Francisella recognition by inflammasomes: differences between mice and men

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Pathogen recognition by intracellular sensors involves the assembly of a caspase-1 activation machine termed the inflammasome. Intracellular pathogens like Francisella that gain access to the cytosolic detection systems are useful tools to uncover the details of caspase-1 activation events. This review overviews Francisella function in the mononuclear phagocyte with particular attention to inflammasome versus pyroptosome roles and outlines differences between mouse and human caspase-1 activation pathways. Specific attention is placed on functional differences between human and murine pyrin as an intracellular recognition molecule for Francisella.

Keywords: AIM2, Francisella, IL-1β, inflammasome, M-CSF, monocytes, pyrin, pyroptosome

OVERVIEW

Understanding the innate host response to infectious challenges is a process greatly accelerated by the discovery of intracellular pathogen sensors and the inflammasome concept, reviewed extensively in this special edition. Much can be learned by focusing on the details of individual pathogen types, which can refine our understanding of innate host functions. In this mini-review, we will focus on the detection of Francisella and how this particular organism has allowed a clearer understanding of intracellular pathogen recognition. We will also briefly review Francisella infection and the peculiarities regarding detection in mouse and man.

Francisella tularensis is a zoonotic agent circulating in the Northern Hemisphere. It is transmitted from host to host by several pathways where arthropod vectors predominate (Ellis et al., 2002; Sjostedt, 2007). F. tularensis represents four subspecies differing in geographical location, infectivity, and virulence to the various hosts (Oyston, 2008). The most virulent species is F. tularensis subsp. tularensis (F. tularensis) with median lethal dose <10 CFU for humans, mice, and rabbits. However, F. tularensis subsp. holarctica (F. holarctica) has more broad virulence with calculated LD50 ranging from ~1 CFU for mice, <10^3 CFU for humans, and >10^3 CFU for rabbits. F. tularensis subsp. novicida (F. novicida) while still highly virulent in mice (LD50 < 10^3 CFU), shows low virulence for humans and rabbits (reviewed in Ellis et al., 2002).

Abbreviations: AIM2, absent in melanoma 2; ASC, apoptosis-associated speck-like protein containing a CARD; CARD, caspase recruitment domain; DAMP, danger-associated molecular pattern; NLR, NOD-like receptor (nucleotide-binding domain, leucine-rich repeat containing protein); NLRC, NOD-like receptor containing a CARD; NLRP, NOD-like receptor containing a PYD; PAMP, pathogen-associated molecular pattern; PRR, pattern recognition receptor; PYD, pyrin domain; RIG-I, retinoic acid inducible gene I.

The disease caused by F. tularensis is called tularemia, otherwise known as rabbit fever (Ellis et al., 2002). The characteristics of tularemia include fever, aches, and signs of toxicity lasting for several days (Parmely et al., 2009). If left untreated, tularemia may result in high mortality, reaching up to 60% with the pneumonic form of the disease (reviewed in Parmely et al., 2009).

Because Francisella has a low dose requirement, high virulence, and high morbidity, it is an attractive organism for bacteriological weapons (Sjostedt, 2007). Since F. tularensis was included in the list of select agents as a potential bioterrorist weapon (Ellis et al., 2002; Rotz et al., 2002), research into the pathogenicity of this bacterium in host organisms has been greatly accelerated. Large data sets have been generated regarding the specific details of bacteria–host interactions, albeit with some controversy. The controversy may be explained by differences in models used, including Francisella subspecies, host organisms, or cell types.

Francisella belongs to a select group of bacteria, including Listeria, Shigella, Burkholderia, and Rickettsia spp., which proliferate within the host cell cytosol (Ray et al., 2009). Francisella evolved to replicate in macrophages, which function to provide the first line of defense against pathogens. To survive in this biological niche, Francisella developed a strategy to avoid recognition and destruction by the immune system in order to utilize the host resources and multiply. The employment of this strategy by different F. tularensis subspecies dictates the differences in bacterial pathogenicity and evolutionary success.

Accordingly, a delay in recognition and immune system activation should be beneficial. F. tularensis (Schu4, type A) is the most virulent subspecies for both mice and humans, suppressing all pro-inflammatory responses for at least 72 h following infection (Kirimanjeswara et al., 2008). Since the Schu4 strain requires a BSL3 facility, more work has been done with the less virulent...
strains, *F. holarctica* (LVS, type B) and *F. novicida*. However, these two subspecies differ in their relative virulence for mice versus human – i.e., virulent in mice and attenuated in humans.

Could this difference in virulence be related to distinctions in recognition? Based on differences in virulence between *Francisella* subspecies and host organisms, it is reasonable to suggest that differences in *Francisella* recognition are based upon host response differences. Notably, virulent and attenuated *Francisella* subspecies have escape capacities that drive release from the phagosome in both humans and mice (Golovliov et al., 2003; Mariathasan et al., 2005; Gavrilin et al., 2006). Upon phagosomal escape *Francisella* may be also recognized by intracellular pathogen sensors, many of which belong to the NLR family of proteins. The focus of current review is to highlight *Francisella* recognition by different hosts.

**INFLAMMASOME VERSUS PYROPTOSOME**

**INFLAMMASOME**

*Francisella* recognition induces fever and cell death and is characterized by a wave of pro-inflammatory cytokines, where IL-1β plays a major role. This cytokine is induced upon mononuclear cell stimulation and synthesized as a biologically inactive proIL-1β (Dinarello, 1998). Its conversion to the biologically active 17 kDa form, which includes cleavage and release from cells, is tightly regulated by caspase-1 which, in turn, also requires an activation event (Yamin et al., 1996). Caspase-1 activation depends upon assembly of a multi-protein complex called the inflammasome (Martinon et al., 2002). All protein partners in the inflammasome possess either CARD (caspase recruitment domain) or PYD (pyrin domain) and assemble via CARD–CARD and PYD–PYD interactions (Martinon and Tschopp, 2006; Martinon et al., 2009). The prototypical inflammasome consists of CARD containing caspase-1, CARD, and PYD containing adaptor molecule ASC, and an NLR sensor of pathogen– or danger-associated molecule patterns (PAMPs or DAMPs). Depending on the presence of CARD or PYD domains, these sensors are subdivided as NLRC or NLRP family members, respectively (Ting et al., 2008). Because caspase-1 is the central protein/component of every inflammasome, while ASC is present in the majority of them, inflammasomes are named based on the participating pattern recognition receptor – PRR (NLR or other CARD or PYD containing protein). As such, several inflammasome structures have been described. These include the NLRP1 inflammasome (NLRP1, ASC, caspase-1, and caspase-5; Martinon et al., 2002), the NLRP3 inflammasome (NLRP3, CARD8, ASC, caspase-1; Agostini et al., 2004), the NLRC4 (IPAF) inflammasome (NLRC4, ASC, caspase-1; Mariathasan et al., 2004), and the NLRC5 inflammasome (NLRC5, ASC, caspase-1; Davis et al., 2010). However, it was proposed that several non-NLR proteins may also initiate the assembly of inflammasomes. For example, the RIG-I inflammasome (RIG-I, ASC, caspase-1; Poeck et al., 2010), the AIM2 inflammasome (AIM2, ASC, caspase-1; Burckstummer et al., 2009; Fernandes-Alnemri et al., 2009; Hornung et al., 2009; Roberts et al., 2009), and the pyrin inflammasome (pyrin, ASC, caspase-1; Yu et al., 2006; Gavrilin et al., 2009).

**PYROPTOSOME**

In addition to the widely accepted inflammasome concept, there is another caspase-1 activation platform, which may be distinct from the inflammasome, called a pyroptosome (Fernandes-Alnemri et al., 2007). The pyroptosome is a single (one per cell) 1–2 μm supra-molecular speck which contains ASC and caspase-1. This speck is different from the *in vitro* reconstituted inflammasome, which forms a ring-like structure with an outer diameter of ∼13 nm and an inner diameter of ∼4 nm (Faustin et al., 2007). Pyroptosome assembly is associated with host cell death that occurs within minutes of the ASC speck complex formation (Fernandes-Alnemri et al., 2007). This type of cell death is uniquely dependent on caspase-1 activation and, because of its ability to induce IL-1β dependent fever, named pyroptosis (Fink and Cookson, 2005). Host cell pyroptosis results in ASC pyroptosome aggregation, followed by cell rupture and the accompanied release of IL-1β (Fernandes-Alnemri et al., 2007). However, IL-1β and caspase-1 accumulation in cell culture media is often seen in the absence of cell death following LPS or bacterial infection of mononuclear phagocytes which suggests that inflammasomes may be distinct from pyroptosomes (Mangan et al., 1991; Gavrilin et al., 2009; Santic et al., 2010). In this regard, at least some of the IL-1β and caspase-1 (as well as ASC and some other PRR) are released from live monocytes in the form of microvesicles (MacKenzie et al., 2001; Sarkar et al., 2009). Since the size of exosomal microvesicles is 80–100 nm, these inflammasomes are likely distinct from the micrometer sized pyroptosome. Thus, pyroptosome formation accompanying pyroptosis may be a radically different way to eliminate a cytosolic bacterial burden compared to microvesicular IL-1β and the inflammasome release pathway. How host cell death is regulated by caspase-1 and how it may differ with respect to caspase-1 activation pathways is an area of particular interest for future studies. The summary of the inflammasome versus pyroptosome is shown in Table 1.

**MURINE RECOGNITION OF INTRACELLULAR FRANCISELLA**

**INTERFERON SIGNALING**

It is clear in mice that type I interferon signaling is necessary for inflammasome activation, IL-1β/IL-18 release, and the extensive cell death that follows *Francisella* infection (Henry et al., 2007). Murine macrophages showed 10-fold increase of Ifnb mRNA 8 h after infection with *F. novicida* (Henry et al., 2007). The 100-fold induction of the Ifnb gene was also observed after mouse macrophage infection with the LVS *Francisella* strain, but in contrast to Il1b gene expression which peaks at 4 h post-infection, induction of the Ifnb gene
was observed much later (starting at 12 h post-infection; Cole et al., 2008). In response to intracellular Francisella, mouse macrophages secrete IFN-β, which in an autocrine fashion activates caspase-1 and induces extensive cell death (Henry et al., 2007). In a paracrine fashion, IFN-β negatively regulates the number of IL-17A(+) gamma delta T cells and resistance to infection (Henry et al., 2010). However, while many researchers agree that IFN-β activates the inflammasome, there is a controversy surrounding its effect on the expression of some potential Francisella-sensing PRR. In particular, it was reported that Aim2 expression is either induced (Jones et al., 2010) or not affected (Alnemri, 2010; Fernandes-Alnemri et al., 2010) in murine macrophages by IFN-β treatment. Thus, IFN-β’s role in inflammasome function requires additional study.

**AIM2 AS FRANCISIELLA-SENSING PRR**

Knockout animals provide an extremely powerful tool to evaluate the effects of specific innate molecules on biological processes. In experiments with knockout mice, the importance of the caspase-1/ASC axis activation following Francisella escape into the cytosol was clearly shown (Mariathasan et al., 2005). It was also shown in mice Nlrp1, Nlrp3, and Nlrc4 do not recognize Francisella (Weiss et al., 2007; Huang et al., 2010). Recently, several groups showed that absent in melanoma 2 (Aim2), an interferon-inducible protein, is responsible for intracellular Francisella detection in murine macrophages (Fernandes-Alnemri et al., 2010; Jones et al., 2010; Rathinam et al., 2010). Aim2 belongs to the HIN-200 (PYHIN) protein family (Ludlow et al., 2005). Mice and humans share 55% amino acid identity of Aim2 (Chouby et al., 2010). Structurally, Aim2 has a HIN-200 amino acid repeat, which can bind double strand DNA, and a PYD domain, which is able to homo-dimerize with other PYD domains (Burckstummer et al., 2009; Chouby et al., 2010). The PYD domain of Aim2 may recruit the PYD of the inflammasome adaptor ASC, as shown by co-localization after dual co-expression (Burckstummer et al., 2009). This co-localization is functional, as over-expression of mouse Aim2 induces ASC-dependent activation of an NF-κB reporter gene (Rathinam et al., 2010) and promotes caspase-1 activation (Fernandes-Alnemri et al., 2009). When Aim2 is expressed in the cell, ASC oligomerization is induced by variety of factors, such as double stranded DNA (Burckstummer et al., 2009; Fernandes-Alnemri et al., 2009; Roberts et al., 2009). Thus, Aim2-ASC-caspase-1 may form an Aim2 inflammasome or pyroptosome, whereas bacterial DNA serves as the DAMP which induces this assembly. In support of this conclusion, it was recently shown that F. novicida and LVS strains activate murine caspase-1 in an Aim2 dependent manner (Fernandes-Alnemri et al., 2010; Jones et al., 2010; Rathinam et al., 2010). Caspase-1 activation, IL-1β secretion, and cell death were absent in Aim2-/- macrophages in response to F. novicida infection (Fernandes-Alnemri et al., 2010), while bacterial burden was increased (Jones et al., 2010). It is proposed that bacterial DNA could leak into the cytoplasm during escape from the phagosome as a result of Francisella lysis and phagosome acidification (Fernandes-Alnemri et al., 2010). Thus far, there is no consensus as to whether the Aim2-ASC speck represents the inflammasome or the pyroptosome. To avoid confusion, some researchers refer to the speck as an “ASC focus,” with the belief that assembly of this focus leads to secretion of mature IL-1β and cell death (Jones et al., 2010).

**INTRACELLULAR FRANCISIELLA RECOGNITION IN HUMAN PHAGOCYTES**

**INTERFERON SIGNALING AND AIM2 EXPRESSION**

Although Aim2 is clearly important in murine responses to Francisella, the process of Francisella recognition in human mononuclear cells is less clear. In sharp contrast to the mouse macrophages, infection of human monocytes with attenuated *F. novicida* does not lead to the up-regulation of type I interferon signaling (Butchar et al., 2008). However, virulent *F. tularensis* Schu4 induces some IFNβ expression (Butchar et al., 2008). At the same time, *F. novicida* induces significantly higher IL-1β release in human monocytes compared to Schu4 or LVS (Gavrilin et al., 2006; Butchar et al., 2008). Of note, we are unable to find detectable expression of AIM2 in primary human CD14+ monocytes by Western blot (unpublished results). However, it has been shown that human PBMC express detectable amounts of AIM2 (Meixenberger et al., 2010). Human AIM2 gene is expressed in the spleen, small intestine, and peripheral leukocytes (DeYoung et al., 1997). Because PBMC contain lymphocytes as well as CD14+ monocytes, lymphocyte contributions may explain this difference in AIM2 detection. Also, in agreement with our data are reports that human monocytes do not express AIM2 except after interferon induction (Hornung et al., 2009; Rathinam et al., 2010). However, the inability of *F. novicida* to induce IFN-β in human monocytes (Butchar et al., 2008), may explain the failure to detect human monocyte AIM2. Concomitantly, human monocytes release large amounts of mature IL-1β following infection with *F. novicida*, suggesting that human monocytes may utilize a PRR other than AIM2 for intracellular Francisella recognition (Gavrilin et al., 2009). Interestingly, Francisella-infected human CD14+ monocytes show high induction of interferon gamma gene (IFNG), i.e., type II interferon signaling (Butchar et al., 2008). This induction may represent a defense mechanism by which IFN-γ restricts cytosolic growth of Francisella without affecting its phagosomal escape (Edwards et al., 2010). Of note, it was shown that IFN-γ also may induce AIM2 expression in human monocytic cell line THP-1 (Fernandes-Alnemri et al., 2009) as well as pyrin in human CD14+ monocytes (Centola et al., 2000).

**PECULIARITIES OF PYRIN FUNCTION IN HUMAN VERSUS MURINE CELLS**

Pyrin, the familial Mediterranean fever (FMF) protein, belongs to the tripartite motif (TRIM) family of proteins, all of which share structural homology (Nisole et al., 2005). Although pyrin has an amino terminal PYD, it is not a member of NLR family of intracellular sensors of pathogens. Nonetheless, pyrin has been shown to interact via its PYD with the adaptor protein ASC (Richards et al., 2001). Importantly, the CARD of the ASC molecule can interact directly with caspase-1 via the amino terminal CARD. As a result, these proteins could assemble into an inflammasome. We observed that pyrin, ASC, and caspase-1, tagged to various fluorescent proteins and expressed in HEK 293 cells, co-localize into one speck (unpublished data), which is similar to the AIM2 pyroptosome discussed earlier.

The hypothetical model of pyrin inflammasome with its C-terminal B30.2 domain as a pathogen sensor was first proposed by Alnemri and colleagues (Yu et al., 2006). Importantly, this lab showed that the *Mefv* knockout animals still respond to *F. novicida*, whereas Aim2-/- animals do not (Fernandes-Alnemri et al., 2008).
et al., 2010). This implies that Aim2, but not pyrin, is responsible for *Francisella*-induced caspase-1 activation and IL-1β release in murine macrophages.

Our work suggests that the human response to *Francisella* may be quite distinct from the murine response. For example, pyrin is an important component of caspase-1 activation and IL-1β release after challenging human mononuclear cells with *F. novicida* (Gavrilin et al., 2009). At first glance, this result is difficult to reconcile with the *Mefv* murine knockout results (Fernandes-Alnemri et al., 2010). However, it is important to highlight an important difference between pyrin protein encoded by *MEFV* in human and mice (Figure 1). Although these two species share 47.6% identity and 65.5% similarity in pyrin amino acid sequence, murine pyrin lacks the complete B30.2 domain, a potential pathogen sensing region (Chae et al., 2000). Importantly, mutations responsible for FMF in humans are mostly located in the B30.2 domain (Ting and Davis, 2005). Thus, it is possible that mice have lost their ability to detect *Francisella* via pyrin due to genetic changes in murine pyrin gene.

Pyrin's role in caspase-1 function has also been a subject of controversy. Discovered as a gene mutation that leads to bouts of inflammation, several groups immediately suggested an anti-inflammatory role for pyrin by showing that pyrin can inhibit IL-1β activation (Chae et al., 2003, 2006; Papin et al., 2007). In contrast, other groups provide evidence that pyrin is a pro-inflammatory protein, activating caspase-1 dependent IL-1β cleavage (Yu et al., 2006, 2007; Seshadri et al., 2007; Gavrilin et al., 2009). Working with human primary CD14+ monocytes and their derived macrophages, we observed a correlation between pyrin levels and the ability of the cell to process and release IL-1β in response to *Francisella* infection. Fresh human monocytes show high levels of pyrin and readily release large quantities of IL-1β, whereas macrophages are relatively deficient in pyrin and defective in processing, but not in the synthesis of proIL-1β. Alveolar macrophages also follow this pattern and do not readily process and release IL-1β (Wewers and Herzyk, 1989). In addition, the modulation of intracellular pyrin levels in mononuclear phagocytes in various stages of differentiation correlates with the ability of pyrin to affect IL-1β release in response to *Francisella* infection (Gavrilin et al., 2006, 2009; Seshadri et al., 2007).

**M-CSF**

Particularly noticeable is the striking effect of M-CSF on pyrin levels in human cells. Macrophages matured in the presence of M-CSF maintain pyrin levels and readily release IL-1β followed by infection with *Francisella*, similar to freshly harvested CD14+ human monocytes. Moreover, macrophages matured in the absence of growth factors, while defective in proIL-1β processing and release, re-induce pyrin and the ability to process IL-1β with M-CSF treatments (Gavrilin et al., 2009). Thus, the local microenvironment may modulate a broad-range of macrophage phenotypes that extend well beyond M1/M2 classification models.

**THP-1 CELL LINE AS A MODEL OF HOST–PATHOGEN INTERACTION**

In discussing the differences in AIM2 and pyrin expression and function between mice and primary human mononuclear cells, it is important to mention the human pre-monocytic cell line THP-1 because a plethora of experiments have been performed with this cell line. THP-1 cells are phagocytic, can be infected with *Francisella*, and show activation of caspase-1 (Mohapatra et al., 2007; Zakharova et al., 2010). It was shown that stimulation of THP-1 cells with PMA, Sendai virus, or INF-γ induces AIM2 expression (Burckstummer et al., 2009; Fernandes-Alnemri et al., 2009; Hornung et al., 2009). PMA treated THP-1 cells also release IL-1β after infection with *F. novicida*. During THP-1 differentiation to THP-derived macrophages (TDM), pyrin disappears as is also observed with CD14+ monocyte differentiation to monocyte-derived macrophages (MDM). Consistent with a role for pyrin in caspase-1 activation after *F. novicida* infection, TDMs transduced to express pyrin release about sixfold more IL-1β upon infection with *F. novicida* as compared to TDM controls (Gavrilin et al., 2009). Nevertheless, even in the absence of pyrin, TDM release more IL-1β than do fresh THP-1 cells, suggesting the existence of a non-pyrin activation mechanism as well. Thus, it is interesting to speculate that pyrin and AIM2 may both be sensors of *Francisella* and that these two sensors may work in concert with each other. This speculation is more important in human cells with fully intact pyrin B30.2 domains.

In keeping with the pyrin/AIM2 dual detection hypothesis, it is notable that each PRR may recognize certain DAMP or PAMP covering discreet molecular patterns, such that overlapping regions of recognition may lead to more precise detection of pathogen types. The example of such composite detection is shown for *Listeria*, where mouse Nlrc4, Nlrp3, and Aim2 work in cooperation (Tsuchiya et al., 2010; Warren et al., 2010; Wu et al., 2010) while human PBMC appear to detect *Listeria* only via NLRP3 and not in cooperation with AIM2, NLRC4, and other tested NLRs (Meixenberger et al., 2010). The newly described NLRC5 inflammasome also demonstrated cooperativity with NLRP3 when THP-1 cells were stimulated with panel of PAMPs and DAMPs, typically activating NLRP3 inflammasome (Davis et al., 2010).

Regarding *Francisella* detection, we cannot rule out the possibility that AIM2 expression in human monocytes exists below the limits of confident detection, but still in amounts sufficient to trigger ASC oligomerization and caspase-1 activation upon DNA recognition. Apparently, NLRs may be able to act at low concentrations relative to ASC and caspase-1, triggering an initial node of activation. For example, NLRP3 initiates ASC pyroptosome oligomerization at the molar ratio of 1:500 (Fernandes-Alnemri et al., 2007). Following

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**FIGURE 1 | Differences between pyrin in human and mice.**

- **Human Pyrin 781 aa**
  - PYD
  - BB
  - CC
  - PRY
  - SPRY

- **Mouse Pyrin 767 aa**
  - PYD
  - BB
  - CC

**Human Pyrin 781 aa**

- PYD
- BB
- CC
- PRY
- SPRY

**Mouse Pyrin 767 aa**

- PYD
- BB
- CC

**PRY domain, associated with SPRY domain**

**SPRY domain**
initial dimerization, ASC oligomerization may become independent of NLR requirements, having reached a critical threshold concentration of pyroptosome formation and function.

CONCLUSION

In summary, the study of specific pathogens continues to uncover particulars of inflammasome regulation. In the case of Francisella, human mononuclear cells recognize Francisella before its internalization resulting in activation of multiple signaling pathways, including proIL-1β synthesis. Thus intracellular Francisella, after escape from the phagosome, may be detected by several pattern recognition receptors, where AIM2 and pyrin may trigger formation of the pyroptosome or the inflammasome platform for caspase-1 activation. This event results in the initiation of cell death and/or IL-1β processing and release. However, comparisons between mice and humans reveal that there are important evolutionary changes in pathogen sensing that are likely to have been driven by environmental pressures. Understanding the peculiarities of pathogen-host sensing is dependent upon powerful animal models but must continue to be interpreted alongside human studies.

ACKNOWLEDGMENTS

This work was supported by the NIH/NIAID Regional Center of Excellence for Bio-defense and Emerging Infectious Diseases Research (RCE) Program U54-AI-057153, by NIH HL089440 and HL76278.

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Conflict of Interest Statement: The authors declare that the research was con-
ducted in the absence of any commercial or
financial relationships that could be con-
strued as a potential conflict of interest.

Received: 16 November 2010; paper pend-
ing published: 28 December 2010; accepted:
19 January 2011; published online: 09
February 2011.

Citation: Gavrilin MA and Wewers MD
(2011) Francisella recognition by inflam-
asomes: differences between mice and
men. Front. Microbiol. 2:11. doi: 10.3389/
fnmic.2011.00011
This article was submitted to Frontiers in
Cellular and Infection Microbiology, a spe-
cialty of Frontiers in Microbiology.

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