophages infection experiments, cells were lysed to enumerate bacteria that had been taken up by the macrophages. For studies to quantify net replication over time after this point, the infected cells were maintained in culture medium containing 20 µg/mL kanamycin or 10 µg/mL gentamicin. At the indicated time points, the intracellular bacteria were released by adding 0.1% Triton X-100 (Sigma-Aldrich, USA). The number of intracellular bacteria recovered were quantified by serial dilution and plating on tryptic soy agar (TSA). Bacterial colonies were counted after 24–48 h of incubation at 37°C.

**Intracellular Oxidative Burst and Nitric Oxide Detection Assays**

Intracellular superoxide anion levels were determined by measuring deposition of formazan inside macrophages following addition of nitroblue tetrazolium (NBT) (Rook et al., 1985). Briefly, IFN-γ (50 U/mL) activated infected cells were treated with freshly prepared 2 mg/mL NBT (Sigma-Aldrich, USA) and incubated at 37°C for 1 h. As a positive control, macrophages were triggered with 0.5 µg/mL of phorbol myristate acetate (PMA; Sigma-Aldrich, USA) for 1 h. To terminate the reaction, the supernatant was removed and the cells were washed twice with methanol. The reduced NBT formazan deposits inside the cells were solubilized by addition of 120 µL of 2 M potassium hydroxide (KOH; Merck) and 140 µL of dimethyl sulfoxide (DMSO; Sigma-Aldrich, USA). Absorbance of the solution was determined at 630 nm. Background values were determined by incubation of macrophages in medium including NBT without PMA. Cell numbers were determined colorimetrically using 0.5% crystal violet in 50% ethyl alcohol and measuring absorbance at 590 nm. Oxidative burst-generated superoxide anion was represented as a ratio of formazan/crystal violet (A₄₅₀/A₅₉₀).

The generation of reactive nitrogen species (RNS) was measured as nitrite levels in cell culture supernatants, which represent the stable oxidation end-product of inducible nitric oxide synthase (iNOS)-generated nitric oxide (NO) (Breitbach et al., 2011). Briefly, IFN-γ (50 U/mL) stimulated macrophages were infected with bacteria as above. At 16 h post-infection (p.i.) nitrite levels were measured by addition of the Griess reagent (Molecular Probes, USA) and absorbance was read at 570 nm. The concentration of nitrite was determined from an established standard curve of sodium nitrite from 0 to 200 µM.

**Detection of Bacterial Virulence Gene Expression in Infected Macrophages**

Transcription of the gene encoding the *Burkholderia* T6SS-1 cluster encoded effector Hcp1 (BPSS1498 for *B. pseudomallei* and BTH II0868 for *B. thailandensis*), the gene encoding *Burkholderia* intracellular motility A (bimA₉₈ BPSS1492; bimA₂₃ BTH II0875), and the gene encoding the Bsa T3SS-3 effector protein BopE and BsaU structural component (bopE₄₁₇ BPSS1525; bsuU₈₉₈ BPSS1539) in macrophages activated with 50 U/mL IFN-γ was determined by reverse transcription (RT)-PCR analysis. In brief, intracellular *Burkholderia* were released from the infected macrophages by isotonic lysis using deionized water and differential centrifugation. The pelleted bacteria were collected and total bacterial RNA was extracted using PrestoTM Mini RNA kit (Geneaid, Taiwan) according to manufacturer’s instruction. The conversion of total RNA to cDNA was performed using SuperScript® III First-Strand Synthesis System (Invitrogen, USA). The cDNA was subjected to DNA amplification with specific primers (Table 1). PCR reactions were performed under the following conditions: 97°C, 5 min and 30 cycles of 97°C for 45 s, 55°C for 45 s, and 72°C for 1 min, followed by incubation at 72°C for 5 min. Each assay was performed in duplicate. Amplification of *B. pseudomallei* and *B. thailandensis* 23S rRNA were used as an internal normalization control. The image densities of amplicons
Quantification of the Intracellular Iron Content in Infected Macrophages

Total intracellular iron content of IFN-γ activated *Burkholderia*-infected macrophages was measured by Graphite Furnace Atomic Absorption Spectrometry (GFAAS), as previously described (Nairz et al., 2008), using AA-6800G (GFA-EX-7), atomic-absorption spectrometer with extended lifetime graphite tubes (Shimadzu corporation, Japan). *Burkholderia*-infected cells were washed three times in normal saline. The lysis of cells was carried out in 0.1% actual analytical nitric acid (Mallinckrodt Chemical, Ireland) and 0.2% Triton-X100 and ultrasonication for 40 s. A pyrolysis temperature of 1000°C and an atomization temperature of 2000°C were used. The concentration of total cellular protein was determined using Pierce™ bicinchoninic acid (BCA) protein assay kit (Thermo Scientific, USA) from an established standard curve of protein ranged from 0 to 2,000 µg/mL.

Confocal Analysis of Bacterial Co-localization with LAMP1

To assess the extent of *Burkholderia* escape from phagosomes of activated (50 U/mL IFN-γ) Nramp1+ or Nramp1− macrophages, we examined co-localization of bacteria with lysosome-associated membrane protein 1 (LAMP1) at 4 h p.i. The bacteria were stained red with mouse anti-*Burkholderia* monoclonal antibody (9D5; 1:20; from the laboratory of Dr. Narisara Chantratita) and Alexa Fluor 568-goat anti-mouse IgG (1:500; Invitrogen, USA). LAMP1 was stained green with rat monoclonal antibody (1D4B; 1:200; Abcam, USA) and Alexa Fluor 488 goat anti-rat IgG antibody (1:500; Molecular Probes, USA), and nuclei were stained blue with 4,6-diamidino-2-phenylindole (DAPI; 1:500). All of the staining procedures were performed for 1 h at 37°C. Cells were examined by a laser-scanning confocal microscope equipped with LSM5 Image Browser (LSM 510 META, Carl Zeiss, Germany). The percentage of intracellular *Burkholderia* associated with LAMP1 was determined as the number of bacteria co-localized with LAMP1/total number of bacteria within macrophages × 100. The association of *Burkholderia* with LAMP1 was considered when the red fluorescent bacteria co-localized with the green fluorescence of LAMP1-positive vacuoles, represented as an area of yellow staining.

Multinucleated Giant Cell (MNGC) Formation Assay

*Burkholderia*-infected macrophages were examined for MNGC formation according to a previously described method (Kespichayawattana et al., 2000). In brief, IFN-γ activated macrophages were infected as described above and at 4, 6, 8, and 10 h p.i., fixed and stained with Giemsa (Merck, Germany). By light microscopy at least 1,000 nuclei per slide were examined for MNGC formation. MNGC was defined as an infected cell with three or more nuclei per cell (Kespichayawattana et al., 2000). The percentage of MNGC formation was calculated as follows: (number of nuclei within multinucleated cells/total number of nuclei counted) × 100 (Kespichayawattana et al., 2000).

Cytokine Detection Assay

Supernatants harvested from bacteria-infected macrophages were collected at indicated time point and filtered through a 0.45-µm membrane. Pro-inflammatory cytokines, including tumor necrosis factor alpha (TNF-α), interleukin-6 (IL-6) and IL-1β, were determined in triplicate by Luminex Bio-Plex™ (Bio-Rad, USA) according to the instructions of the manufacturer. As a positive control for induction of pro-inflammatory cytokines, 20 µg/mL lipopolysaccharide (LPS; List Biological Labs, USA) from *Escherichia coli* was used.

Statistical Analysis

All assays were conducted at least in triplicate and unpaired t-tests of independent experiments were performed using Graph Pad Prism6 software. Results were considered significant at a two-tailed P-value of ≤ 0.05.

RESULTS

Nramp1 Controls Replication of Non-pathogenic *B. thailandensis*, but Not Pathogenic *B. pseudomallei*, within Stably Transfected RAW264.7 Macrophages

Since little was known about the effect of Nramp1 on *Burkholderia* spp. infection, we first compared the survival of *B. pseudomallei* with its closely related non-pathogenic bacteria *B. thailandensis* in activated (50 U/mL IFN-γ) RAW264.7 macrophages stably transfected with wild-type (Nramp1+) or mutant (Nramp1−) Nramp1 (White et al., 2004). Net
intracellular survival of the bacteria was determined at 4, 8, and 12 h post-infection (p.i.). We observed that *B. thailandensis*-infected Nramp1$^+$ cells exhibited a significantly reduced number of viable intracellular bacteria as compared with infected Nramp1$^-$ cells at 4, 8, and 12 h p.i. (**Figure 1A**), indicating that Nramp1$^+$ wild-type macrophages were more effective at killing *B. thailandensis* than Nramp1$^-$ mutant macrophages.

Measurement of bacterial uptake showed no significant difference between Nramp1$^+$ and Nramp1$^-$ cells (**Figure 1B**), suggesting that the observed difference in killing was not due to differences in bacterial uptake between the macrophage lines. Moreover, we found that killing efficiency of Nramp1$^+$ cells against *B. thailandensis* at 6 h p.i. was correlated with the concentration of IFN-γ used to stimulate the macrophages (from 10 to 200 U/mL) in a dose-dependent manner (**Figure 1C**), indicating that the influence of Nramp1 on controlling *B. thailandensis* survival is associated with IFN-γ stimulation. In contrast to these results with *B. thailandensis*, no significant difference in *B. pseudomallei* net intracellular replication was observed when comparing activated (50 U/mL IFN-γ) Nramp1$^-$ and Nramp1$^+$ macrophages (**Figure 1D**). The numbers of intracellular *B. pseudomallei* increased over the time intervals investigated in both types of macrophages suggesting that *B. pseudomallei* can escape the killing activity of functional Nramp1 in stably transfected RAW 264.7 macrophages.

**Nramp1 Restricts *B. thailandensis* Growth via an NADPH Oxidase-Mediated Mechanism**

Previous studies have shown that activation of Nramp1$^+$ macrophages is associated with enhanced activation of the oxidative burst through ROS-mediated NF-κB activity and Lcn2 transcription (Fritsche et al., 2012). To determine whether infection with *B. thailandensis* is accompanied by enhanced oxidative burst activity in Nramp1$^+$ macrophages, we measured the generation of superoxide anion. As shown in **Figure 2A**, *B. thailandensis*-infected or lipopolysaccharide (LPS)-stimulated Nramp1$^+$ cells produce higher levels of superoxide anion than Nramp1$^-$ cells at 4 h p.i., indicating a more effective induction of the oxidative burst by Nramp1$^+$ cells. Superoxide anion is the precursor of more potent microbicidal ROS regulated by the NADPH oxidase complex. To determine whether production of ROS is important for Nramp1-mediated killing of *B. thailandensis*, we performed further experiments in the presence of the NADPH oxidase blocking agent
Nramp1-Mediated Restriction of *B. thailandensis* Does Not Involve Nitric Oxide (NO)

Previous studies have also indicated differences in induction of iNOS and associated NO production in activated Nramp1+ and Nramp1− macrophages (Fritsche et al., 2008; Song et al., 2013). To determine the role of iNOS-induced RNS killing of *B. thailandensis* in Nramp1+ macrophages, we first measured the end product nitrite. We found that activated (50 U/mL IFN-γ) Nramp1+ cells infected with *B. thailandensis* or *S. Typhimurium* produced a significantly higher level of nitrite than activated Nramp1− cells over 16 h p.i. (Figure 2D). However, while NO production was reduced by specific blocking of iNOS enzyme activity with aminoguanidine (AG) (Figure 2D), this had no statistically significant effect on the ability of Nramp1+ cells to control *B. thailandensis* (Figure 2E). These results indicate that the ability of Nramp1+ macrophages to kill *B. thailandensis* does not require iNOS-generated NO.

*B. thailandensis* Induces Pro-inflammatory Cytokine Responses in Nramp1+ Macrophages

Since Nramp1 is well-known to have pleiotropic effects on release of pro-inflammatory cytokines, including IL1-β, TNF-α and IL-6, in response to bacterial uptake (Valdez et al., 2008), we also investigated the effects of Nramp1 on production of pro-inflammatory cytokines in response to *Burkholderia* infection. Activated (50 U/mL IFN-γ) *B. thailandensis*-infected Nramp1+ cells exhibited an increased level of TNF-α and IL-6 than...

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**FIGURE 2** | Effect of Nramp1 on ROS-mediated oxidative burst, RNS-mediated NO and killing of *B. thailandensis* within infected macrophages. (A) ROS-mediated oxidative burst in activated (50 U/mL IFN-γ) *B. thailandensis*-infected macrophages. Oxidative burst-generated superoxide anion production was determined by NBT reduction assay at 4 p.i. and represented as a ratio of formazan:crystal violet. Nramp1+ or Nramp1− activated (50 U/mL IFN-γ) macrophages were treated with/without NADPH oxidase inhibitor (50 µg/mL of DPI) for 1 h prior to *B. thailandensis* infection. (B,C) show net intracellular survival of *B. thailandensis* in Nramp1+ cells or Nramp1− cells, respectively, with/without inhibition (DPI) of the oxidative burst. (D) RNS-mediated NO of *B. thailandensis*-infected macrophages. The production of NO in infected macrophages was examined by Griess reagent assay. Nramp1+ or Nramp1− cells were treated with/without iNOS enzyme inhibitor (250 µg/mL of AG) for 1 h prior to infection. (E) Net intracellular survival of *B. thailandensis* in Nramp1+ cells with/without AG inhibition of the iNOS pathway. Results represent the mean ± SEM of three independent experiments. Asterisk indicates statistical significance (*P* ≤ 0.05, t-test) when comparing Nramp1+ cells with Nramp1− cells under the same conditions or when comparing infected cells treated with and without inhibitors.
B. pseudomallei, but Not B. thailandensis, Efficiently Escapes from the Phagosome of Nramp1+ Macrophages

One hypothesis to account for the difference in influence of Nramp1 on B. pseudomallei compared to B. thailandensis could relate to the ability of the bacteria to escape from the phagosome. We therefore used confocal microscopy to determine the level of escape from phagosomes by scoring colocalization of bacteria with LAMP1 (Figure 4A). At 4 h p.i. we found that c. 69% of intracellular B. thailandensis were associated with LAMP1+ vesicles in Nramp1+ cells (Figure 4B), this being a far greater proportion than the c. 9% of B. pseudomallei associated with LAMP1 in Nramp1+ cells at the equivalent time (Figure 4B). A B. pseudomallei mutant lacking the Bsa T3SS-3 translocon component BipB was included as control, as bipB mutant is known to exhibit delayed phagosome escape (Suparak et al., 2005). Of the intracellular bipB mutant bacteria analyzed, 95% were associated with LAMP1 (Figure 4B). These results suggested that B. thailandensis is less able to escape phagosome of Nramp1+ cells than B. pseudomallei. Significantly fewer B. thailandensis (27%; Figures 4A,B) were observed to be associated with LAMP1 in Nramp1− cells, possibly as the bacteria are not controlled in this compartment in macrophages lacking functional Nramp1 and are therefore able to escape to the cytosol.

Consistent with the hypothesis that retention of Burkholderia in phagosomes leads to more effective killing of the bacteria, we observed that the B. pseudomallei bipB mutant was recovered from Nramp1+ macrophages at comparable levels to B. thailandensis (Figure 4C), The bipB mutant also exhibited moderate attenuation in Nramp1− cells (Figure 4C), but the extent to which this reflects other cellular pathways for control of intracellular bacteria or other activities of T3SS-3 during intracellular life will be challenging to separate.

MNGC Formation in Nramp1+ and Nramp1− Macrophages

MNGC formation was measured in activated (50 U/mL IFN-γ) B. pseudomallei- and B. thailandensis-infected Nramp1+ and Nramp1− macrophages (Figure 5). At early stages of infection (4 and 6 h p.i.) we observed a significant difference in the percentage of nuclei in MNGC in Nramp1+ and Nramp1− B. pseudomallei infected macrophages, with MNGC formation being higher in mutant macrophages (Figure 5A). This difference had waned at 8 and 10 h p.i., indicating an increase in MNGC formation in Nramp1+ macrophages that may be associated with increasing...
escape of \textit{B. pseudomallei} from phagosomes over time. In contrast, the difference in MNGC formation between Nramp1\textsuperscript{+} and Nramp1\textsuperscript{−} macrophages was maintained throughout the 10 h infection period in \textit{B. thailandensis}-infected macrophages (Figure 5B).

Transcription of Virulence-Associated Genes of \textit{B. pseudomallei} Is Upregulated in Nramp1\textsuperscript{+} Macrophages

Our study demonstrates that \textit{B. pseudomallei} is better able to evade Nramp1-mediated control of bacterial replication than \textit{B. thailandensis}. Among the bacterial factors that have been shown to be important virulence determinants of \textit{B. pseudomallei} in murine models of melioidosis are Type VI secretion system cluster 1 (T6SS-1) (Burtnick et al., 2011), the Bsa Type III secretion system (T3SS-3) (Stevens et al., 2004) and BimA protein required for actin-based motility (Lazar Adler et al., 2015). Promoter-trap studies have indicated that Type VI secretion is induced during macrophage infection (Shalomi et al., 2007). Negative regulation of \textit{B. pseudomallei} T6SS-1 has been demonstrated in iron- and zinc-supplemented culture medium (Burtnick and Brett, 2013), but the role of these divalent cations in regulating expression upon interaction with host cells has not been studied.

To determine the influence of functional wild-type Nramp1 on the induction of T6SS-1, transcription of \textit{hcp1} was examined in activated (50 U/mL IFN-\gamma) \textit{Burkholderia}-infected Nramp1\textsuperscript{+} and Nramp1\textsuperscript{−} cells. As shown in Figure 6A, the level of mRNA of \textit{B. pseudomallei} \textit{hcp1} in Nramp1\textsuperscript{+} cells was higher than in Nramp1\textsuperscript{−} cells at 3 and 6 h p.i., despite our observation that \textit{B. pseudomallei} enters the cytosol at comparable levels in Nramp1\textsuperscript{+} and Nramp1\textsuperscript{−} cells as gauged by LAMP1 co-localization (Figures 4A,B). Similarly, transcription of \textit{B. pseudomallei} \textit{bimA}, located upstream of the T6SS-1 locus and co-regulated by the VirAG two-component sensory system, was induced in infected Nramp1\textsuperscript{+} cells when compared to Nramp1\textsuperscript{−} cells (Figure 6A). Bacteria cultured in rich LB broth or iron-limited minimal medium M9 supplemented with a carbon source and casamino acids (M9CG) were found to be
negative and positive for transcription of these genes, respectively (Figure 6A). Transcription of the B. pseudomallei bsaU gene encoding a structural component of T3SS-3 appeared elevated at 3 h p.i in Nramp1+ cells compared to macrophages lacking functional Nramp1, though equivalent levels of the transcript were detected at 6 h p.i. (Figure 6A). Levels of the transcript for the T3SS-3 effectors bopE were slightly greater in Nramp1+ cells. However, quantitative analysis of amplified bopE gene band density indicated no significantly different between expression in Nramp1+ and Nramp1− cells (Figure 6A). Transcription of the gene encoding 23S rRNA was observed under all condition tested, and showed no change over time. The absence of DNA contamination in RNA extracted samples was verified by PCR prior to cDNA synthesis (data not shown).

Unlike pathogenic B. pseudomallei, the expression of the B. thailandensis hcp1 and bimA genes at 3h p.i. was at the lower limit of detection, and showed no differences between Nramp1+ and Nramp1− at 6 h p.i. (Figure 6B) This suggests that B. thailandensis did not respond differently to the environment within Nramp1+ cells as the pathogenic strain did, despite the presence of 90% nucleotide sequences identity in the coding and promoter regions between B. pseudomallei T6SS-1 (Accession no. BPSS1496-1511) and B. thailandensis (Accession no. BTH_II0855-0870). It has been reported in B. pseudomallei that hcp1 and bimA genes were co-regulated (Burtnick et al., 2011). In Figure 6B, the expression levels of hcp1 and bimA genes are similar suggesting that both genes may be co-regulated. This may imply the conservation of hcp1 and bimA genes organization in B. thailandensis and B. pseudomallei, even though their precise location is unknown.

Measurement of Intracellular Iron Levels in Nramp1+ and Nramp1− Macrophages

Previous studies have shown that once B. pseudomallei escapes from the phagosome and enters the cytosol, highly reducing...
conditions induce expression of T6SS-1 via VirAG regulation activated by host glutathione (GSH) (Wong et al., 2015). We hypothesized that upregulation of Burkholderia T6SS-1 may be a consequence of differing levels of intracellular iron within Nramp1+ compared to Nramp1− cells. Total intracellular iron content of Nramp1+ and Nramp1− cells in response to Burkholderia infection was quantified by atomic absorption spectrometry and normalized to total protein concentration. Activated (50 U/mL IFN-γ) B. pseudomallei-infected Nramp1+ cells showed a significant reduction of the total cellular iron level when compared to Nramp1− cells (Figure 7). Since iron limitation is a known regulator of hcp1 (Burtnick and Brett, 2013), it is plausible that the reduced iron concentration detected in Nramp1+ macrophages may be a potential explanation for activation of the VirAG-coregulated hcp1 and bimA genes.

**DISCUSSION**

Previous studies have demonstrated that control of B. pseudomallei infection involves innate immune responses, a key component of which is activation of macrophages and neutrophils to kill bacteria (Lazar Adler et al., 2009). The Nramp1 is known to have many pleiotropic effects on macrophage activation, leading to efficient bacterial clearance for a number of intracellular bacterial and protozoan pathogens (Wessling-Resnick, 2015). Here we present data on the impact of some of these Nramp1-regulated pleiotropic effects on clearance of Burkholderia spp., in particular in relation to the differences in virulence of B. pseudomallei compared to B. thailandensis as summarized in Figure 8. Nramp1 controls growth of B. thailandensis within the phagoslysome via an NADPH oxidase-mediated mechanism (Figure 8A). In contrast, B. pseudomallei are able to evade Nramp1- and NAPDH oxidase-mediated killing in macrophages by escaping more rapidly from the phagolysosome (Figure 8B). Increased levels of transcription of virulence-associated genes occur in B. pseudomallei-infected Nramp1+ cells, possibly as a consequence of escaping to the cytosol to a greater extent that B. thailandensis and encountering a more iron-restricted environment.
First we show that functional wild-type Nramp1 can control the intracellular replication of *B. thailandensis*, but not *B. pseudomallei*, within IFN-γ stimulated macrophages. The action of Nramp1 in determining the response to macrophage activation by IFN-γ is well established (Atkinson et al., 1997; Barrera et al., 1997; Fritsche et al., 2003; Monack et al., 2004). Furthermore, the growth restriction of *B. thailandensis* in macrophages expressing wild-type Nramp1 was dependent upon the presence of functional NADPH oxidase but not inducible nitric oxide synthase. It is acknowledged that whilst DPI inhibits NADPH oxidase-mediated ROS production it may also affect other flavor enzymes, such as xanthine oxidase and nitric oxide synthase (NOS) (O’Donnell et al., 1993). We consider it unlikely that indirect effects on NOS explain the impact of DPI on control of *B. thailandensis* in Nramp1⁺ macrophages as the iNOS inhibitor aminoguanine had no significant effect. The absence of a role for NOS in control if intracellular *Leishmania donovani* resistance to *Burkholderia pseudomallei* is well established (Stevens et al., 2002). Instead, our studies have shown that *B. pseudomallei* T6SS-1 effector Hcp1 and the BimA protein mediating actin-based motility, is required for cell-to-cell spreading and MNGC formation, a hallmark of *B. pseudomallei* infection (Kespichayawattana et al., 2000; Stevens et al., 2005). Earlier studies have shown that *B. pseudomallei* T6SS-1 gene expression is negatively regulated under iron rich conditions during *in vitro* growth (Burtnick and Brett, 2013). Nramp1⁺ cells become iron limited due to a significant reduction in transferrin receptor 1 (TIR1)-mediated iron uptake and ferroportin 1 (Fpn1)-mediated iron export (Fritsche et al., 2007). In contrast, Nramp1⁻ cells remain iron sufficient. Our data confirm a reduction in intracellular iron in Nramp1⁺ compared to Nramp1⁻ macrophages, which may be associated with the enhanced induction of the T6SS-1-encoded *hcp1* gene in *B. pseudomallei*-infected Nramp1⁺ macrophages. Induction of *bimA* (which is co-regulated with T6SS-1 by VirAG) was also observed in Nramp1⁺ macrophages infected with *B. pseudomallei* at 6 h p.i., when escape into the cytosol is known to have occurred (Stevens et al., 2002). It is important to note that induction of *hcp1* and *bimA* occurred to a greater extent in Nramp1⁺ macrophages, even though the extent of co-localization of *B. pseudomallei* with LAMP1 was comparable in macrophages expressing wild-type and mutant Nramp1. This implies that in both cell types the bacteria escape to the cytosol at an equivalent level, and that the differences in gene expression may therefore reflect Nramp1-dependent differences in the environmental cues they encounter.

It is plausible that the reduced levels of intracellular iron detected in Nramp1⁺ cells are responsible for the observed induction of *hcp1* and *bimA* expression, however further studies will be required to determine the cytosolic concentration of iron in cells expressing wild-type or mutated Nramp1 and test whether the differences observed are sufficient to trigger a change in *hcp1* and *bimA* expression *in vitro*. Moreover, studies with *hcp1* or *bimA* mutants in Nramp1⁺ or Nramp1⁻ macrophages may help to elucidate whether the observed differences in MNGC formation occur through Type VI secretion- and/or actin-based motility-dependent pathways. However, it should be noted that at many time intervals
cell fusion occurred to a greater extent in macrophages lacking functional Nramp1 and at no time interval tested was MNNGC formation greater in Nramp1+ cells. Nevertheless, our study indicates for the first time that Nramp1- and NAPDH oxidase-mediated control of intracellular *Burkholderia* differs between non-pathogenic and pathogenic species, and indicates that Nramp1 can affect the expression of virulence-associated genes in a manner that appears independent of escape from phagosomes.

**AUTHOR CONTRIBUTIONS**

VM, JB, and SK designed the research. VM performed experiments and analyzed data. VM, PW, and VS performed infection experiments with *B. pseudomallei*. NC developed monoclonal antibody to *B. pseudomallei* and *B. thailandensis* for immunofluorescence detection. MS critically revised the manuscript. VM, JB, and SK wrote the manuscript. JB and SK conceptualized the study. All authors read and approved the final version of the manuscript.

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