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Source / Izvornik: Microbes and Infection, 2010, 12, 126 - 134

Journal article, Accepted version Rad u časopisu, Završna verzija rukopisa prihvaćena za objavljivanje (postprint)

https://doi.org/10.1016/j.micinf.2009.11.003

Permanent link / Trajna poveznica: https://urn.nsk.hr/urn:nbn:hr:184:393035

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Download date / Datum preuzimanja: 2025-03-28



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Author Manuscript

Microbes Infect. Author manuscript; available in PMC 2011 February 1

Published in final edited form as:

Microbes Infect. 2010 February ; 12(2): 126. doi:10.1016/j.micinf.2009.11.003.

Regulation of apoptosis and anti-apoptosis signalling by *Francisella tularensis*

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Abstract

Francisella tularensis induces apoptosis within macrophages but the temporal and spatial modulation through activation of caspase-1, caspase-3, and the anti-apoptosis nuclear transcription factor B (NF- κ B) is not known. Whether escape of the bacteria into the cytosol is sufficient and/or essential for activation of NF- κ B is not known. Our results show that *F. tularensis* subsp. *novicida* induces sustained nuclear translocation of NF- κ B at early time points after infection of human monocytes derived macrophages (hMDMs). The sustained nuclear translocation of NF- κ B is defective in the *iglC* mutant that fails to escape into the cytosol of macrophages. Nuclear translocation of NF- κ B by the wild type strain is abolished upon treatment with the NF- κ B inhibitor caffein acid phenyl ester. While the wild type strain triggers caspase-3 and caspase-1 activation by 6 h post-infection the *iglC* mutant is defective in triggering both caspases. In hMDMs treated with the apoptosis-inducing agent, staurosporin, there is an induction of cell death in the *iglC* mutant-infected macrophages are resistant to cell death-induced agent. We conclude that although caspase-1 and caspase-3 are triggered within *F. tularensis*-infected hMDMs during early stages of infection, cell death is delayed, which is correlated with simultaneous activation of NF- κ B.

Keywords

tularemia; iglC; cell death

1. Introduction

Francisella tularensis is a gram-negative intracellular bacterium and the etiological agent of tularemia. There are four closely related subspecies of *F. tularensis (tularensis, holarctica, mediasiatica* and *novicida*), and subspecies *tularensis* is the most virulent to humans [1,2]. *F. tularensis* subsp. *novicida* is attenuated in humans but causes disease in mice similar to that

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caused by subsp. *tularensis*, and is an attractive model to study pathogenesis of tularenia [3]. Relatively little is known about intracellular trafficking of *F. tularensis*. The *F. tularensis*-containing phagosome (FCP) of subsp *tularensis*, *holarctica*, and *novicida* matures to a late endosome-like phagosome that has a limited fusion to the lysosomes [1,3,4]. The FCP acquires the proton vATPase pump and is acidified within 15 min of its biogenesis, and the acidification is essential for disruption of the FCP and escape of *F. tularensis* into the macrophage cytosol, similar to *L. monocytogenes* [1,5]. The *Francisella* pathogenicity island (FPI) gene *iglC*, and its regulator MglA are essential for bacterial escape into the cytosol of hMDMs, since mutants in both genes are defective in escape into the cytosol and their phagosomes fuse to lysosomes [1,6].

Several bacterial pathogens induce apoptosis in mammalian cells by activating specific components of the apoptotic pathways [7]. Caspases play essential roles in apoptotic cell death [8]. Two major apoptotic pathways, designated the extrinsic and intrinsic pathways, mediate the activation of apoptotic caspases in response to various extra- and intracellular apoptotic stimuli [8]. The two main apoptotic pathways converge on the activation of caspase-3 [9]. The expression of many anti-apoptotic genes is regulated by the nuclear transcription factor kappa-B (NF- κ B) that triggers the expression of two families of genes involved in inflammation and anti-apoptosis [10]. In activated macrophages, the ability of *Francisella* to escape the phagosome and replicate in the cytosol correlates with the activation of the inflammasome, which contains the host proteins ASC and caspase-1 [1,11]. Inflammasome activation is critical for innate host defense and leads to the induction of cell death in infected cells and the concomitant release of the proinflammatory cytokines IL-1 β and IL-18 [1,11,12]. *F. tularensis* LVS induces apoptosis in the J774A.1 murine macrophage cell line through a pathway partly resembling the intrinsic apoptotic pathway and is not thought to involve caspase 1, caspase 8, Bcl-2, or Bid [1,13,14].

The transcription factor NF- κ B plays a crucial role in regulation of apoptosis by triggering expression of various anti-apoptotic genes [15]. NF- κ B represents a family of homo- and heterodimer transcription factors, and the p65/p50 heterodimer is the most predominant active complex in mammalian cells [15]. In resting cells, NF- κ B proteins are predominantly sequestered in the cytoplasm by the NF- κ B inhibitory proteins (IkBs) [16]. The I κ B kinase mediates phosphorylation of I κ Bs, followed by ubiquitination and proteosomal degradation, which is crucial to the activation and nuclear translocation of NF- κ B [16]. Inhibition of NF- κ B results in a differential effect on the levels of pro- and anti-inflammatory cytokine production in response to infection by the LVS strain of *F. tularensis* [1,17,18]. Inhibition of PI3K/Akt results in suppression of *F. tularensis* subsp *novicida*–induced cytokine production through the inhibition of NF- κ B [19].

In this study, we examined the temporal and spatial modulation of *F. tularensis* induced apoptosis through activation of caspase-1, caspase-3, and the anti-apoptosis nuclear transcription factor B (NF- κ B), and the role of phagosomal escape in modulation of these cellular processes at various stages of infection. Our data show that *F. tularensis* subsp. *novicida* induces sustained nuclear translocation of NF- κ B, initiated at early time points after infection of hMDMs, which correlates with down-regulation of cell death despite robust activation of the executioner caspase-1 and caspase-3, tipping the balance in favour of survival of the infected cells till termination of intracellular proliferation.

2. Material and methods

2.1. Bacteria and macrophages

The wild-type (wt) *F. tularensis* subsp *novicida* strain U112 and it isogenic *iglC* mutant has been described previously [20]. The tetracyclin resistant plasmid pKK214 encodes gfp was

introduced to the *F. tularensis* strains. *Legionella pneumophila* strain AA100 expressing GFP has been grown as described previously [21]. To prepare hMDMs, peripheral blood monocytes were isolated from healthy volunteers with no history of tularemia or Legionnaires' disease and hMDMs were prepared as we described previously. Obtaining blood was approved by the institutional IRB with a consent form according to standard federal laws.

2.2. Confocal microscopy

In all confocal microscopy experiments, the cells were infected with bacteria at a MOI of 10 for 1 h, followed by 3x washing and 1 h of gentamicin treatment (50 μ g ml⁻¹) to remove extracellular bacteria. All experiments were done in triplicate and a minimum of 100 cells per sample were examined from different coverslips. On average, 8–15 0.2 μ m serial z sections of each image were captured and stored for further analyses, using Adobe Photoshop 6.0 (Adobe Photoshop).

2.3. Apoptosis resistance of F. tularensis infected cells

To examine the sensitivity of *F. tularensis*-infected hMDMs to staurosporin-induced apoptosis, hMDMs were infected with the wt-GFP or the mutant strain *iglC*-GFP and either left untreated or treated with Staurosporin (50 nM) (Sigma, St Louis, MO). Treatment with staurosporine was carried out for 4 h starting at 8 h after infection. Control uninfected monolayers were either left untreated or treated with the above mentioned staurosporine for the same time periods. To examine the sensitivity of *F. tularensis*-infected hMDMs cells to CAPE-induced apoptosis, hMDMs were treated with 20 μ g/ml CAPE for 30 min prior to infection as previously described [22].

2.4. Nuclear translocation of the p65 subunit of NF-kB in F. tularensis-infected cells

In all experiments, uninfected untreated cells, *E. coli* LPS-treated cells ($1 \mu g m l^{-1}$ for 20 min) and *L. pneumophla* AA100 infected cells (Sigma, St Louis, MO) were used as a negative and positive controls, respectively, for nuclear translocation of the p65 subunit of NF- κ B. To examine nuclear translocation of the p65 subunit in *F. tularensis*-infected cells, monolayers of hMDMs were infected with either wt-GFP or *iglC*-GFP and proceed by the protocol described previously [22].

2.5. Caspase 1 and Caspase 3 activation

To assess active caspase-1 or active caspase-3 staining by confocal microscopy, hMDMs on glass coverslips were infected with either wt-GFP or *iglC*-GFP strains at an MOI of 10 for 1, 6, and 24 h and proceed as described previously [22]. Briefly, macrophages were stained for 1 h with 6-carboxyfluorescein-YVAD fluoromethylketone (Immunochemistry Technologies) and caspase-3 inhibitor (BD, San Diego, CA) as recommended by the manufacturer. As a positive control for the hMDMs, macrophages were treated with 10 mM simvastatin (Calbiochem). For labeling of the caspase 3 positive cells, rabbit polyclonal antiactive caspase-3 antiserum (BD, San Diego, CA) and caspase-1 inhibitor were used (Santa Cruz Biotechnology, Santa Cruz, CA) for 1 h, and then incubated for 1 h with a goat anti-rabbit immunoglobulin G secondary antibody conjugated to Alexa red (Molecular Probes, Inc., Eugene, OR). Apoptotic nuclei were labled using an apoptosis detection kit (TUNEL) according to the manufacturer's instructions (Boehringer Mannheim Corporation, Indianapolis, IN).

3. Results

3.1. F. tularensis subsp novicida induces sustained nuclear translocation of NF-kB in hMDMs

In mammalian cells, the most predominant active complex of NF- κ B is the p65/p50 heterodimer [16]. Although activation of NF- κ B has been reported during *F. tularensis* LVS infection [17,18], the temporal and spatial signaling mechanisms involved in modulation of macrophage survival and death are not well described. Thus, we examined whether *F. tularensis* triggered nuclear translocation of the p65 subunit of NF- κ B in hMDMs. The *iglC* mutant was used as a control, since it does not escape from the phagosome [6]. That would allow us to decipher whether bacterial escape from the phagosome is essential for modulation of various cellular processes involved in survival of the infected host cell.

Using laser scanning confocal microscopy our results showed that NF- κ B was translocated to the nuclei of only 7% of uninfected macrophages, while NF- κ B was translocated to the nucleus in 85–95% of strain AA00 of *L. pneumophila*-infected hMDMs at all time points examined (Fig. 1A and B). The p65 subunit of NF- κ B was translocated to the nucleus in ~85% of *E. coli* LPS-treated macrophages (Fig. 1A and B). By 1h after infection, 80–95% of the wt-infected macrophages showed nuclear translocation of NF- κ B (Fig. 1 A and B). In contrast, only 25% of the *iglC* mutant-infected macrophages exhibited nuclear translocation of NF- κ B, at 1h after infection (Fig. 1 A and B).

By 6–48 h after infection, 35–45% of the wt strain-infected macrophages showed sustained nuclear translocation of NF- κ B (Fig. 1A and B), but only 8–15% of the *iglC* mutant-infected macrophages showed nuclear translocation of NF- κ B (Fig. 1 A and B). There was statistically significant difference between nuclear translocation of NF- κ B by the wt strain and the *iglC* mutant strain at all time points during infection (Student *t*-test, *p*<0.001). We conclude that bacterial escape into the host cell cytosol by 60 min post-infection is essential for early nuclear translocation of NF- κ B.

To examine whether *F. tularensis* subsp. *novicida* LPS is involved in *F. tularensis* induced nuclear translocation of NF- κ B, formalin-killed bacteria were used for infection. At 30 min after infection, nuclear translocation of NF- κ B was detected in only 5 % of hMDMs infected with the formalin-killed WT strain (Fig. 1 A and B), which was similar to uninfected hMDMs. These data may suggest that the *F. tularensis* subsp. *novicida* LPS is unlikely to be involved in the induction of nuclear translocation of NF- κ B. However, it can't be excluded that formalin-sensitive LPS components may be involved.

3.2. CAPE blocks sustained activation of NF-kB in human macrophages

We utilized caffeic acid phenethyl ester, which inhibits NF- κ B activation in response to various stimuli, such as LPS to examine its effect on the infection by *F. tularensis* subsp. *novicida* [23]. We used *L. pneumophila* as a positive control, since CAPE does not inhibit nuclear translocation of NF- κ B in *L. pneumophila*-infected macrophages [22]. Since CAPE blocks the LPS mediated activation of NF- κ B, LPS-treated cells in presence of CAPE were used as a control.

Our data showed that at 30 min after infection in the presence of CAPE, nuclear translocation of NF- κ B was detected only in 20 % of the wt strain infected macrophages while 80% of strain AA100 of *L. pneumophila* infected hMDMs treated with CAPE exhibited nuclear translocation (Fig. 2 A and B).

By 1 and 6 h after infection, only ~10% of the wt strain-infected hMDMs that were treated with CAPE exhibited nuclear translocation of NF- κ B (Student *t*-test, *p*<0.001) (Fig. 2 A and

B). The data indicate CAPE blocks NF-κB activation by *F. tularensis* subsp. *novicida*, which is very distinct from *L. pneumophila*.

3.3. Modulation of apoptosis via nuclear translocation of NF-kB by *F. tularensis* subsp. *novicida*

It has been shown that *F. tularensis* induce apoptosis in murine macrophages [13,24]. We examined the role of sustained nuclear translocation of NF- κ B activation in modulation of apoptosis in *F. tularensis*-infected human macrophages. We used *L. pneumophila* as a control, since nuclear translocation of NF- κ B in *L. pneumophila*-infected macrophages is required for resistance to apoptosis-inducing agents [22].

Our data showed that only 5–10% of untreated uninfected, wt-infected and *iglC* mutantinfected cells were TUNEL-positive by 1–6 h after infection (Fig. 3B). In CAPE treated cells, 23–40% of uninfected, wt-infected and *iglC* mutant-infected cells were TUNEL-positive, which was significantly different from untreated cells (Student *t*-test, p<0.01) (Fig. 3 A and B). As expected, CAPE did not enhance apoptosis in *L. pneumophila*-infected control cells [22]. Importantly, CAPE does not activate caspases directly.

By 24 h after infection, 70% of the untreated cells infected with the wt strain of *F. tularensis* were TUNEL-positive while 85% of CAPE-treated infected cells underwent apoptosis (Student *t*-test, p<0.01) (Fig. 3 A and B). In untreated cells infected with the *iglC* mutant 16% of the cells were TUNEL-positive. In contrast, in CAPE treated cells 45% of the *iglC* mutant-infected cells underwent cell death, similar to *L. pneumophila*. We conclude that sustained nuclear translocation of NF- κ B by the wt strain contributes modestly but significantly to down-regulation of cell death in *F. tularensis* subsp. *novicida* infected human macrophages.

3.4. The FPI-encoded protein IgIC plays an important role in cell death through caspase-1 activation

F. tularensis escapes from the phagosome into the cytosol of macrophages within 30–60 min of entry into the cells (See [1] for a recent review). *F. tularensis* activates the inflammasome though caspase-1, and the *iglC* mutant is defective in caspase-1 activation within mice macrophages [25]. It is not known whether bacterial escape by itself or subsequent replication within the cytosol after phagosomal escape of *F. tularensis* is essential for triggering caspase-1 activation within human macrophages. Therefore, we used single cell analyses to determine, simultaneously, the kinetics of activation of caspase-1 and apoptosis in hMDMs infected by the *iglC* mutant and compared it to the wt strain.

The data showed at 1 h after infection, the wt strain of *F. tularensis* subsp. *novicida* induced caspase-1 activation in 45% of the infected cells but there were no signs of cell death at that time point (Fig. 4 A and B). In contrast, the *iglC* mutant did not trigger caspase-1 activation at 1 h after infection (Student *t*-test, p<0.001) (Fig. 4 A and B).

By 6 h after infection with the wt strain, 60% of the cells were active for caspase-1 but only 20% of the cells were also TUNEL-positive; thus, positive for both TUNEL and caspase-1 (Fig. 4 A and B). At 6 h after infection only ~10 % of the cells infected with the *iglC* mutant exhibited caspase-1 activation (Fig. 4 A and B) (Student *t*-test, p), and were also TUNEL-positive (Student*t*-test, <math>p < 0.001) (Fig. 4 A and B). Therefore, bacterial escape into the cytosol is essential for caspase-1 activation during early stages of the infection as also showed by Mariathasan et al. [11]

By 24 h after infection by the wt strain, 70% of the cells were positive for both caspase-1 and TUNEL (Fig. 4 A and B), but only 16% of the *iglC* mutant-infected cells induced caspase-1 activation and 10% were also TUNEL-positive (Fig. 4 A and B) (Student *t*-test, p<0.0002).

Our results showed that *F. tularensis* subsp *novicida* induced activation of caspase-1 during early stages of infection of hMDMs which was associated with the induction of delayed cell death by 24 h. The FPI-encoded protein IglC plays an important role in cell death through caspase-1 activation. Cell death is likely to be mediated by multiple processes including caspase-1, caspase-3, and other caspases-independent processes.

3.5. Bacterial escape is essential for caspase-3 activation in F. tularensis infected hMDMs

Although caspase-3 has been shown to be triggered by *F. tularensis* within the J744.1 murine macrophage cell line [24], it is not known whether this occurs in primary cells and whether escape and replication of the bacteria in the cytosol is also required for triggering caspase-3 activation.

The data showed that at 1h after infection approximately 20% of the cells infected by the wt strain exhibited activation of caspase-3, which was significantly different from the *iglC* mutant or uninfected cells (Student *t*-test, p<0.001). Only a few cells were positive for TUNEL at this time point (Fig. 5 B).

At 6 h post-infection, 46% of wt-infected hMDMs exhibited caspase-3 activation, which was significantly different from cells infected by the *iglC* mutant (13%) (Student *t*-test, *p*<0.0009). Only 20% of the wt strain-infected cells that exhibited caspase-3 activation became TUNEL-positive (Fig. 5 B). We conclude that bacterial escape into the cytosol is sufficient and essential for early activation of caspase-3 prior to bacterial proliferation within the cytosol. Importantly, despite caspase-3 activation, cell death is not triggered till later stages of the infection.

By 24 h after infection by the wt strain, a large number of the cells were lysed, and 80% of the remaining infected cells exhibited caspase-3 activation but only 45% were also TUNEL-positive (Fig. 5 A and B). In contrast, the cells infected with the *iglC* mutant strains exhibited minimal frequency of caspase-3 activation or apoptosis (Fig. 5 A and B).

We conclude that bacterial escape into the cytosol is essential and sufficient for the induction of caspase-3 activation during early stages of the infection. Although there is robust caspase-3 activation in *F. tularensis* infected hMDMs, cell death is largely delayed, suggesting potential anti-apoptotic signalling in the infected cells. The delayed cells death is likely mediated by caspases as well as caspases-independent processes.

3.6. Resistance of infected macrophages to apoptotic stimuli

Caspase-3 is the executioner of apoptosis and its activation signals the point of 'no return" in the apoptotic pathways. Since many infected cells exhibited robust activation of caspase-3 and caspase-1 but were not apoptotic, the delay in cell death of *F. tularensis*-infected cells may be correlated with up-regulation of anti-apoptotic signalling.

At 6 h after infection, more then 70% of the wt strain-infected cells treated with 50 nM staurosporin (ST) showed activated caspase-3 and were also TUNEL-positive, which was significantly different from uninfected cells (Student *t*-test, p<0.01). At all time points after infection, less then 50% of the *iglC* mutant infected cells exhibited activation of caspase-3 and were also TUNEL-positive (Fig. 6 A and B). These data showed that *F. tularensis*-infected cells exhibited modest but significant resistance to potent external apoptotic stimuli, compared to uninfected cells, and this correlated with the activation of anti-apoptotic signalling.

4. Discussion

Francisella tularensis is a highly contagious, facultative intracellular bacterium that causes tularaemia in humans and animals [1]. *F. tularensis* rapidly escape the phagosome and replicate

in the cytosol of host cells [1]. It is not known whether escape of *F. tularensis* from the phagosome into the cytosol and subsequent intracellular proliferation are critical for innate host defence mechanisms such as modulation of apoptosis through caspases and NF-k β . In addition, since *F. tularensis* triggers caspase-1 and caspase-3 activation along with apoptosis and activation of NF- κ B [1], the temporal and spatial modulation of these cellular processes is not known. Therefore, our study is the first to determine all of these processes in the same system under the same experimental conditions. Our data clearly show that despite activation of caspase-1 and caspase-3 during early stages of the infection by *F. tularensis*, cell death is delayed, which is correlated with the activation of NF- κ B.

Fine-tuning caspases activation along with modulation of pro-apoptotic, anti-apoptotic, as well as proinflammatory processes through temporal modulation of NF-κB is likely to be an important aspect of the pathogenic evolution of *F. tularensis*. Several intracellular pathogens, such as *Legionella pneumophila*, *Rickettsia rickettsii* and, *Toxoplasma gondii* inhibit apoptosis of their host cells by a mechanism that involves NF- κB activation [22,26,27]. *F. tularensis* subsp *novicida mglA* mutant displays enhanced NF-κB activation and enhanced proinflammatory cytokine production compared to cells infected with the wild type strain [28]. Our results have shown that *F. tularensis* subsp. *novicida* induces nuclear translocation of NF-κB at early time points after infection of hMDMs. The sustained nuclear translocation of NF-κB in *F. tularensis* infected macrophages is defective in the *iglC* mutant that fails to escape into the cytosol at the early time point of infection. Therefore, bacterial escape into the cytosol is required, and is sufficient, for nuclear translocation of NF-κB in *F. tularensis*infected macrophages.

Some bacteria induce apoptosis in mammalian cells by activating specific components of the apoptotic pathways, including the activation of caspases. The cytosolic pathogens *F*. *tularensis* triggers activation of caspase-1 through the adaptors ASC or Ipaf [1,11]. In turn, caspase-1 activation leads to the release of two potent proinflammatory cytokines, IL-1 α and -18, and eventually leads to cell death [11]. Previous work has shown that processing of caspase-1 does not occur in macrophages infected with the *mglA* mutant of *F*. *tularensis* [12], which is defective in a global regulator involved in stress and metabolic responses as well as bacterial escape into the cytosol. Our data show that the *iglC* mutant is defective in triggering caspase-1, indicating that bacterial escape into the cytosol is sufficient and essential for activation of caspase-1.

Our data have shown that by 6 h post-infection, the wt strain-infected hMDMs exhibited caspase-3 activation, but the *iglC* mutant strain is defective in triggering caspase-3 activation. Therefore, bacterial escape into the cytosol is essential and sufficient for triggering caspase-3 activation. Despite robust caspase-3 activation, cell death is not apparent till late stages of the infection, concomitant with the termination of intracellular replication. Cell death is most likely mediated by multiple process including caspas-1, caspase-3, and caspases-independent processes.

Our studies show for the first time simultaneous modulation of caspase-1, caspase-3, and activation of NF- κ B, and the delicate balance between them to maintain viability of the infected cell. It is remarkable that both caspase-1 and caspase-3 are activated during initial stages of the infection, yet viability of the infected cell is largely maintained, which is correlated with simultaneous activation of the anti-apoptotic transcription factor NF- κ B.

Acknowledgments

YAK is supported by Public Health Service Awards R01AI43965 and R01AI069321 from NIH and by the commonwealth of Kentucky Research Challenge Trust Fund and by Ministry of Science Educatuion and Sports Republic of Croatia (Grant number: 062-0621273-0950).

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Fig. 1.

p65 nuclear translocation in *F. tularensis*-infected macrophages. (A) Representative confocal laser scanning microscopy images for nuclear translocation of the p65 subunit of NF- κ B. Monolayers of hMDMs were infected with live or formalin-killed (FK) wt-GFP, *iglC*-GFP, AA100-GFP at moi of 10 for 1 h. At several time points after infection, the cells were labelled for the p65 subunit of NF- κ B and nuclei. Uninfected monolayers and *E. coli* LPS-treated monolayers were included as negative and positive controls. Approximately 100 infected macrophages were analyzed from different coverslips. (B) Quantitative analysis of p65 nuclear translocation in *F. tularensis*-infected macrophages. The results are representative of three independent experiments and error bars represent standard deviations. The asterisks represent statistically significant difference.

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Fig. 2.

CAPE blocks NF- κ B activation human macrophages. (A) Representative confocal laser scanning microscopy images for nuclear translocation of the p65 subunit of NF- κ B in CAPE treated cells. Monolayers of hMDMs were treated with 20 µg ml⁻¹ of CAPE 30 min prior to infection. Cells were infected with the wt strain of *F. tularensis* subsp. *novicida* at a moi 10. The parental strain AA100 of *L. pneumophila* was used as positive control. Uninfected and *E. coli* LPS-treated (1 µg ml⁻¹ for 20 min) monolayers were included as a negative and positive control respectively. (B) Quantitative analysis of p65 nuclear translocation in *F. tularensis*-infected macrophages in CAPE treated cells. The results are representative of three independent experiments and error bars represent standard deviations. The asterisks represent statistically significant difference.





Fig. 3.

The protection from cell death correlates with nuclear translocation of NF- κ B in *F. tularensis*-infected cells. (A) Representative confocal laser scanning microscopy images of apoptosis in CAPE treated cells 6h post-infection. Monolayers of hMDMs were either untreated or treated with 20 µg ml⁻¹ of CAPE 30 min prior to infection. Cells were infected with the wt strain of *F. tularensis* subsp. *novicida* or with the *iglC* mutant strain at a moi 10. The parental strain AA100 of *L. pneumophila* was used as control. At 1, 6 and 24 h after infection, cell death was determined by TUNEL. (B) Quantitative analysis of cell death detected by TUNEL after CAPE treatment. Approximately 100 macrophages were analysed by confocal microscopy from different coverslips. The asterisks represent statistically significant difference.



Fig. 4.

F. tularensis induces caspase 1 activation in hMDMs. (A) Representative confocal laser scanning microscopy images of hMDMs infected with the wt strain or *iglC* mutant. The activation of caspase 1 (C1) and apoptosis (TUNEL) were determined at 1, 6 and 24 h post infection. (B) Quantitative analysis of C1 activation in C3 inhibited cells stained for cell death. Approximately 100 macrophages were analysed by confocal microscopy from different coverslips. The data are representative of three independent experiments and error bars represent standard deviations. The asterisks represent statistically significant difference.

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Fig. 5.

F. tularensis induces caspase 3 activation in the late stage of infection of hMDMs. (A) Representative confocal laser scanning microscopy images of hMDMs infected with the wt strain or the *iglC* mutant. The activation of caspase 3 (C3) and TUNEL were determined at 1, 6 and 24 h post-infection. (B) Quantitative analysis of C3 activation in C1 inhibited cells stained for cell death. Approximately 100 macrophages were analysed by confocal microscopy from different coverslips. The data are representative of three independent experiments and error bars represent standard deviations. The asterisks represent statistically significant difference.



Fig. 6.

Resistance to apoptosis-inducing agents. (A) Representative confocal microscopy images of hMDMs macrophages infected with either *F. tularensis subsp. novicida* wild type strain or *iglC* mutant strain. Staurosporin was used to induce artificially apoptosis of the macrophages. The cells were stained with both the TUNEL assay labelling (red) and the active Caspase-3 antibodies (blue). (B) Quantitative analysis of cell death in staurosporin treated cells. Approximately 100 macrophages were analysed by confocal microscopy from different coverslips. The asterisks represent statistically significant difference.