Yersinia pestis Can Bypass Protective Antibodies to LcrV and Activation with Gamma Interferon To Survive and Induce Apoptosis in Murine Macrophages

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Yersinia pestis, the agent of plague, uses a type III secretion injectisome to deliver Yop proteins into macrophages to counteract phagocytosis and induce apoptosis. Additionally, internalized Y. pestis can survive in the phagosomes of naive or gamma interferon (IFN-γ)-activated macrophages by blocking vacuole acidification. The Y. pestis LcrV protein is a target of protective antibodies. The binding of antibodies to LcrV at the injectisome tip results in neutralization of the apoptosis of Y. pestis-infected macrophages and is used as an in vitro correlate of protective immunity. The cytokines IFN-γ and tumor necrosis factor alpha can cooperate with anti-LcrV to promote protection against lethal Y. pestis infection in mice. It is not known if these phagocyte-activating cytokines cooperate with anti-LcrV to increase the killing of the pathogen and decrease apoptosis in macrophages. We investigated how anti-LcrV and IFN-γ impact bacterial survival and apoptosis in cultured murine macrophages infected with Y. pestis KIM5. Y. pestis KIM5 opsonized with polyclonal or monoclonal anti-LcrV was used to infect macrophages treated with or without IFN-γ. The phagocytosis and survival of KIM5 and the apoptosis of macrophages were measured at different time points postinfection. The results show that anti-LcrV reduced apoptosis at an early time point (5 h) but not at a later time point (24 h). Polyclonal anti-LcrV was unable to inhibit apoptosis at either time point in IFN-γ-activated macrophages. Additionally, anti-LcrV was ineffective at promoting the killing of KIM5 in naive or activated macrophages. We conclude that Y. pestis can bypass protective antibodies to LcrV and activation with IFN-γ to survive and induce apoptosis in murine macrophages.

There are three human pathogenic Yersinia species: Y. pestis, Y. pseudotuberculosis, and Y. enterocolitica (27). All three species contain an approximately 70-kb virulence plasmid (pCD1 in Y. pestis and pYV in Y. pseudotuberculosis and Y. enterocolitica) that encodes a type III secretion system (T3SS) and effector proteins termed Yersinia outer proteins or Yops. Y. pestis is the causative agent of pneumonic and bubonic plague and the latter two cause gastroenteritis. Y. pestis is thought to be closely related to Y. pseudotuberculosis (1). In addition to carrying pCD1, Y. pestis harbors two additional plasmids, pMT1 and pPCP1, that give it increased virulence compared to Y. pseudotuberculosis (27). Historically, Y. pestis has had a major impact on society, killing large numbers of people worldwide. Today, with the development of antibiotics and improved sanitary conditions, bubonic and pneumonic plague are no longer major public health concerns. However, there are still rodent populations infected with plague, and small numbers of humans within the population are infected annually (27). It is important to further study Y. pestis to create a safe and effective vaccine, both because there is still a natural reservoir and because there is the potential danger that pneumonic plague may be used for acts of bioterrorism.

The pCD1 plasmid encodes a T3SS composed of the secretion apparatus, chaperones, Yops (9), and the translocator proteins (YopB, YopD, and LcrV). Six effector Yops have been identified: YopH, YopO/YpkA, YopP/YopJ, YopE, YopM, and YopT. YopJ (YopP in Y. enterocolitica) inhibits mitogen-activated protein kinase signaling cascades as well as the activation of NF-κB, a transcription factor important for the expression of proinflammatory cytokines such as tumor necrosis factor alpha (TNF-α), by preventing the phosphorylation and subsequent degradation of the NF-κB inhibitor (IκB) (25). As a result, YopJ inhibits the expression of TNF-α, diminishing the inflammatory response. YopJ has also been shown to induce apoptosis, due to the loss of mitogen-activated protein kinase and NF-κB activity (25).

The phagocytosis of an opsonized pathogen by a macrophage is mediated by surface receptors, such as the Fc receptor. During the process of phagocytosis several signaling cascades are activated. Rho GTPase family members (Rac-1, RhoA, and Cdc42) that are localized in the plasma membrane become activated through the binding of GTP. These GTPases regulate the polymerization of actin to drive phagocytosis (2). Several of the Yops have antiphagocytic effects, through the direct inhibition of Rho GTPases, including YopO (also known as Yersinia protein kinase A), YopT, and YopE (25, 30). YopH has been shown to inhibit phagocytosis and the expression of monocyte chemoattractant protein 1, a chemokine involved in macrophage recruitment, and diminish the Fc-mediated oxidative burst in neutrophils and macrophages (6, 25).

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The expression of the T3SS and the regulation of Yop translocation are dependent on temperature, calcium levels, and host cell contact. At 28°C, the expression of the T3SS is down-regulated. At 37°C, the T3SS is maximally induced (9), and a needle-like surface structure, the Ysc injectisome, is formed. Upon contact with a host cell, the T3SS is systematically activated. The translocators YopB and YopD are believed to form a channel in the host cell membrane, allowing the delivery of the effector Yops. The effector Yops are translocated into the host cell cytoplasm, where they disrupt host cell signaling (9).

In addition to YopB and YopD, the LcrV protein is necessary to deliver the effector Yops into the host cell (28). The mechanism by which LcrV mediates translocation is not fully understood, but it appears to be important for the correct assembly of the translocation channel (23). LcrV has been shown to localize to the tip of the injectisome (23). LcrV, also known as V antigen, has many other important roles. It has a regulatory role in Yop secretion within the bacterium (27). LcrV is also a soluble protein and is an important protective antigen (24, 42).

Y. pestis is efficiently phagocytosed and survives within the phagosomes of naïve murine macrophages when the bacteria are grown at 28°C prior to infection (13, 14, 34, 41). Y. pestis can block phagosome acidification, which may be important for survival in macrophages (34). The growth of Y. pestis at 37°C prior to infection promotes Yop delivery during phagocytosis, and as a result, the efficiency of bacterial uptake by macrophages is reduced. However, ~20 to 35% of 37°C-grown Y. pestis bacteria that associate with macrophages are internalized (10, 43). Yop-expressing Y. pestis that are internalized by naïve macrophages are able to survive intracellularly (21). In addition, macrophages infected with 37°C-grown Y. pestis die of YopJ-induced apoptosis (12, 21, 43). Thus, Yop-expressing Y. pestis can counteract the antibacterial functions of naïve macrophages by intracellular survival and the induction of apoptosis if they are unable to avoid phagocytosis. Łukaszewski et al. showed that naïve mice infected with Y. pestis could harbor Y. pestis within CD11b+ spleen macrophages for several days postinfection (p.i.) and that a significant percentage of these phagocytes died of apoptosis during this time period (22).

Mice can be protected against lethal Y. pestis infection by passive immunization with anti-LcrV antibodies (15–17, 19, 38, 39, 42, 44). Opsonization with anti-LcrV antibodies increases the phagocytosis of Y. pestis by macrophages (10, 29, 43). The increased phagocytosis of Y. pestis mediated by anti-LcrV antibody opsonization is associated with reduced Yop translocation (10, 29) and reduced apoptosis (10, 29, 43). The ability of anti-LcrV antibodies to inhibit apoptosis in macrophages infected with Y. pestis is commonly used as a measure of neutralizing activity (3, 43–45).

In addition to antibodies, the cytokines gamma interferon (IFN-γ) and TNF-α are important for protective immune responses against Y. pestis infection (11, 19, 24, 26, 38, 39). Y. pestis can survive in macrophages activated with IFN-γ (22, 32–34), but a significantly reduced intracellular persistence of the bacteria is observed when macrophages are exposed to both IFN-γ and TNF-α (22).

Macrophages are important for anti-LcrV-mediated protection against Y. pestis in the livers of infected mice (10, 29). In addition to protecting macrophages from apoptosis, it is possible that opsonization with anti-LcrV decreases the survival of Y. pestis in phagocytes, due to the ability of Fc receptors to activate bactericidal processes. The activation of macrophages with IFN-γ is known to upregulate Fc receptor expression and function, which could further increase intracellular killing, following the uptake of anti-V-opsonized Y. pestis (19, 38, 39). It is also possible that activation per se could decrease the sensitivity of Y. pestis-infected macrophages to undergo apoptosis.

In this study, we investigated how opsonization with anti-LcrV antibodies, and activation with IFN-γ, impacts the ability of Y. pestis to survive within and kill macrophages. The findings have implications for understanding how anti-LcrV antibodies and cytokines function to protect against plague.

### MATERIALS AND METHODS

**Bacterial strains.** The strains used are shown in Table 1. Y. pestis pgm mutants KIM5 and KIM5/GFP (where GFP is green fluorescent protein) contain pCD1Ap. KIM5/GFP also contains an isopropyl-β-D-thiogalactopyranoside (IPTG)-inducible plasmid encoding GFP (pMMB207gfp3.1). The YopB Δ-thiogalactopyranoside mutant contains an in-frame deletion of yopB in pCD1Ap as described by Lilje et al. (21). The KIM5 phoP mutant contains pCD1Ap and an in-frame deletion of phoP as described by Grabenstein et al. (13). Y. pseudotuberculosis serogroup 1 strain 32777, formerly known as IP2777, contains pYV.

**Sera and antibodies.** Anti-LcrV serum and control serum were obtained from LcrV-immunized or adjuvant-only injected mice, respectively, as described by Ivanov et al. (18). The monoclonal antibody (MAb) specific for LcrV (MAb 7.3) has been described previously (17). A hybridoma producing anti-YopD MAb 248.19 was generated by the Hybridoma Facility at Stony Brook University. BALB/c mice were immunized with YopBDE as described by Grabenstein et al. (13). The MAb 248.19 was generated by the Hybridoma Facility at Stony Brook University. BALB/c mice were immunized with YopBDE antigen (17). Following the fusion of spleen cells from immunized mice, hybridoma clones producing anti-YopD MAb were identified by enzyme-linked immunosorbent assay (ELISA) and immunohotting. One subclone producing anti-YopD MAb was designated 248.19, and the MAb was isolated immunoglobulin G1 (IgG1). The MAb was purified by the ammonium sulfate precipitation of hybridoma supernatants. The precipitated protein was dissolved in phosphate-buffered saline (PBS) and dialyzed against PBS.

**Primary macrophage cell culture.** Bone marrow-derived macrophages (BMDMs) were isolated from the femurs of C57BL/6 mice (Jackson Laboratory) and prepared as previously described (31). BMDMs were seeded in 24-well cell culture plates at 1.5 × 10⁶ cells per well for 24 h prior to infection in Dulbecco’s modified Eagle’s medium (Gibco) with 10% fetal bovine serum (HyClone), 1% sodium pyruvate, 2 mM glutamate, and 15% L-cell-conditioned medium.

### Infection conditions.

- **Strain Characteristic(s)**
  - **Reference**

| Strain | Characteristic(s) | Reference |
|--------|------------------|-----------|
| Y. pestis | KIM5 | pCD1Ap, pMT1°, pPCP1°, pgm mutant, Amp'| 21 |
| KIM5/GFP | KIM5/pMMB207gfp3.1, Amp', Cam° | 13 |
| KIM5 phoP | pCD1Ap phoP in-frame deletion | 13 |
| KIM5 yopB | pCD1Ap yopB in-frame deletion of nucleotides 496 to 774, Amp° | 21 |
| Y. pseudotuberculosis | 32777 | pYV° | 37 |

° Amp', ampicillin resistance; Cam°, chloramphenicol resistance.
culture medium and added to the macrophages (multiplicity of infection of 10 bacteria per macrophage). The plates were centrifuged for 5 min at 95 × g to facilitate contact between the macrophages and the bacteria and incubated for 20 min at 37°C with 5% CO₂.

**Phagocytosis assay.** RAW 264.7 mouse macrophage-like cells (ATCC TIB-71) were grown in Dulbecco’s modified Eagle’s medium plus Glutamax (Gibco BRL) with 10% heat-inactivated fetal bovine serum (HyClone) and 1 mM sodium pyruvate at 37°C with 5% CO₂. Macrophages were seeded on glass coverslips at 1.5 × 10⁵ cells in 1 ml medium in 24-well cell culture plates and incubated overnight. Macrophages were infected as described above with opsonized or nonopsonized KIM5/GFP expressing GFP. The plates were centrifuged at 95 × g for 5 min to facilitate contact between the macrophages and the bacteria and incubated for 20 min at 37°C with 5% CO₂. The wells were then washed with prewarmed PBS once and fixed with 2.5% paraformaldehyde (PFA) in PBS at room temperature for 30 min. After being washed with PBS, coverslips were blocked with 3% bovine serum albumin in PBS for 20 min. Extracellular bacteria were labeled with rabbit anti-Yersinia antiserum SB349 (5) and with goat anti-rabbit antibody conjugated to Alexa Fluor 594 (Molecular Probes). The coverslips were washed and mounted with ProLong Gold antifade reagent (Invitrogen) onto slides. The slides were examined by epifluorescence microscopy using a Zeiss Axioplan2 microscope. Pictures of three fields per coverslip in at least three independent experiments were taken. The number of macrophage-associated intracellular bacteria (green only) and the number of macrophage-associated extracellular bacteria (red and green overlay) were counted, and the percent internalization was determined as follows: [no. of intracellular bacteria/(no. of intracellular bacteria + no. of extracellular bacteria)] × 100.

**LDH release assay.** Lactate dehydrogenase (LDH) release from the supernatants collected from Y. pestis-infected BMDMs at 5 and 24 h p.i. was determined using the CytoTox-96 nonradioactive cytotoxicity assay (Promega). The supernatants were collected and centrifuged to remove cellular debris, and LDH levels were determined in triplicate. Total LDH release was determined from supernatants from freeze-thaw-lysed BMDMs. Spontaneous LDH release was determined from supernatants obtained from uninfected cells. The percentage of LDH release was calculated as follows: % LDH release = [(infected cell LDH release - spontaneous LDH release)/(total LDH release – spontaneous LDH release)] × 100.

**CFU assay.** At 25 min, 1.5 h, and 5 h p.i., Y. pestis-infected BMDMs were washed with PBS and lysed with 500 µl of 0.1% Triton X-100 in PBS. The plate was then incubated at 37°C with 5% CO₂ for 10 min. Wells were scraped and lysates collected in microcentrifuge tubes. A total of 500 µl PBS were used to wash the wells. Lysates and washes were combined and used for serial dilutions. Dilutions were plated on HI-plus-Amp plates and incubated at 28°C for 2 days. The plates were enumerated, and the log₁₀ CFU per ml were determined from the results of three independent experiments.

**TNF-α and IL-1β ELISA.** At 24 h p.i., the supernatants from wells containing infected BMDMs were collected, centrifuged for 10 min at 200 × g to remove cellular debris, and transferred to new tubes. The supernatants were diluted appropriately, and 50 µl of each diluted sample was analyzed. The concentration of TNF-α and IL-1β (IL-1β) in the supernatant was measured using the Quantikine mouse TNF-α and IL-1β immunoassay kits (R&D Systems).

**Phagosome acidification.** J774A.1 murine macrophage-like cells (ATCC TIB-67) seeded in 24-well plates on glass coverslips at 1.5 × 10⁵ cells/well were infected with KIM5 expressing GFP as described above. Determination of the colocalization of phagosomes containing GFP-positive Y. pestis with LysoTracker Red DND 99 was performed by fluorescence microscopy (34). As a positive control for the colocalization with LysoTracker Red DND 99, we used KIM5/GFP fixed in 2.5% PFA (34). One hour prior to fixation, the cell culture medium was removed from the wells, and cell culture medium containing 50 nM LysoTracker Red DND 99 was added. Coverslips were fixed with 2.5% PFA and mounted onto the slides. The percentage of KIM5/GFP that colocalized with LysoTracker Red was determined as described previously (34).

**Statistical analysis.** Experiments were performed at least three times. The results were subjected to analysis of variance and the Tukey posttest using Prism (GraphPad). The results were considered significantly different if the probability (P) values were less than 0.05.

## RESULTS AND DISCUSSION

**Opsonization with anti-V serum increases the phagocytosis of Y. pestis by naïve but not activated macrophages.** In order to study how anti-LcrV antibodies and IFN-γ impact the ability of Y. pestis to survive within and kill macrophages, we utilized mouse anti-LcrV immune serum (anti-V), previously shown to be protective (18), and cultured mouse macrophages (RAW 264.7 macrophage-like cell line or primary BMDMs). To confirm that the anti-V used in this study contained antibodies that could increase the uptake of Y. pestis by macrophages, a phagocytosis assay was performed. A culture of Y. pestis KIM5/GFP (Table 1) was shifted from 28°C to 37°C in the presence of IPTG to induce the expression of the T3SS and GFP. Samples of the bacteria were preincubated with normal mouse control serum, anti-V or PBS (nonopsonized) for 10 min prior to infection. Naïve RAW 264.7 cells were then infected with the bacteria at a multiplicity of infection of 10 for 25 min and then fixed. Phagocytosis was measured by fluorescence-based microscopy. The percent internalization of nonopsonized KIM5/GFP was ~30%, which was significantly lower than that under the control serum condition (~45%), likely due to the absence of complement proteins in the nonopsonization condition (Fig. 1A). KIM5/GFP preincubated with anti-V were phagocytosed to a significantly higher level (~65%) than bacteria preincubated with control serum (Fig. 1A), showing that this serum contained anti-LcrV antibodies that could increase the phagocytosis of Y. pestis by macrophages.

We next examined the interaction between IFN-γ-stimulated macrophages and KIM5/GFP opsonized with or without anti-V. The phagocytosis assay was performed as above except...
that RAW 264.7 cells were activated with IFN-γ (100 U/ml) for 24 h before infection. With IFN-γ-activated macrophages, there was an overall increase in bacterial uptake, but the increase was most dramatic for the nonopsonized and control serum-opsonized conditions (Fig. 1B). Approximately 55 to 60% of the nonopsonized or control-opsonized KIM5/GFP was internalized. Anti-V-opsonized KIM5/GFP was internalized at a slightly higher level (~75%), but this increase was not significant compared to that of the control. These results showed that the activation state of the macrophage was important for the ability of anti-V to increase the phagocytosis of \textit{Y. pestis}. The fact that an increased phagocytosis of \textit{Y. pestis} was observed following exposure to IFN-γ indicated that the cytokine treatment was upregulating antibacterial functions in macrophages.

The opsonization of \textit{Y. pestis} with anti-V serum decreases apoptosis in naïve but not in activated macrophages at an early stage of infection. We next examined the effect of anti-V sera on YopJ-induced macrophage death using an LDH release assay. Naïve BMDMs were infected at a multiplicity of infection of 10 with KIM5 (Table 1) grown and opsonized as described above. BMDMs were also infected in parallel with the noncytotoxic KIM5 \textit{yopB} mutant (Table 1) as a control. After phagocytosis was allowed to occur for 25 min, the survival of extracellular bacteria was prevented by the addition of a low concentration of gentamicin to the tissue culture medium. This infection protocol was used because it allows for the measurement of intracellular survival and apoptosis under the same infection conditions. KIM5, being a \textit{pgm} mutant, lacks the \textit{ripCBA} genes required for the growth of \textit{Y. pestis} in activated macrophages (33), and therefore, to avoid decreases in survival due to the absence of the \textit{ripCBA} genes, the CFU-based survival assays (see Fig. 3) were limited to a short time period (5 h). LDH release was initially determined at 5 h p.i. The results showed that macrophages infected with the nonopsonized \textit{yopB} mutant, deficient in Yop translocation, released very small amounts of LDH, about 2.5% of the total, by 5 h of infection (Fig. 2A). This was a significantly lower LDH release than that of BMDMs infected with control serum-opsonized KIM5 (~18%). Compared to BMDMs infected with control serum-opsonized KIM5, LDH was also significantly lower when KIM5 was opsonized with anti-V (3%) but not when the bacteria were left unopsonized (12.5%). These results were consistent with previous studies, which showed that rabbit polyclonal anti-LcrV antibodies could reduce Yop translocation and decrease apoptosis in naïve J774A.1 macrophage-like cells infected with \textit{Y. pestis} (10, 29, 43).

When the LDH assay was performed with IFN-γ-activated macrophages, only the BMDMs infected with the nonopsonized \textit{yopB} mutant released significantly lower levels of LDH than the control serum (2.3% versus 18%) (Fig. 2C). Thus, the ability of anti-V to inhibit apoptosis was diminished in activated macrophages.

Opsonization with anti-V does not decrease the survival of \textit{Y. pestis} in activated macrophages. To determine if the opsonization of \textit{Y. pestis} with anti-V would result in increased killing within BMDMs, we performed a CFU assay at various times p.i. Naïve BMDMs were infected with opsonized or nonopsonized bacteria as above for the LDH assay. As a preliminary test to confirm that the BMDMs used were bactericidal under...
the conditions of the assay, a CFU assay was performed with *Y. pseudotuberculosis* grown alongside KIM5 at 37°C. The internalization of nonopsonized 37°C-grown *Y. pseudotuberculosis* into BMDMs triggers a bactericidal process that requires macrophage sensing of the T3SS (35, 48). As shown in Fig. 3A, by 5 h p.i there was a significant decrease in the number of CFU for *Y. pseudotuberculosis* 32777. As reported previously (21), there was no significant decrease in the number of CFU over the same time period in BMDMs infected with nonopsonized KIM5 (Fig. 3A). Although it is unclear why the T3SS of *Y. pestis* KIM5 did not stimulate intracellular killing in BMDMs, these results confirmed that the macrophages used were bactericidal. Furthermore, because the T3SS of KIM5 did not stimulate intracellular killing in BMDMs, we did not face the problem of multiple bactericidal processes occurring simultaneously, which could complicate the analysis of the potential role of anti-V opsonization and IFN-γ activation in the decreasing survival of *Y. pestis* in macrophages.

Next, CFU assays were carried out using opsonized or nonopsonized KIM5. A KIM5 phoP mutant (Table 1), shown previously to be highly defective for intracellular survival (13), was included as a control. As shown in Fig. 3B, by 5 h p.i., there was a significant (~2-log) decrease in the number of intracellular CFU that could be recovered from the macrophages infected with nonopsonized KIM5 phoP. In contrast, no significant decrease in the number of CFU was observed for the nonopsonized KIM5 yopB mutant, or KIM5 under any condition (nonopsonized, anti-V, or control serum) (Fig. 3B). Based on these results, we concluded that opsonization with anti-V does not lead to intracellular killing of KIM5 by naïve BMDMs.

Seeing no evidence for the intracellular killing of anti-V-opsonized KIM5 in naïve BMDMs, a CFU assay was performed with macrophages activated with IFN-γ (Fig. 3C). As with the naïve BMDMs, we used KIM5 phoP as a control for intracellular killing. The results obtained with IFN-γ-stimulated BMDMs were similar to those seen with naïve BMDMs, as only KIM5 phoP showed a significant decrease in the number of CFU by 5 h p.i. (Fig. 3C). Therefore, *Y. pestis* that was opsonized with anti-V prior to uptake was able to survive in macrophages that were activated with IFN-γ.

**The opsonization of *Y. pestis* with anti-V serum does not decrease apoptosis in naïve or activated macrophages at a late stage of infection.** Under the infection conditions used in this study, in which internalized *Y. pestis* survives within macrophages (Fig. 3) (21), the kinetics of apoptosis is extended, such that LDH release is first detected by 5 h p.i. (early stage) (Fig. 2), and continues to increase over the next 19 h (late stage) (21). To determine if opsonization with anti-V would affect apoptosis at a late stage of infection, an LDH release assay was performed on *Y. pestis*-infected naïve or activated macrophages. The results with naïve macrophages was unexpected, since we had observed a ~sixfold decrease in apoptosis in BMDMs infected with anti-V-opsonized KIM5 compared to that with control-opsonized KIM5 at 5 h p.i. (Fig. 2A) and assumed this proportional difference would be maintained to the 24-h time point (Fig. 2B). Instead, the results suggested that apoptosis was accelerated in BMDMs infected with anti-V-opsonized *Y. pestis* between 5 and 24 h compared to that with the control infection. Since anti-V-opsonized KIM5 was internalized by naïve macrophages at a higher level than control-opsonized KIM5 (Fig. 1A), it was possible that larger numbers of internalized bacteria allowed for higher levels of cell death between 5 and 24 h p.i.

Macrophages treated with IFN-γ appeared to be less sensitive to YopJ-dependent cell death, as the overall levels of LDH released from KIM5-infected BMDMs were lower in activated cells than the naïve phagocytes (compare Fig. 2B and D).
Activation with lipopolysaccharide desensitizes macrophages to YopJ/YopP-induced apoptosis during infection with *Y. pseudotuberculosis* or *Y. enterocolitica* (4, 36), and a similar phenomenon may be operating in IFN-γ-activated BMDMs infected with *Y. pestis*.

The opsonization of *Y. pestis* with anti-V serum does not alter the secretion of TNF-α by infected macrophages. TNF-α has been shown to cooperate with IFN-γ to promote protective immunity against *Y. pestis* (39). In addition, macrophages activated with TNF-α and IFN-γ are more bactericidal against *Y. pestis* than macrophages treated with IFN-γ alone (22). Previous studies have shown that *Y. pestis* can partially suppress the secretion of TNF-α from infected macrophages and that YopJ is required for this suppression (21, 46). Lilo et al. showed that after 24 h of infection, ~2,000 pg/ml TNF-α was released by KIM5-infected BMDMs (21). We wanted to determine if TNF-α was secreted during our infections with KIM5 and to what degree opsonization affected TNF-α levels. In parallel, we determined levels of another cytokine (IL-1β) that is secreted from KIM5-infected macrophages in a YopJ-dependent manner (21). BMDMs were infected with opsonized or nonopsonized KIM5 or nonopsonized KIM5 *yopB* (Fig. 4). Supernatants were collected at 24 h, and ELISA was performed to measure TNF-α and IL-1β levels. We found that with BMDMs infected with KIM5 *yopB*, there was ~5,000 pg/ml TNF-α released (Fig. 4A). Consistent with the concept that YopJ can interfere with the production of TNF-α, the level of TNF-α secreted was diminished to ~2,500 pg/ml in KIM5-infected macrophages. When KIM5 was opsonized with control serum or anti-V, the levels of TNF-α that were secreted from the infected macrophages were similar to those secreted from macrophages infected with nonopsonized KIM5 (Fig. 4A), showing that anti-V did not significantly affect the secretion of TNF-α. Additionally, opsonization with anti-V did not affect the secretion of IL-1β from BMDMs infected with KIM5 (Fig. 4B).

The opsonization of *Y. pestis* with anti-LcrV MAb 7.3 does not decrease survival in macrophages or inhibit late-stage apoptosis. Polyclonal antibodies and MAbs that are specific for the same antigen can exhibit different activities with respect to neutralizing function (7). The anti-V used in these experiments had been shown to passively protect mice against *Y. pestis* KIM5, a conditionally virulent strain, in an intravenous infection model (18). However, it was desirable to repeat the above experiments using a well-characterized protective MAb specific for LcrV. The opsonization of *Y. pestis* with anti-LcrV IgG1 MAb 7.3 increases phagocytosis and decreases apoptosis in naïve J774A.1 cells (43). In addition, MAb 7.3 is known to provide protection against fully virulent *Y. pestis* in murine bubonic and pneumonic plague infections (15–17). We opsonized KIM5/GFP or KIM5 with MAb 7.3 and repeated the phagocytosis, intracellular survival, and LDH release assays. In parallel, macrophages were infected with *Y. pestis* that was left nonopsonized or incubated with an IgG1 isotype control MAb specific for YopD (MAb 248.19). Figure 5 shows that MAb 7.3...
The colocalization of mouse macrophage-like cells and used microscopy to measure would increase phagosome acidification, we infected J774A.1 phagosomes in macrophages have used 28°C-grown nonopsonized Y. pestis have shown a block in the acidification of Y. pestis (Fig. 7C).

Anti-V opsonization does not overcome the block in acidification of Y. pestis-containing phagosomes. To date, studies that have shown a block in the acidification of Y. pestis-containing phagosomes in macrophages have used 28°C-grown nonopsonized bacteria (34). To determine if opsonization with anti-V would increase phagosome acidification, we infected J774A.1 mouse macrophage-like cells and used microscopy to measure the colocalization of Y. pestis-containing phagosomes with the acidotropic probe LysoTracker Red DND-99 (34). Prior to infection, KIM5/GFP was grown at 28°C for 2 h or at 37°C for 2 h and then not opsonized or opsonized with control serum or anti-V. The colocalization of GFP-expressing KIM5 and LysoTracker Red was determined at 1.25 h p.i. As a positive control for phagosome acidification, we used KIM5/GFP fixed with PFA. Only phagosomes containing fixed KIM5/GFP (both grown at 28°C and 37°C) showed significantly increased colocalization with LysoTracker Red (>90%) than the standard condition (nonopsonized, 28°C) (Fig. 8). We concluded that opsonization with anti-V does not overcome the block in phagosome acidification imposed by Y. pestis in macrophages.

Conclusions. This study focused on investigating the potential cooperative effect of anti-V opsonization and IFN-γ activation in allowing macrophages to protect themselves from being colonized and killed by Y. pestis. We conclude that Y. pestis can efficiently evade the combined effects of anti-V opsonization and IFN-γ activation in macrophages due to its ability to block phagosome acidification. In fact, we obtained evidence that activation with IFN-γ could be counterproductive for antibody-mediated protection, as anti-V was unable to significantly reduce apoptosis at the early time point in activated macrophages infected with KIM5 (although this effect was not seen with MAb 7.3). On the other hand, activation with IFN-γ increased overall levels of bacterial phagocytosis and decreased apoptosis at late stages, illustrating a protective activity of this cytokine.

We also obtained evidence that the ability of anti-V to inhibit apoptosis decreased as the time of the infection was extended. This suggests that intracellular bacteria can contribute to apoptosis and that anti-V is not neutralizing once the bacteria are internalized. It is not clear if the intracellular survival of Y. pestis is important for late-stage apoptosis. KIM5 phoP does not survive intracellularly (Fig. 3B and C); however, BMDMs infected with this mutant released levels of LDH at 5 h and 24 h p.i. similar to those of BMDMs infected with KIM5 (data not shown). In the case of Y. pseudotuberculosis, the results suggest that protein synthesis is required for intracellular bacteria to induce apoptosis (48). Perhaps internalized KIM5 causes increased apoptosis by continuing to synthesize and translocate YopJ for a limited period of time within nascent phagosomes, but bacterial survival is not required per se. The possibility that Y. pestis can synthesize and translocate small amounts of YopJ from within the phagosome is not necessarily in conflict with a previous study showing that no delivery of YopH could be detected by immunoblotting after the first 30 to 45 min of infection of macrophages (10). Immunoblotting may not be sensitive enough to detect the translocation of biologically significant amounts of a Yop. Alternatively, some Yops may be more efficiently translocated from an intracellular location (e.g., YopJ) than other Yops (e.g., YopH).

In our studies, we activated macrophages using IFN-γ only. There is evidence that TNF-α cooperates with IFN-γ to kill Y. pestis in macrophages in the absence of anti-V (22). Perhaps the addition of TNF-α is required along with anti-V and IFN-γ in macrophages to increase the killing of Y. pestis. While we did not add exogenous TNF-α in our CFU assays, we note that KIM5-infected macrophages secrete detectable levels of TNF-α by 4 h p.i. (21) and that ~2,500 pg/ml TNF-α was detected in supernatants by 24 h p.i. (Fig. 4). Therefore, we can conclude that Y. pestis can bypass protection mediated by...
anti-V antibodies, preactivation with IFN-γ, and autocrine activation mediated by endogenous TNF-α produced during infection with KIM5. An additional consideration is that the experiments described here utilized the KIM5 pgm mutant, which lacks the ripCBA genes needed for the long-term survival and growth of Y. pestis in IFN-γ-activated macrophages (33). Because of this, our intracellular survival assays were restricted to a maximum of a 5-h time period. Use of a fully virulent pgm+/H11001 strain would have allowed for the use of longer time points in CFU assays, and a role for anti-V in the killing of Y. pestis in IFN-γ-activated macrophages might be revealed at later time periods.

These results have implications for understanding the mechanism of protective immunity to plague mediated by anti-V and phagocyte-activating cytokines. As our results suggest that opsonization with anti-V does not dramatically increase the killing of Y. pestis by activated macrophages, it is possible that neutrophils may be more critical for protection than macrophages (10, 29). Consistent with this idea is the demonstration that macrophages are not essential for anti-V-mediated protection against Y. pestis in the spleens of infected mice (10, 29), while neutrophils appear to be important for anti-V-mediated protection in both the spleens and livers of mice challenged with plague (10). It is possible that TNF-α cooperates with IFN-γ to upregulate microbicidal processes in neutrophils. Anti-V increases the phagocytosis of Y. pestis by murine neutrophils (10). Y. pestis is unable to survive in rodent (8) or human (40) neutrophils. It has been shown that Yop delivery does not protect intracellular Y. pestis from neutrophil killing.

FIG. 7. Effect of MAbs on YopJ-mediated cell death through determination of LDH release. Naïve BMDMs (A and B) or IFN-γ-stimulated BMDMs (C and D) were infected with the indicated strains of KIM5 with or without preincubation with MAbs or left uninfected. LDH levels in the supernatants of the BMDMs at 5 h (A and C) and 24 h (B and D) p.i. were measured as indicated in the legend to Fig. 2. Results were taken from three independent experiments and averaged. Error bars show standard deviations. *, P < 0.05; **, P < 0.01 compared to MAb YopD.

FIG. 8. Effect of bacterial growth, temperature, and serum opsonization on phagosome acidification within naïve J774A.1 cells. J774A.1 cells were infected with fixed or live KIM5/GFP pregrown at 28°C or 37°C and left unopsonized or opsonized as indicated. Lyso-Tracker Red DND 99 was used to label acidic compartments, and colocalization with KIM5/GFP was determined at 1.25 h p.i. by fluorescence microscopy. Results were taken from three independent experiments and averaged. Error bars show standard deviations. ***, P < 0.001 compared to the unopsonized KIM5 28°C subculture.

FIG. 8. Effect of bacterial growth, temperature, and serum opsonization on phagosome acidification within naïve J774A.1 cells. J774A.1 cells were infected with fixed or live KIM5/GFP pregrown at 28°C or 37°C and left unopsonized or opsonized as indicated. Lyso-Tracker Red DND 99 was used to label acidic compartments, and colocalization with KIM5/GFP was determined at 1.25 h p.i. by fluorescence microscopy. Results were taken from three independent experiments and averaged. Error bars show standard deviations. ***, P < 0.001 compared to the unopsonized KIM5 28°C subculture.
(40). However, whether anti-V opsonization accelerates the killing of *Y. pestis* in neutrophils has not been examined.

The neutralization of apoptosis in *Y. pestis*-infected macrophages by anti-V is commonly used as an in vitro correlate of protective immunity (3, 43–45). Although this assay has many positive features, including ease of use, reproducibility, and a quantifiable readout, we feel the following points should be considered with respect to improving the in vitro correlate of protective immunity assay. First, our results showing that anti-V can promote the uptake of *Y. pestis* into macrophages without killing the bacteria, and that anti-V may not inhibit the translocation of YopJ by the intracellular population, indicate that the time point at which apoptosis levels are quantified is critically important. In addition, the activation status of the macrophage is an important variable, as we found that IFN-γ-activated macrophages are less sensitive to apoptosis. Finally, *Y. pestis* strains differ in their ability to induce apoptosis (KIM has high activity [21]) and not all strains require apoptosis for virulence (20, 47). Therefore, there continues to be a need to develop an anti-LcrV neutralization assay that does not rely on macrophage apoptosis as a readout.

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