Virulent *Francisella tularensis* Destabilize Host mRNA to Rapidly Suppress Inflammation

Timothy J. Bauler Jennifer C. Chase Tara D. Wehrly Catharine M. Bosio

Immunity to Pulmonary Pathogens Section, Laboratory of Intracellular Parasites, Rocky Mountain Laboratories,
NIAID, NIH, Hamilton, Mont., USA

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mRNA stability · Intracellular bacteria · Inflammation ·
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Abstract

Highly virulent bacterial pathogens have evolved rapid means to suppress host inflammatory responses by unknown mechanisms. Here, we use virulent *Francisella tularensis*, the cause of lethal tularemia in humans, as a model to elucidate these mechanisms. We show that following infection of murine macrophages *F. tularensis* rapidly and selectively destabilizes mRNA containing adenylate-uridylate-rich elements that encode for cytokines and chemokines important in controlling bacterial infection. Degradation of host mRNA encoding interleukin (IL)-1 β , IL-6 and CXCL1 did not require viable bacteria or de novo protein synthesis, but did require escape of intracellular organisms from endocytic vesicles into the host cytosol. The specific targeting of host mRNA encoding inflammatory cytokines and chemokines for decay by a bacterial pathogen has not been previously reported. Thus, our findings represent a novel strategy by which a highly virulent pathogen modulates host inflammatory responses critical to the evasion of innate immunity.

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Introduction

Rapid induction of inflammatory responses is required for the development of effective innate and adaptive immunity against invading pathogens. The host has a wide array of strategies to detect, respond and eliminate microorganisms. These include pattern recognition receptors, compartmentalization in toxic vacuoles and production of proinflammatory cytokines and chemokines to directly activate antimicrobial pathways and/or recruit host cells capable of aiding in eradication of the pathogen [1]. Virulent organisms must evade and modulate these host factors to ensure their successful infection, replication and dissemination in the host. Identification of the mechanisms by which highly virulent bacteria negatively regulate host responsiveness to infection is central to identification of novel therapeutics and vaccines.

Francisella tularensis ssp. *tularensis* (Ftt) is a Gram-negative facultative intracellular bacterium and the causative agent of tularemia. Isolates of Ftt are extremely virulent, causing acute lethal disease in humans and mice following inhalation with as few as 10 bacteria [2]. Ftt infects the host through a variety of routes, including inhalation, ingestion, through skin abrasions and by arthropod bites [3]. Following inoculation, Ftt primarily targets macrophages and dendritic cells in the lung and peripheral organs, e.g. the spleen, for intracellular replication [3]. After being engulfed by the host cell via phagocytosis, the bac-

terium quickly escapes the endocytic compartment gaining access to the cytosol, where it replicates to high numbers. The existence of reproducible in vitro and in vivo models that closely replicate features of human disease make Ftt an ideal model organism for the study of immunity to highly pathogenic organisms.

There are several subspecies and strains of *Francisella* that can be categorized by their relative virulence for humans (as reviewed [2]). As stated above Ftt causes an acute, lethal disease in humans. *F. tularensis* ssp. *holarctica* is also capable of causing significant disease in humans, but is not typically fatal. Over 60 years ago, a live vaccine strain (LVS) was generated from *F. tularensis* ssp. *holarctica* following serial passage in mice [4]. As a vaccine strain, LVS is largely attenuated in humans. Lastly, *F. novicida* is an avirulent species of *Francisella*. The avirulent nature of *F. novicida* has made this bacterium useful for incorporation into comparative studies performed with Ftt.

An important aspect of virulence embodied by Ftt, like other highly pathogenic organisms, is its ability to evade detection by the host and suppress induction of inflammatory responses in vitro and in vivo [5–7]. Here we uncovered a unique mechanism by which Ftt downregulates proinflammatory responses. Performing comparative and coinfection studies with avirulent *F. novicida*, we show that Ftt impacted the host inflammatory responses within the first 2 h of infection without the requirement for replication or de novo protein synthesis. Our data identified that the rapid impairment of inflammatory responses by Ftt was due to its ability to induce selective decay of host mRNA encoding cytokines and chemokines required for resolution of infection. This strategy may represent a common mechanism used by other intracellular pathogens to effectively target and dismantle the host inflammatory response.

Materials and Methods

Mice and Generation of Bone Marrow-Derived Macrophages

Specific pathogen-free C57BL/6 NHsd mice were bred at Rocky Mountain Laboratories. C57BL/6J wild-type controls (WT) and Toll-like receptor 2 (TLR2)-deficient mice (TLR2^{-/-}) were purchased from Jackson Laboratories (Bar Harbor, Me., USA). Tristetraprolin, i.e. TTP^{flox/flox} (WT), controls and TTP^{flox/flox} LysM-Cre (TTP^{-/-}) were provided by Dr. Peter Murray (St. Jude Children's Research Hospital, Memphis, Tenn., USA). All research involving animals was conducted in accordance with Animal Care and Use guidelines, and animal protocols were approved by the Animal Care and Use Committee at Rocky Mountain Laboratories. Bone marrow-derived macrophages (BMM) were generated as previously described, except that medium was supplemented with

20 ng/ml recombinant murine macrophage colony-stimulating factor and heat-inactivated fetal bovine serum was used in medium for all BMM cultures (Peprotech, Rocky Hill, N.J., USA) [8]. BMM were primed with either 500 ng/ml Pam₃CSK₄ (P3C) or 25 ng/ml monophosphoryl lipid A (both from Invivogen, San Diego, Calif., USA) as indicated in every experiment 16 h prior to infection.

Bacteria

Stock cultures of Ftt SchuS4 (Dr. Jeannine Peterson, CDC, Fort Collins, Colo., USA), SchuS4 Δ fevR FTT0383 (Dr. Jean Celli, Washington State University, Pullman, Wash., USA), LVS ATCC29684 (Dr. Karen Elkins, CBER/FDA, Rockville, Md., USA), *F. tularensis* ssp. *holarctica* strain FSC200 (Dr. Anders Sjöstedt, University of Umea, Umea, Sweden) and *F. novicida* U112 (Dr. Denise Monack, Stanford University, Stanford, Calif., USA) were generated and utilized as previously described [6]. As indicated, killed SchuS4 was generated following treatment with 1.5% paraformaldehyde (PFA) for 30 min and washed twice prior to inoculation. Alternatively, SchuS4 was killed by γ -irradiation (250 krad, ⁶⁰Co source) and washed once prior to inoculation. Successful killing of SchuS4 was verified by incubation of bacteria 72 h on modified Mueller-Hinton agar. Where specified, SchuS4 was pretreated with chloramphenicol (10 μ g/ml) 30 min prior to infection to prevent de novo bacterial protein synthesis, as previously described [9]. Chloramphenicol was retained in the media for the duration of the experiment. All experiments were performed under approved BSL-3 safety protocols at Rocky Mountain Laboratories.

Infection and Exposure of BMM to Francisella

Bacteria were resuspended at the indicated multiplicity of infection (MOI; or corresponding MOI for killed bacteria) in 250 μ l complete DMEM [cDMEM, DMEM with glutamine, HEPES and nonessential amino acid added (Life Technologies, Grand Island, N.Y., USA), plus 10% heat-inactivated fetal bovine serum (Thermo Fisher, Waltham, Mass., USA)], added to BMM on ice, and plates were centrifuged for 15 min at 850 g at 4°C. MOI were selected to ensure uptake of similar numbers of each strain of bacteria by BMM. Cells were then incubated at 37°C for 45 min, washed twice with DMEM, followed by incubation with gentamicin (50 μ g/ml for 45 min at 37°C) to eliminate extracellular bacteria. Cells were washed once with DMEM and cultured in cDMEM. Intracellular bacteria were enumerated as previously described [10]. Briefly, culture medium was removed, and cells were lysed with sterile water. Cell lysates were serially diluted and plated on modified Mueller-Hinton agar plates for enumeration of colonies. For coinfection experiments, SchuS4 and *F. novicida* were mixed and added to BMM in a total volume of 250 μ l and infection proceeded as above. In coinfecting cultures, approximately 95% of cells were infected with both SchuS4 and *F. novicida*, as determined by confocal microscopy. Where specified, BMM were pretreated with cytochalasin D (10 μ g/ml, A.G. Scientific, San Diego, Calif., USA) or bafilomycin A1 (100 nM, A.G. Scientific) 1 h prior to infection to inhibit phagocytosis and endosomal acidification, respectively. As indicated, the transcription inhibitor actinomycin D (2.5 μ g/ml, Sigma, St. Louis, Mo., USA) or the p38 inhibitor SB203580 (1 μ g/ml, Cell Signaling Technologies, Danvers, Mass., USA) was added at the initiation of infection. All drugs were retained in the media throughout the course of the infection. At the time points assessed, none of the chemicals significantly affected the viability of *Francisella*.

Table 1. Taqman gene expression assays and assay ID numbers used in this study

Gene	ID number
IL-1 β	Mm00434228_m1
IL-6	Mm00446190_m1
CXCL1	Mm04207460_m1
IL-18	Mm00434225_m1
CCL2	Mm00441242_m1
CCL3	Mm00441259_g1
CCL4	Mm00443111_m1
CCL5	Mm01302427_m1
CCL22	Mm00436439_m1
IL-1 α	Mm00439620_m1
IL-12b	Mm00434174_m1
TNF- α	Mm00443258_m1
GAPDH	Mm99999915_g1

Quantification of Cell Death and Cytokines

Cell death was measured using a Cytotox 96 nonradioactive cytotoxicity kit according to the manufacturer's instructions (Promega, Madison, Wisc., USA). Concentrations of interleukin (IL)-1 β , IL-6 and CXCL1 in culture supernatants were determined by ELISA (R&D Systems, Minneapolis, Minn., USA).

Western Blotting and Quantitative Real-Time Polymerase Chain Reaction

At the indicated time points medium was removed from BMM. Cells were lysed in 150 μ l NuPAGE LDS Sample Buffer (Life Technologies) by vigorous pipetting and scraping of the cells. Cell lysates were heated at 95°C and immediately placed on ice prior to loading onto 4–12% SDS-NuPAGE gradient gels (Life Technologies). Western blots were generated from SDS-NuPAGE gels as previously described [11]. Blots were probed with antibodies to phospho-T180/Y182 p38, p38, phospho-T334 MK2 (27B7), MK2 and β -actin (13E5; all from Cell Signaling Technologies). RNA was purified using RNeasy kits (Qiagen), cDNA generated using the Superscript VILO kit (Life Technologies) and quantitative real-time polymerase chain reaction (qRT-PCR) was run using primer/probe sets (table 1; all from Life Technologies) and an ABI 7900HT (Life Technologies). Input RNA was normalized to GAPDH, and n-fold change of the indicated genes as compared to untreated, uninfected controls was quantified as $\Delta\Delta C_T$. The mRNA half-life ($t_{1/2}$) was calculated as $-\ln 2/m$, where m is the slope of the line fit on a semilogarithmic plot of mRNA concentration as a function of time using least-squares regression as previously described [12].

Phagosomal Integrity Assay

To differentiate between vacuolar and cytosolic bacteria, phagosomal integrity assays were performed, as previously described [8]. Anti-*F. tularensis* lipopolysaccharide (US Biological, Salem, Mass., USA) antibodies were conjugated with either Alexa Fluor 546 or Alexa Fluor 488 using Zenon Alexa Fluor 546 mouse IgG2a or Zenon Alexa Fluor 488 mouse IgG2a labeling kits according to the manufacturer's instructions (Invitrogen). Briefly, BMM were treated with 50 μ g/ml digitonin for 1 min to selectively per-

meabilize the plasma membrane, and then stained with Alexa Fluor 546-conjugated anti-*F. tularensis* lipopolysaccharide to label cytoplasmic bacteria. Next, BMM were fixed with 3% PFA for 20 min at 37°C and further permeabilized in 0.1% saponin/10% horse serum in phosphate-buffered saline for 30 min. All intracellular bacteria were labeled using Alexa Fluor 488-conjugated anti-*F. tularensis* lipopolysaccharide (US Biological). Cytoplasmic bacteria exhibited 546/488 dual fluorescence whereas vacuole-bound bacteria exhibited only 488 single fluorescence allowing differential enumeration of organisms. Detection and quantitation of cytoplasmic versus vacuole-bound bacteria was accomplished by visualizing infected cells using an Axio Imager epifluorescence microscope (Zeiss).

Statistical Analysis

Statistical differences between 2 groups were determined using a 2-tailed Student's t test. For comparison between 3 or more groups, analysis was done by 1-way ANOVA followed by Tukey's multiple comparisons test. Significance was determined at $p < 0.05$.

Results

SchuS4 Inhibits Production of IL-1 β , IL-6 and CXCL1 in Previously Activated Cells by Reducing mRNA Transcripts

Ftt SchuS4 both evades provoking an inflammatory response upon infection and suppresses the ability of the host cell to mount an inflammatory response after infection [5, 11]. However, the majority of this work was conducted in resting cells. Given the dramatic ability of SchuS4 to dampen inflammation, we postulated that SchuS4 may also downmodulate inflammatory programs in cells activated prior to infection. We chose to examine IL-1 β , IL-6, CXCL1 and IL-18 as representative cytokines/chemokines produced by macrophages that are important for host defense against bacterial infections [13]. Macrophages were primed with P3C prior to infection resulting in increased phosphorylation of mitogen-activated protein kinase p38, increased expression of genes encoding IL-1 β , IL-6 and CXCL1, and secretion of IL-6 and CXCL1 (online suppl. fig. 1; for all online suppl. material, see www.karger.com/doi/10.1159/000363243). SchuS4 infection of P3C-primed BMM did not induce secretion of IL-1 β , IL-6, CXCL1 or IL-18 that was significantly greater than primed, uninfected cells, despite replication of the bacterium (fig. 1a, b). Rather, SchuS4 infection significantly decreased the amount of IL-6 and CXCL1 present in culture supernatants compared to primed, uninfected controls (fig. 1b). This demonstrated that SchuS4 evades provoking an inflammatory response and readily inhibits an established inflammatory program in previously activated cells.

Fig. 1. SchuS4 infection reduces IL-1 β , IL-6 and CXCL1 RNA transcripts to inhibit cytokine/chemokine production in previously activated BMM. BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI 50, *F. novicida* MOI 10 or coinfecting with both SchuS4 MOI 50 and *F. novicida* MOI 10. Cells were infected with the indicated MOI to ensure equivalent uptake of *F. novicida* and SchuS4. Mock-infected cells served as negative controls. **a** Intracellular bacteria were enumerated at the indicated times after infection. **b** Supernatants were harvested 12 h after infection, and the quantity of IL-1 β , IL-6, CXCL1 and IL-18 was measured by ELISA. * $p < 0.05$; n.s. = not significant when comparing the indicated groups. **c** Supernatants were harvested 12 h after infection, and lactate dehydrogenase levels were quantified to calculate the percentage of cell lysis. n.s. = Not significant compared to the indicated samples. **d** RNA was harvested at the indicated times after infection. qRT-PCR was performed to examine the quantity of the indicated transcripts, normalized to GAPDH. * $p < 0.05$, SchuS4 and *F. novicida*/SchuS4-coinfected cells compared to mock and *F. novicida*-infected cells. Each data point depicts the mean of triplicate samples. Error bars represent SEM. Data are representative of 3 independent experiments.

We next sought to test the ability of SchuS4 to modulate cytokine responses in host cells activated with a more complex stimulus rather than only a single defined ligand. We utilized avirulent *F. novicida* to increase the production of inflammatory cytokines among infected cells. As expected, *F. novicida* infection of primed cells resulted in secretion of significantly greater concentrations of IL-1 β , IL-6, CXCL1 and IL-18 compared to cultures of uninfected and SchuS4-infected cells (fig. 1a, b). BMM coinfecting with SchuS4 and *F. novicida* secreted similar amounts of IL-18 compared to cells infected with *F. novicida* alone (fig. 1b). However, BMM coinfecting with SchuS4 and *F. novicida* did secrete significantly less IL-1 β , IL-6 and CXCL1 than cells infected with *F. novicida* alone (fig. 1b). The dampened release of IL-1 β , IL-6 and CXCL1 observed in SchuS4-*F. novicida*-coinfecting cells could not be attributed to reduced viability of macrophages since cell death in SchuS4-*F. novicida*-coinfecting cultures was similar to that of cells infected with *F. novicida* only (fig. 1c). We next determined if the reduced secretion of IL-1 β , IL-6 and CXCL1 was correlated with a reduction in mRNA transcripts encoding these proteins. All cultures infected with SchuS4 had significantly less IL-1 β , IL-6 and CXCL1 mRNA transcripts compared to uninfected cells or macrophages infected with *F. novicida* alone (fig. 1d). In contrast, and in agreement with similar concentrations of IL-18 detected in culture supernatants of *F. novicida* and SchuS4-*F. novicida*-coinfecting cells, the amount of IL-18 mRNA was unaffected by SchuS4 infection at each time point tested (fig. 1d). Thus, inhibition of IL-1 β , IL-6 and CXCL1 among SchuS4-infected cells was correlated with decreased numbers of mRNA transcripts encoding these proteins. These data also indicated that the impact of SchuS4 infection was selective for specific genes rather than a global reduction of mRNA transcripts among infected cultures. We next tested the ability of other closely related subspecies and strains of *Francisella* to reduce levels of mRNA among BMM. Similar to SchuS4-infected cells, virulent *F. tularensis* ssp. *holarctica* strain FSC200 reduced the number of detectable mRNA transcripts and did not induce secretion of IL-1 β , IL-6 or CXCL1 above that detected in uninfected cells (fig. 2a, b). In contrast, similar to cells infected with *F. novicida*, BMM infected with attenuated LVS increased mRNA transcripts compared to uninfected and SchuS4- or FSC200-infected cells (fig. 2a). LVS and *F. novicida* also triggered secretion of significantly more IL-1 β , IL-6 and CXCL1 compared to uninfected and SchuS4- or FSC200-infected cells (fig. 2b). Therefore, reduction of mRNA transcripts encoding spe-

cific proinflammatory cytokines and chemokines is a feature of infection with virulent, but not avirulent or attenuated, strains of *Francisella*.

Reduction of IL-1 β , IL-6 and CXCL1 mRNA by SchuS4 Requires Phagosomal Escape, but Not Bacterial Protein Synthesis or Viability

We next identified the cellular location from which SchuS4 affected numbers of mRNA transcripts. Upon *Francisella* engagement of receptors at the plasma membrane, the bacterium is phagocytosed into an endosome and then approximately 1 h after infection escapes into the cytoplasm, where it replicates exponentially [14, 15]. SchuS4 could be initiating the observed decrease in IL-1 β , IL-6 and CXCL1 mRNA upon contact with the cell surface, while transiting the endosome, or after escape into the cytosol. Thus, using a combination of chemical inhibitors, BMM with specific mutations in surface receptors, mutagenized SchuS4 and two different techniques to kill the bacterium, we empirically determined the requirements for SchuS4 to inhibit production of IL-1 β , IL-6 and CXCL1.

To determine if SchuS4 interaction with the cell membrane was sufficient to reduce mRNA transcripts, the actin-depolymerizing agent cytochalasin D was used to inhibit bacterial phagocytosis. As expected, SchuS4 infection of cytochalasin D-treated BMM decreased bacterial uptake by >97% compared to untreated cells (data not shown). Cytochalasin D-treated cells infected with SchuS4 did not reduce IL-1 β , IL-6 or CXCL1 transcripts (fig. 3a). Therefore, phagocytosis of the bacterium was necessary to decrease these mRNA.

Previous studies have shown that interaction with the surface receptor TLR2 was required for immunosuppressive mechanisms of SchuS4 [16, 17]. To determine if ligation of TLR2 was required for modulation of secretion of IL-1 β , IL-6 and CXCL1 by SchuS4, we compared the ability of SchuS4 to affect production of these cytokines among WT and TLR2^{-/-} BMM. The absence of TLR2 had no effect on the ability of SchuS4 to either evade induction or dampen secretion of IL-1 β , IL-6 and CXCL1 among infected BMM (fig. 3b).

We next determined if reduction of IL-1 β , IL-6 and CXCL1 transcripts required escape of SchuS4 from the endosome via 2 methods. First, cells were treated with bafilomycin A1, which inhibits SchuS4 escape from the endosome (fig. 3c) [8]. SchuS4 infection of bafilomycin-treated cells did not decrease the number of IL-1 β , IL-6 or CXCL1 mRNA transcripts beyond that observed in treated, uninfected cells (fig. 3a). However, bafilomycin

Fig. 2. Reduction of IL-1 β , IL-6 and CXCL1 mRNA correlates with virulence of *Francisella*. BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI 50, FSC200 MOI 50, LVS MOI 25 or *F. novicida* MOI 10. Cells were infected with the indicated MOI to ensure equivalent uptake. **a** RNA was harvested 5 h after infection. qRT-PCR was performed to examine the quantity of the indicated transcripts, normalized to GAPDH. **b** Supernatants were

harvested 12 h after infection, and the indicated cytokines were quantified by ELISA. * $p < 0.05$: significantly reduced compared to the mock-infected cells; ** $p < 0.05$: significantly greater than uninfected or SchuS4- or FSC200-infected cells; n.s. = not significant compared to the indicated samples. Each data point depicts the mean of triplicate samples. Error bars represent SEM. Data are representative of 2 independent experiments.

treatment alone resulted in reduction of mRNA transcripts independent of infection (fig. 3a), suggesting that bafilomycin had an off target effect on the ability of host to increase transcription of host mRNA. Thus, we utilized another method to confirm our finding that phagosomal escape of SchuS4 was required for the reduction of host mRNA. Next, we used a targeted deletion mutant of SchuS4 which is defective for phagosomal escape, SchuS4 Δ fevR (fig. 3c) [18]. SchuS4 Δ fevR does not exhibit defects in growth in axenic medium and is phagocytosed by primary cells with similar efficiency as WT SchuS4. However, these mutants fail to escape the phagolysosome following engulfment by macrophages [18]. Infection of BMM with SchuS4 Δ fevR did not result in the reduction of IL-1 β , IL-6 or CXCL1 transcripts (fig. 3a). Although it was possible that fevR is important for the expression of bacterial proteins that may contribute to reduction of host mRNA transcripts, taken together with data from cells treated with bafilomycin, these results suggest that

reduction of IL-1 β , IL-6 and CXCL1 mRNA was dependent on SchuS4 escape into the cytosol.

We then assessed whether de novo bacterial protein synthesis was required to decrease IL-1 β , IL-6 and CXCL1 mRNA transcripts following SchuS4 infection. Chloramphenicol was used to inhibit SchuS4 protein synthesis. Chloramphenicol-treated SchuS4 escaped the endosome with similar kinetics as untreated bacteria but were unable to replicate, demonstrating the effectiveness of the antibiotic treatment (fig. 3d and data not shown). There were no significant differences in the reduction of transcripts encoding IL-1 β , IL-6 and CXCL1 among cells infected with chloramphenicol-treated SchuS4 versus untreated SchuS4 (fig. 3e). Thus, de novo protein synthesis was not required for SchuS4 to decrease these mRNA.

To determine if SchuS4 viability was required to reduce IL-1 β , IL-6 and CXCL1 transcripts, we compared mRNA levels among cells infected with live SchuS4 versus cells exposed to SchuS4 killed with either PFA or ir-

Fig. 3. Phagosomal escape, but not bacterial protein synthesis or viability, is required for SchuS4-mediated reduction of IL-1 β , IL-6 and CXCL1 mRNA. **a–c** BMM were primed with P3C (500 ng/ml) and treated with cytochalasin D (CytoD) to inhibit bacterial phagocytosis or bafilomycin (Bafilo) to inhibit bacterial escape from endosomes prior to infection, where indicated. BMM were infected with SchuS4 MOI 50 or SchuS4 Δ fevR MOI 50. Mock-infected cells served as negative controls. **a** RNA was harvested 5 h after infection. qRT-PCR was performed to examine the quantity of the indicated transcripts, normalized to GAPDH. * $p < 0.05$ compared to the indicated samples; n.s. = not significant compared to the indicated samples. **b** WT or TLR2^{-/-} BMM were primed with monophosphoryl lipid A (25 ng/ml) and infected with SchuS4 MOI 50, *F. novicida* MOI 10 or coinfecting with both SchuS4 MOI 50 and *F. novicida* MOI 10. Secretion of IL-1 β , IL-6 and CXCL1 was as-

essed by ELISA. * $p < 0.05$: *F. novicida*-infected compared to *F. novicida*/SchuS4-coinfected cells. **c** Cytoplasmic bacteria were quantified 5 h after infection using a phagosomal-integrity assay. * $p < 0.05$ compared to WT SchuS4 infection of untreated cells. **d**, **e** BMM were primed with P3C (500 ng/ml) and infected with MOI 50 of SchuS4, SchuS4 pretreated with chloramphenicol (Chloramp) to inhibit bacterial protein synthesis, or inoculated with SchuS4 killed by treatment with PFA or irradiation (Irradiate). **d** Cytoplasmic bacteria were quantified 5 h after infection using a phagosomal-integrity assay. * $p < 0.05$ compared to untreated SchuS4. **e** RNA was harvested 5 h after infection. qRT-PCR was performed to examine the quantity of the indicated transcripts, normalized to GAPDH. * $p < 0.05$ compared to uninfected cells. Each data point depicts the mean of triplicate samples. Error bars represent SEM. Data are representative of 3 independent experiments.

radiation. Unlike live SchuS4, PFA-killed SchuS4 were not able to significantly decrease mRNA transcripts of IL-1 β , IL-6 or CXCL1 compared to uninfected controls (fig. 3e). However, PFA-killed SchuS4 were retained in the endosome (fig. 3d). As described above, escape from the phagosome appeared to be essential for the reduction of mRNA transcripts by SchuS4 (fig. 3a). Thus, it was not clear if the effect on mRNA transcripts observed among cells treated with PFA-killed SchuS4 was due to the absence of viable SchuS4, or their retention in the endosome. To resolve this issue, we utilized irradiated SchuS4. Despite their inability to replicate, irradiated SchuS4 escape the endosome, which allowed us to assess the requirement for bacterial viability independently of escape into the cytosol (fig. 3d). Irradiated SchuS4 reduced IL-1 β , IL-6 and CXCL1 mRNA equivalently to live SchuS4 (fig. 3e). Thus, the ability of SchuS4 to decrease IL-1 β , IL-6 and CXCL1 transcripts was not dependent on the viability of the organism.

SchuS4 Infection Decreases IL-1 β , IL-6 and CXCL1 mRNA Concomitantly with p38 Dephosphorylation

The mitogen-activated protein kinases, and specifically p38, play a central role in the production of many pro-inflammatory cytokines/chemokines following bacterial infection, including IL-1 β , IL-6 and CXCL1. Therefore, we determined if SchuS4 infection impacted the activation of p38. We observed a notable dephosphorylation of p38 as early as 1 h after infection in SchuS4 and SchuS4-*F. novicida*-coinfected cells, compared to both uninfected and *F. novicida*-infected cells (fig. 4a). We next confirmed that dephosphorylation of p38 was a consistent phenotype among BMM infected or exposed to preparations of SchuS4 that were associated with reduction of mRNA. Consistently with their ability to reduce IL-1 β , IL-6 and CXCL1 mRNA, infection or exposure of BMM with SchuS4 treated with chloramphenicol or inactivated by irradiation both resulted in dephosphorylation of p38 in BMM (fig. 4b). Further, neither SchuS4 Δ fevR nor PFA-inactivated SchuS4 reduced phosphorylation of p38 compared to mock controls (fig. 4b). The lack of impact on phosphorylation of p38 by these bacteria is consistent with their inability to reduce host mRNA. Thus, dephosphorylation of p38 in BMM correlates with conditions in which SchuS4 was able to reduce host mRNA transcripts. To determine whether inactivation of the p38 pathway was sufficient to reduce IL-1 β , IL-6 and CXCL1 transcript levels and secretion in the absence of infection, we inhibited p38 activity with SB203580. BMM treated with SB203580 exhibited decreased levels of mRNA transcripts

and secretion of these proteins compared to uninfected cells (fig. 4c, d). Similarly, *F. novicida*-infected, SB203580-treated BMM had significantly fewer IL-1 β , IL-6 and CXCL1 mRNA transcripts and reduced protein detected in the culture supernatant compared to mock-treated *F. novicida*-infected controls (fig. 4c, d). Together these data demonstrate that SchuS4 inactivates p38 upon infection, and that decreased activation of p38 is sufficient to reduce IL-1 β , IL-6 and CXCL1 mRNA and secretion.

SchuS4 Infection Destabilizes mRNA Containing Adenylate-Uridylate-Rich Elements

Both transcription and mRNA stability are known to be regulated by p38 [19, 20]. Thus, the reduction in mRNA encoding IL-1 β , IL-6 and CXCL1 following SchuS4 infection could result either from a decreased rate of new transcription or an increased rate of mRNA decay. To differentiate between these two possibilities, actinomycin D was used to inhibit transcription. Actinomycin D is routinely used to determine if different stimuli are affecting mRNA levels during transcription or as a post-transcriptional event [21]. If the ability of SchuS4 to reduce mRNA transcripts in the presence of actinomycin D were lost, it would suggest that SchuS4 was acting to reduce mRNA by inhibiting new transcription. Conversely, if treatment with actinomycin D had no effect on the ability of SchuS4 to reduce the number of mRNA transcripts this would demonstrate that SchuS4 was reducing copies of mRNA via acceleration of mRNA decay. SchuS4 infection of actinomycin D-treated BMM resulted in a similar magnitude of reduction of IL-1 β , IL-6 and CXCL1 transcripts compared to untreated, SchuS4-infected cells (fig. 5a). Accordingly, the half-life of these transcripts in SchuS4-infected, actinomycin D-treated cells was significantly decreased compared to uninfected actinomycin D cells (fig. 5a). These data support that SchuS4-mediated reduction of IL-1 β , IL-6 and CXCL1 mRNA occurs via decreased mRNA stability. To further confirm that SchuS4 is inducing mRNA destabilization, we examined the activation of MK2, which is directly phosphorylated by p38 and phosphorylates mRNA binding proteins to indirectly regulate mRNA stability [22]. Similar to the kinetics of p38 inactivation following SchuS4 infection, dephosphorylation of MK2 was observed as early as 1 h after infection (fig. 5b). Thus, SchuS4-mediated inactivation of the p38-MK2 pathway upon infection results in the destabilization of IL-1 β , IL-6, and CXCL1 mRNA.

The stability of mRNA that encode tightly regulated proteins, such as many cytokines and chemokines, is controlled by the interaction of a variety of mRNA binding

Fig. 4. Dephosphorylation of p38 by SchuS4 infection. BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI 50, chloramphenicol (Chloramp)-treated SchuS4 MOI 50, SchuS4 ΔfevR MOI 50, *F. novicida* MOI 10, or coinfecting with both SchuS4 MOI 50 and *F. novicida* MOI 10. As indicated, BMM were exposed to SchuS4 inactivated with either PFA or irradiation. Where specified, cells were treated with the p38 inhibitor SB203580 at the onset of infection. Mock-infected cells served as negative controls. **a**, **b** Cell lysates were generated at the indicated times after infection

(**a**) or 5 h after infection or exposure to SchuS4 (**b**). Western blots were probed with antibodies to phosphorylated p38 (P-p38) and p38 and actin as loading controls. **c** RNA was harvested from cells 5 h after infection. qRT-PCR was performed to examine the indicated transcripts. **d** Supernatants were harvested 12 h after infection and the quantity of IL-1β, IL-6 and CXCL1 was measured by ELISA. * $p < 0.05$ comparing the indicated samples. Each data point depicts the mean of triplicate samples. Error bars represent SEM. Data are representative of 3 independent experiments.

proteins with mRNA sequence elements. The best-characterized of these are adenylate-uridylylate-rich elements (ARE), found in the 3'-untranslated region of many inherently unstable transcripts. ARE consist of one or more AUUUA pentamers which interact with a variety of host proteins which regulate the stability of host mRNA [23].

IL-1β, IL-6 and CXCL1 mRNA contain ARE, whereas IL-18 transcripts, which were not reduced after SchuS4 infection, do not. This suggested that mRNA containing ARE were targeted for degradation during SchuS4 infection [24]. To confirm that SchuS4 infection was inducing selective destabilization of ARE-containing transcripts,

5

(For legend see next page.)

we assessed a variety of mRNA transcripts encoding chemokines and cytokines produced by macrophages following the encounter with microorganisms. CCL2, CCL3, CCL4, IL-1 α , IL-12b and tumor necrosis factor α mRNA, which contain ARE, were all reduced 5 h after SchuS4 infection (fig. 5c). In contrast, CCL5 and CCL22 transcripts, which do not have ARE, remain unchanged upon SchuS4 infection (fig. 5c). Thus, SchuS4 infection resulted in selective destabilization of ARE-containing mRNA to limit production of key proinflammatory cytokines and chemokines.

There are many proteins that control the stabilization or decay of host mRNA via interaction with the 3'-ARE. The best-studied of these is TTP. Phosphorylation of TTP is dependent on p38. Phosphorylated TTP does not interact with host mRNA. Upon dephosphorylation, TTP interacts with AU-rich sequences in the 3'-region of host mRNA facilitating degradation of the sequence [25]. Given the observed dephosphorylation of p38 among SchuS4-infected cells and the selective targeting of ARE-containing mRNA for decay, we hypothesized that SchuS4 may be facilitating TTP activity to degrade host mRNA. However, we did not observe reconstitution of host mRNA stabilization among SchuS4-infected TTP-deficient macrophages (fig. 5d). This suggests that alternative, p38-dependent, mechanisms controlling host mRNA stabilization or degradation are targeted by SchuS4 in primary macrophages.

Discussion

It has been well established that SchuS4 both evades and suppresses inflammatory responses among resting resident host cells [5, 6, 11]. Here, we found that SchuS4

similarly inhibited proinflammatory responses among cells activated prior to challenge, as measured by secretion of key cytokines and chemokines. Further, we identified that a major mechanism by which SchuS4 rapidly inhibited production of these host products was to induce decay of ARE-containing mRNA transcripts.

We first established that ligation of surface receptors and/or phagocytosis of the bacterium was not sufficient for SchuS4 to execute suppression of host responses. Rather, we determined that mRNA decay relied on escape of SchuS4 into the cytosol. Our finding that interaction with surface receptors and uptake of the organisms alone were not adequate for SchuS4-mediated degradation of host mRNA was initially surprising. Previous studies showed that engagement of surface receptors, e.g. TLR2 [16] and CR3 [17], was required for some of the immunosuppressive mechanisms of SchuS4. The involvement of CR3 in mediating inhibition of inflammatory responses was dependent on opsonization of bacteria with serum prior to infection. We did not utilize opsonized bacteria in our study. Thus, it is unlikely that CR3 had a role in the degradation of mRNA. To confirm TLR2 was not required for modulation of IL-1 β , IL-6 or CXCL1, we verified that SchuS4 was still capable of inhibiting secretion of these cytokines following coinfection with avirulent *F. novicida* among TLR2^{-/-} cells (fig. 3c). Thus, to our knowledge, this data represents the first immunosuppressive mechanism induced by SchuS4 that dissociates *Francisella*-mediated inhibition of inflammatory responses from TLR signaling pathways.

We also determined that neither de novo bacterial protein synthesis nor viability was required for mRNA destabilization following infection, but that escape into the host cytosol was necessary to induce decay of host mRNA. *Francisella* escape from the endosome requires the *Fran-*

Fig. 5. Destabilization of ARE-containing mRNA following SchuS4 infection. **a** BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI 50 and treated with actinomycin D (ActD) at the onset of infection to inhibit transcription. RNA was harvested from cells at the indicated times after infection, and qRT-PCR was performed to quantify the indicated mRNA transcripts. The mRNA half-life ($t_{1/2}$) in uninfected and SchuS4-infected actinomycin D-treated samples is indicated. * $p < 0.05$ comparing the indicated sample to both uninfected and only actinomycin D-treated cells. **b, c** BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI 50, *F. novicida* MOI 10 or coinfecting with both SchuS4 MOI 50 and *F. novicida* MOI 10. **b** Cell lysates were generated at the indicated times after infection. Western blots were probed with antibodies to phosphorylated MK2 (P-MK2), and MK2 and actin as loading controls. Mock-

infected cells served as negative controls. **c** RNA was harvested from cells 5 h after infection, and qRT-PCR was performed to quantify the indicated mRNA transcripts. Mock-infected cells served as negative controls. TNF = Tumor necrosis factor; UTR = untranslated region. * $p < 0.05$ comparing the indicated sample to both uninfected and only *F. novicida*-infected cells. n.s. = Not significantly different. **d** WT controls and TTP^{-/-} BMM were primed with P3C (500 ng/ml) and infected with SchuS4 MOI, *F. novicida* MOI 10 or coinfecting with both SchuS4 MOI 50 and *F. novicida* MOI 10. RNA was harvested 5 h after infection. qRT-PCR was performed to examine the quantity of the indicated transcripts, normalized to GAPDH. * $p < 0.05$ comparing the indicated groups. Each data point depicts the mean of triplicate samples. Error bars represent SEM. Data are representative of 3 independent experiments.

cisella Pathogenicity Island, therefore it appears likely that the *Francisella* Pathogenicity Island and putative type VI secretion system of *Francisella* remain functional in the absence of new protein synthesis and in the majority of irradiated bacteria [26]. Together these data suggest that molecules required by the bacteria for both endosomal escape and suppression of proinflammatory responses via mRNA decay are preformed in SchuS4. Interestingly, bacteria killed using PFA did not escape the endosome and as a result also failed to accelerate host mRNA decay. Inactivation of bacteria with PFA or irradiation does not induce significant lysis of bacteria [27, 28]. However, PFA renders the bacteria completely inactive whereas γ -irradiation can prevent bacterial replication while leaving bacterial metabolism intact [29]. Thus, it is possible that although neither PFA-killed or irradiated bacteria were able to replicate, irradiated bacteria may have retained an active metabolism allowing for a functional type IV secretion system and escape from the endosome. Regardless of the mechanism of endosomal escape, our data show that the immediate availability of bacterial products capable of inhibiting inflammatory responses is consistent with the rapid ability of SchuS4 to modulate host responses. Identification of the bacterial component(s) required for mRNA decay is currently under investigation by our laboratory.

There are a number of means by which stability of host mRNA is controlled in the cell. The most commonly studied of these is the interaction of stabilizing/destabilizing proteins with ARE present in the 3'-untranslated region of host mRNA transcripts. There are several excellent reviews discussing the biology of ARE and their role in stabilization of host mRNA [23, 30, 31]. Briefly, ARE are pentamers of AUUUA. They are highly variable in their abundance and sequence and are divided into specific classes based on the number of ARE pentamers present in the 3'-untranslated region. ARE are not present in every mRNA and, thus, represent a mechanism by which the host can selectively control posttranscriptional production of potent cytokines and chemokines. Stabilization of host mRNA that contain ARE occurs through direct and/or indirect interaction of stabilizing or destabilizing proteins present in the host cell. Phosphorylation of these host proteins by kinases, e.g. by p38, dictates their activity. Modulation of ARE-interacting proteins has been noted as a mechanism by which viruses influence inflammatory responses in host cells [32, 33].

However, destabilization of cytokine mRNA during infection with intact bacteria has not been described. Rather, there is a precedent in the literature for increasing mRNA stability by bacteria and/or bacterial components

as a mechanism to sustain inflammatory responses. For example, avirulent *F. novicida* elicits a strong proinflammatory response following infection of host cells and it has been proposed that *F. novicida* inactivates TTP as a mechanism to enhance proinflammatory responses [34–36]. Specifically, an *F. novicida* Δ lpcC mutant provokes an even greater inflammatory response compared to WT *F. novicida*. This elevated inflammatory response corresponded with increased deactivation of TTP compared to cells infected with WT *F. novicida*. In contrast to infections mediated by *F. novicida*, we show that SchuS4 suppressed inflammatory responses by enhanced decay of host ARE-containing mRNA. Thus, it was possible that, as opposed to *F. novicida*, SchuS4 may further activate TTP to destabilize host mRNA. However, we did not observe a difference in the ability of SchuS4 to reduce ARE-containing mRNA among TTP-deficient BMM compared to littermate control BMM (fig. 5d). This suggested that SchuS4 infection may alter the activity of other ARE binding proteins known to interact with MK2, e.g. poly-A binding protein, HuR and BAF1, which regulate the stability of mRNA [37–39]. We are currently examining the ability of SchuS4 to modulate function of multiple ARE binding proteins following infection.

In this report, we describe a novel mechanism used by a highly virulent pathogen to rapidly and effectively dampen components of the host inflammatory response via destabilization of cellular ARE-containing mRNA. Importantly, this mechanism did not require synthesis of bacterial components following infection. Rather, effective induction of mRNA decay was initiated with only the proteins, lipids and carbohydrates present in the bacterium at the outset of infection. Our data represent an important advance in understanding the pathogenesis of disease following infection with Ftt. Destabilization of host cell mRNA may also represent a common mechanism of virulence employed by other similarly pathogenic bacteria. Continued characterization of virulence mechanisms restricted to highly virulent pathogens such as Ftt will likely be a fruitful approach to understanding the failure of the immune system to properly mount a protective response to these and other intracellular pathogens.

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