Shigella Type III Secretion Protein MxiI Is Recognized by Naip2 to Induce Nlrc4 Inflammasome Activation Independently of Pkc\(\delta\)

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Abstract

Recognition of intracellular pathogenic bacteria by members of the nucleotide-binding domain and leucine-rich repeat containing (NLR) family triggers immune responses against bacterial infection. A major response induced by several Gram-negative bacteria is the activation of caspase-1 via the Nlrc4 inflammasome. Upon activation, caspase-1 regulates the processing of pro-IL-1\(\beta\) and pro-IL-18 leading to the release of mature IL-1\(\beta\) and IL-18, and induction of pyroptosis. The activation of the Nlrc4 inflammasome requires the presence of an intact type III or IV secretion system that mediates the translocation of small amounts of flagellin or PrgJ-like rod proteins into the host cytosol to induce Nlrc4 activation. Using the Salmonella system, it was shown that Naip2 and Naip5 link flagellin and the rod protein PrgJ, respectively, to Nlrc4. Furthermore, phosphorylation of Nlrc4 at Ser533 by Pkc\(\delta\) was found to be critical for the activation of the Nlrc4 inflammasome. Here, we show that Naip2 recognizes the Shigella T3SS inner rod protein MxiI and induces Nlrc4 inflammasome activation. The expression of MxiI in primary macrophages was sufficient to induce pyroptosis and IL-1\(\beta\) release, which were prevented in macrophages deficient in Nlrc4. In the presence of MxiI or Shigella infection, MxiI associated with Naip2, and Naip2 interacted with Nlrc4. siRNA-mediated knockdown of Naip2, but not Naip5, inhibited Shigella-induced caspase-1 activation, IL-1\(\beta\) maturation and Asc pyroptosome formation. Notably, the Pkc\(\delta\) kinase was dispensable for caspase-1 activation and secretion of IL-1\(\beta\) induced by Shigella or Salmonella infection. These results indicate that activation of caspase-1 by Shigella is triggered by the rod protein MxiI that interacts with Naip2 to induce activation of the Nlrc4 inflammasome independently of the Pkc\(\delta\) kinase.

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Introduction

Recognition of intracellular pathogenic bacteria by members of the nucleotide-binding domain and leucine-rich repeat containing (NLR) family triggers immune responses against bacterial infection [1,2]. A major response against several pathogenic Gram-negative bacteria, including Salmonella, Legionella, and Shigella is the activation of caspase-1 via Nlrc4 in macrophages [1,3]. Upon bacterial stimulation, Nlrc4 mediates the formation of a multi-protein complex termed the inflammasome that induces the activation of caspase-1 leading to the proteolytic maturation of pro-IL-1\(\beta\) and pro-IL-18 as well as the induction of pyroptotic cell death in macrophages [4–6]. Many Gram-negative bacteria encode a type III secretion system (T3SS) with conserved structural features that promote virulence by injecting bacterial effector proteins directly into the cytosol of host cells [7,8]. In macrophages infected with Salmonella, the cytosolic delivery of flagellin or the bacterial rod protein PrgJ through the T3SS is recognized by Nlrc4 leading to inflammasome activation [9]. Recently, Naips (NLR family, apoptosis inhibitory proteins) have been shown to act as adaptor molecules that connect flagellin or the bacterial rod protein PrgJ to Nlrc4 [10,11]. Specifically, Naip5 and Naip6 associate with flagellin to promote Nlrc4 oligomerization and inflammasome activation, whereas Naip2 links PrgJ to Nlrc4 [10–12]. These findings suggest a model in which certain Naips specifically recognize flagellin or PrgJ to mediate Nlrc4 inflammasome activation. Recent studies, however, have revealed...
that the activation of Nlrc4 is more complex in that phosphorylation of Nlrc4 at Ser533 was found to be critical for the activation of the inflammasome [13]. Furthermore, it was suggested that Pkcδ is the major Nlrc4 kinase responsible for Nlrc4 phosphorylation and inflammasome activation [13].

Shigella are non-flagellated bacterial pathogens that contain highly evolved invasion systems that enable them to invade host cells and colonize the epithelium of the large intestine, which ultimately leads to a severe form of colitis called bacillary dysentery [14]. After uptake of Shigella by intestinal macrophages, the bacterium delivers a subset of effector proteins via the T3SS apparatus into the host cytosol [7,8,15]. The inner rod of the T3SS needle complex forms a conduit for protein transport through the periplasm which is assembled by the polymerization of PrgJ in Salmonella and its homologue MxiI in Shigella [16,17]. Because of the homology of Salmonella PrgJ with Shigella MxiI, it can be predicted that Shigella induces activation of Nlrc4 via the sensing of MxiI by host macrophages. Consistent with this notion, the T3SS of Shigella is required to induce IL-1β secretion and pyroptosis via the Nlrc4 inflammasome [18]. Furthermore, ectopic expression of MxiI reduced the viability of macrophages and this was inhibited in the absence of Nlrc4 [9]. However, the mechanism by which Shigella MxiI induces activation of the Nlrc4

**Figure 1.** Expression of *Shigella* rod protein MxiI induces activation of the Nlrc4 inflammasome in macrophages. WT or Nlrc4−/− BMDMs were nucleofected with MSCV-IRES-GFP (GFP) or MSCV-IRES-GFP encoding *Shigella* MxiI (MxiI-GFP). After 20 hrs, the percentage of GFP-positive viable cells in the total cell population was analyzed by fluorescence microscopy (A) and the production of IL-1β in cell free supernatants by ELISA (B). * p<0.0001. (A and B) Results represent mean ± SD and are representative of three independent experiments.
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**Author Summary**

*Shigella* are bacterial pathogens that are the cause of bacillary dysentery. An important feature of *Shigella* is their ability to invade the cytoplasm of host epithelial cells and macrophages. A major component of host recognition of *Shigella* invasion is the activation of the inflammasome, a molecular platform that drives the activation of caspase-1 in macrophages. Although *Shigella* is known to induce the activation of the Nlrc4 inflammasome, the mechanism by which the bacterium activates Nlrc4 is largely unknown. We discovered that the *Shigella* T3SS inner rod protein MxiI induces Nlrc4 inflammasome activation through the interaction with host Naip2, which promoted the association of Naip2 with Nlrc4 in macrophages. Expression of MxiI induced caspase-1 activation, Asc oligomerization, pyroptosis and IL-1β release which required Naip2, but not Naip5. Significantly, caspase-1 activation induced by *Shigella* infection was unaffected by deficiency of the Pkcδ kinase. This study elucidates the microbial-host interactions that drive the activation of the Nlrc4 inflammasome in *Shigella*-infected macrophages.
inflammasome remains unknown. In this study, we provide evidence that MxiI mediates the activation of the Nlrc4 inflammasome through interactions with Naip2. Furthermore, we demonstrate that Naip2, but not Naip5, is critical for the interaction of MxiI with Nlrc4 and the activation of the inflammasome in macrophages infected with *Shigella*. Finally, we show that Pkcδ is dispensable for Nlrc4 activation.

**Results**

**Expression of *Shigella* rod protein MxiI induces activation of the Nlrc4 inflammasome**

In the case of flagellated pathogenic bacteria, flagellin is a major and potent stimulator of the Nlrc4 inflammasome. In addition, *Salmonella* T3SS rod protein PrgJ is sensed by Nlrc4 to activate caspase-1. Because *Shigella* are unflagellated bacteria, we hypothesized that the *Shigella* T3SS rod protein MxiI, a homologue of *Salmonella* PrgJ, induces the activation of the Nlrc4 inflammasome. To test this hypothesis, we expressed MxiI in wild-type (WT) and Nlrc4-deficient bone marrow-derived macrophages (BMDM) using a MCV-IRES-GFP retroviral vector and assessed cell viability by the numbers of viable green fluorescence protein (GFP)-positive cells. After overnight culture, the viability of WT macrophages was dramatically decreased by MxiI-GFP expression when compared to expression of GFP (Figure 1A). Importantly, the decrease in cell viability was inhibited in Nlrc4−/− macrophages (Figure 1A). Consistently, expression of MxiI-GFP, but not GFP, induced the release of IL-1β in WT macrophages, which was abolished in macrophages lacking Nlrc4 (Figure 1B). These results indicate that expression of MxiI induces the activation of the Nlrc4 inflammasome.

**Shigella MxiI interacts with Naip2 and promotes the interaction of Naip2 with Nlrc4**

We next tested whether the rod protein MxiI interacts with Naip2 or Naip5 in macrophages. Because expression of MxiI in macrophages causes cell death (Figure 1A), we used macrophages from caspase-1-deficient mice to assess the interaction of MxiI with Naip proteins by immunoprecipitation. In these experiments, we expressed T7-tagged MxiI in the presence of HA-tagged Naip2, HA-tagged Naip3 or control plasmid. Immunoprecipitation analysis showed that MxiI associated with Naip2, but much less with Naip5 as revealed by immunoblotting with anti-HA antibody (Figure 2A). Next, we investigated the interaction between Nlrc4 and Naip2 in *Shigella*-infected macrophages. To assess this, we expressed T7-tagged Nlrc4 and HA-tagged Naip2 or Naip5, or control empty vector in uninfected or caspase-1-deficient macrophages infected with WT or an isogenic vector in uninfected or caspase-1-deficient macrophages infected with WT or T3SS-deficient *Shigella* for 3 hrs and inflammasome activation was analyzed by immunoblotting with an antibody specific for mature IL-1β p17. In the absence of exogenous Naip2 or Naip5, infection with WT *Shigella* enhanced the processing of pro-IL-1β into IL-1β p17 (Figure S1A). The formation of IL-1β p17 was further enhanced by Naip2, but not Naip5 in *Shigella*-infected cells (Figure S1A). In this reconstitution system, the enhancement of IL-1β p17 formation by Naip2 in cells infected with WT *Shigella* required Nlrc4, Asc and caspase-1 (Figure S1B).

**Naip2 promotes the processing of IL-1β in an inflammasome reconstitution system**

We next performed additional studies to verify that *Shigella* infection promoted the activation of Nlrc4 via Naip2. To confirm the preferential effect of Naip2 on Nlrc4 activation, we performed reconstitution experiments by expressing Nlrc4, Asc, caspase-1, pro-IL-1β and Naip2 or Naip5 in 293T cells. One day after transfection, cells were infected with WT or T3SS-deficient *Shigella* for 6 hrs and inflammasome activation was analyzed by immunoblotting with an antibody specific for mature IL-1β p17. In the absence of exogenous Naip2 or Naip5, infection with WT *Shigella* induced processing of pro-IL-1β into IL-1β p17 (Figure S1A). The formation of IL-1β p17 was further enhanced by Naip2, but not Naip5 in *Shigella*-infected cells (Figure S1A). In this reconstitution system, the enhancement of IL-1β p17 formation by Naip2 in infected with WT *Shigella* required Nlrc4, Asc and caspase-1 (Figure S1B).

**Shigella induces caspase-1 activation via Nlrc4, Asc and Naip2 in macrophages**

*Shigella* infection stimulates Nlrc4- and Asc-dependent inflammasome activation in macrophages [18]. However, *Shigella* was also shown to induce macrophage cell death via Nlrp3 after 2–6 hrs of infection at a bacteria/macrophage ratio of 50:1 [24]. To verify these seemingly contradictory results, we reassessed the role of Asc, Nlrc4 and Nlrp3 in *Shigella*-induced caspase-1 activation. In these experiments, LPS-primed BMDM were infected with the *Shigella* WT or S325 (T3SS-deficient mutant) at a bacteria/macrophage ratio of 10:1 for 30 min. As expected, WT, but not mutant *Shigella*, induced processing of procaspase-1 into the p20 subunit of caspase-1 (Figure S2A). The inability of the mutant bacterium to activate caspase-1 could not be explained by reduced uptake by macrophages (Figure S3). Importantly, caspase-1 activation, IL-1β release, and pyroptosis required Nlrc4 and Asc, but not Nlrp3 (Figure S2A-C). Because previous studies showed that Asc was not required for pyroptosis induced by *Shigella* in BMDM differentiated for 5 days [18], we assessed cell death induced by *Shigella* in BMDM differentiated for 3, 4 and 5 days in culture (Figure S4). Consistent with previous studies [18], Asc was not required for pyroptosis in macrophages differentiated for 5 days (Figure S4). In macrophages differentiated for 3 or 4 days, however, cell death induced by *Shigella* was enhanced in WT macrophages and impaired in Asc-deficient macrophages (Figure S2C and S4) which is in line with the results presented in Figure S2C.

Next, we investigated the role of Naip2 and Naip5 in caspase-1 activation induced by *Shigella*. We used siRNA-mediated knockdown to reduce the expression of Naip2 and Naip5 in macrophages (Figure 3A). Notably, caspase-1 activation induced by *Shigella* was attenuated by inhibiting the expression of Naip2, but not Naip5 (Figure 3B). Importantly, the ability of individual siRNA to inhibit caspase-1 activation correlated with reduction of Naip2 expression (Figure 3A, B). In addition, knockdown of Naip2, but not Naip5, reduced the release of IL-1β and IL-18 induced by *Shigella* infection at 1 or 2 hrs post-infection (Figure 3B, C). In control experiments, knockdown of Naip2 did not affect the production of IL-6 or CXCL2 in macrophages infected with WT or S325 mutant *Shigella* (Figure 3C). These results suggest that *Shigella* induces Nlrc4-dependent inflammasome activation via Naip2 in macrophages.
Figure 2. *Shigella* MxiI interacts with Naip2 and promotes the interaction of Naip2 with Nlrc4. (A) T7-tagged MxiI was co-expressed with HA-tagged Naip2 or Naip5 or control empty vector in caspase-1-deficient BMDMs. Cell lysates were immunoprecipitated with anti-T7 antibody and the interaction between MxiI and Naip2/5 was analyzed by immunoblotting with anti-HA antibody. (B) T7-tagged Nlrc4 or empty vector was co-expressed with HA-tagged Naip2, Naip5, or empty vector in caspase-1-deficient BMDMs. After 16 hrs cells were infected with *Shigella* WT or S325 mutant for 2 hr at a bacteria/macrophage ratio of 10:1. Cell lysates were immunoprecipitated with anti-T7 beads and the interaction between Nlrc4 and Naip2/5 was analyzed by immunoblotting with anti-HA antibody. (C) T7-tagged MxiI or empty vector was co-expressed with HA-tagged Naip2. Cell lysates were immunoprecipitated with anti-HA beads and the interaction of Naip2 with MxiI and endogenous Nlrc4 was analyzed by immunoblotting with anti-T7 or anti-Nlrc4 antibody. (A–C) Results are representative of three independent experiments.

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Figure 3. *Shigella* induces caspase-1 activation via Nlrc4, Asc and Naip2 in macrophages. (A–C). BMDM were nucleofected with siRNA targeting Naip2 or Naip5 for 48 hrs and then infected, or not, with *Shigella* WT or S325 mutant for additional 1–3 hrs at a bacteria/cell ratio of 10:1. (A) Knockdown efficiency was evaluated by RT-PCR after 48 hrs in uninfected cells. (B and C) The activation of caspase-1 was evaluated by detecting cleaved caspase-1 (p20) by immunoblotting, and the level of casp1 p20 was further quantified by densitometry (B). Cytokines production was analyzed by ELISA in cell free supernatants (B–C). * p<0.05, ** p<0.001; NS, not significant. (A–C) Results represent mean ± SD and are representative of three independent experiments.

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**Shigella or MxiI expression induces Naip2-dependent Asc pyroptosome formation in macrophages**

The Asc pyroptosome is a molecular platform that is thought to be important for the recruitment and activation of caspase-1 [25–27]. Infection of macrophages with WT, but not T3SS-deficient, *Shigella* induced the formation of the Asc pyroptosome which was detected in the cell cytoplasm by staining with an antibody that recognizes Asc (Figure 4A, B). The Asc pyroptosome induced by *Shigella* infection co-localized with FLICA staining that labels activated caspase-1 (Figure 4A). Importantly, knockdown of Naip2 by siRNA reduced Asc pyroptosome formation whereas Naip5 did not (Figure 4C, D). To provide direct biochemical evidence that the Asc pyroptosome is formed, we cross-linked the insoluble Asc protein complexes from *Shigella* or *Salmonella* infected macrophages and subjected them to immunoblotting with anti-Asc antibody. Immunoblotting analysis revealed that infection with WT *Shigella* or *Salmonella* induces prominent Asc dimer formation in WT, but not Asc-deficient macrophages (Figure 5A, upper panel). The induction of Asc dimers correlated with IL-1β release in culture supernatants (Figure 5A, lower panel). In contrast, Shigella deficient in T3SS and the fliC-deficient *Salmonella* mutant were impaired in the induction of Asc dimer formation (Figure 5A). Notably, expression of MxiI was sufficient to induce the formation of Asc dimers in caspase-1-deficient macrophages in the absence of *Shigella* infection (Figure 5B). Furthermore, knockdown of Naip2 by siRNA, but not Naip5, inhibited Asc dimer formation (Figure 5C). These results indicate that *Shigella* MxiI and Naip2 are important in Asc pyroptosome formation which is associated with inflammasome activation.

**Discussion**

The intracellular sensing of flagellin is the major trigger for the activation of the Nlr4 inflammasome in macrophages infected with *Salmonella* [4]. Because *Shigella* is non-flagellated, the current studies were aimed at understanding the mechanism by which *Shigella* induces the activation of Nlr4 in macrophages. We show here that *Shigella* induces the activation of the Nlr4 inflammasome through MxiI, an inner rod protein of the T3SS. MxiI associated with Naip2 and was sufficient to induce Nlr4-dependent IL-1β secretion and the interaction with Nlr4. Importantly, inhibition of Naip2 expression impaired the activation of the Nlr4 inflammasome and IL-1β/IL-18 release in *Shigella*-infected macrophages. Because IL-1β secretion induced by *Shigella* was not abolished by Naip2 knockdown, it is possible that *Shigella* also activates another inflammasome pathway that is minor and only unmasked by the inhibition of the Naip2-Nlr4 pathway. Alternatively, it is possible that the partial inhibition of IL-1β secretion reflects residual Naip2 protein expression in macrophages.

Our work is consistent with a model in which the T3SS inner rod proteins including PrgJ in *Salmonella* and MxiI in *Shigella* are recognized by Naip2 and this interaction leads to the recruitment and activation of Nlr4. Consistent with this model, we show that expression of MxiI promotes the association of Naip2 with Nlr4 and induces the oligomerization of Asc in macrophages. Furthermore, WT, but not T3SS-deficient *Shigella*, enhances the association of Naip2 and Nlr4 in macrophages. The failure of mutant *Shigella* to induce the interaction between Naip2 and Nlr4 is presumably explained by the inability of the T3SS mutant to release MxiI into the host cytosol. A measure of inflammasome activation is the formation of Asc oligomers [25–27]. Importantly, Asc oligomerization induced by MxiI was observed in caspase-1-deficient macrophages, indicating that this critical event is not a secondary event of caspase-1 activation. MxiI is composed of 97 amino acids and is predicted to be a soluble protein using publically available tools ([http://www.psort.org/psortb](http://www.psort.org/psortb)). It has been shown that MxiI is secreted into the culture medium by *Shigella* in a T3SS dependent manner [19–21]. Thus, as it was suggested for *Salmonella* PrgJ [9,22], we propose that small amounts of MxiI are leaked into the host cytosol via the T3SS during *Shigella* infection to induce the activation of Nlr4.

Recent studies showed that Nlr4 phosphorylation by Pkcδ is critical for inflammasome activation induced by *Salmonella* infection [13]. Thus, we assessed whether inflammasome activation caused by *Shigella* infection also requires Pkcδ. In these experiments, LPS-primed BMDM from WT and Pkcδ-deficient mice were infected with WT or S325 (T3SS-deficient mutant) *Shigella*, and IL-1β release was evaluated at different time points and bacterial/macrophage ratios after infection. As expected, expression of Pkcδ was induced by LPS stimulation in WT, but not Pkcδ-deficient macrophages (Figure 6A). Importantly, Pkcδ was not required for IL-1β secretion induced by *Shigella* or *Salmonella* (Figure 6B–D). In fact, Pkcδ deficiency enhanced IL-1β secretion in response to *Shigella* and *Salmonella* infection (Figure 6B–D). Furthermore, Pkcδ-deficient macrophages produced higher amounts of IL-1α and CCL5, but not CXCL2 than WT macrophages in response to infection (Figure 6D). The increased production of cytokines in Pkcδ-deficient macrophages was not associated with enhanced NF-κB or MAPK activation after *Shigella* infection (Figure 5S). Notably, induction of apoptosis in *Shigella*-infected macrophages was inhibited in macrophages deficient in Pkcδ (Figure 5S). Furthermore, treatment with z-DEVD-fmk, a cell permeable caspase-3 inhibitor, increased the production of IL-1β in WT macrophages infected with *Shigella* (Figure 5S), suggesting that increased production of IL-1β in Pkcδ-deficient macrophages is mediated, at least in part, by inhibition of apoptosis in *Shigella*-infected macrophages. Importantly, caspase-1 activation induced by *Shigella* or *Salmonella* was unimpaired in macrophages deficient in Pkcδ (Figure 6E), whereas it was abolished in macrophages deficient in Nlr4 (Figure 6E). These results indicate that Pkcδ is not essential for inflammasome activation induced by *Shigella* or *Salmonella* infection.
Figure 4. *Shigella* induces Naip2-dependent Asc pyroptosome formation in macrophages. WT (A–D), Asc−/− (A–D), Naip2-deficient (siRNA) or Naip5-deficient (siRNA) (C) BMDM were infected with *Shigella* WT or S325 mutant for up to 90 min (A–D). Cells were fixed and analyzed by confocal microscopy (A–D) and the percentage of cells containing Asc pyroptosomes was evaluated (D). Caspase-1 activation was detected using FLICA reagent (A) (green), Asc localization with anti-Asc antibody (red in A, green in B and C) and nuclei with DAPI (A) (blue). Arrows denotes Asc pyroptosomes. * p<0.01. Results represent mean ± SD and are representative of three independent experiments. doi:10.1371/journal.ppat.1003926.g004
that caspase-1 activation induced by *Shigella* or *Salmonella* infection is not impaired in Pkcδ-deficient macrophages. We do not have a clear explanation for the difference in results between our studies and previous results by Qu et al. These authors showed that in addition to Pkcδ, Pak2 was capable of phosphorylating Nlrc4 at the critical Ser533, although the results suggested that Pak2 was a minor Nlrc4-phosphorylating kinase [13]. Thus, it is conceivable that the difference in results could be explained by kinase redundancy and subtle variation in the expression of Nlrc4-phosphorylating kinases in different macrophage preparations. Regardless of the explanation, findings within this investigation clearly show that Pkcδ is dispensable for Nlrc4 activation. Thus, our results challenge the notion that Pkcδ is critical for inflammasome activation and indicate that further work is needed to understand the mechanism and role of Nlrc4 phosphorylation in inflammasome activation.

*Shigella* MxiI associates with Naip2 to induce the interaction of Naip2 with Nlrc4, which presumably leads to Nlrc4 oligomerization and inflammasome activation. In the *Salmonella* system, cytosolic flagellin binds to Naip5 and induces the association of Naip5 with Nlrc4 [10–12]. Reconstitution experiments with purified flagellin, Naip5 and Nlrc4 revealed that these components are sufficient to induce the formation of a disk-like complex composed of 11 or 12 proteins including Nlrc4 and Naip5, although the exact ratio of Naip5 and Nlrc4 in the complex remains unclear [12]. Based on the latter observations, we suggest that *Shigella* MxiI induces the oligomerization of Nlrc4 via their interaction with Naip2. Consistent with this model, we found that MxiI induced the interaction of Naip2 with Nlrc4 and the oligomerization of Asc. Furthermore, Naip2, but not Naip5, was critical for caspase-1 activation, pyroptosome formation, Asc oligomerization and IL-1β secretion. Collectively, these results
Figure 6. Pkcδ is not required for inflammasome activation caused by Shigella infection. (A) BMDMs from WT and Prkcd−/− mice were stimulated with LPS and the expression of Pkcδ was evaluated by immunoblotting. (B–D) BMDMs from WT and Prkcd−/− mice were infected with Shigella WT or S325 Shigella mutant (B–E) or Salmonella (C and E) at a bacteria/macrophage ratio of 10:1 for various time points (B) or at the indicated bacteria/macrophage ratio (C) for 1 hr, or with indicated bacteria/macrophage ratios for 2 hrs (D) or 30 min (E). The production of cytokines in cell free supernatant was analyzed by ELISA (B–D) and the activation of caspase-1 was evaluated by detecting cleaved caspase-1 (p20) by immunoblotting (E). *p<0.02. Results represent mean ± SD. Results are representative of at least three independent experiments.

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support a model in which distinct Naip family members act as sensors of flagellin and T3SS inner rod proteins and oligomerized Nlrc4 provides a platform for the recruitment and activation of caspase-1. While Naip2 knockdown reduced inflammasome activation, Naip5 knockdown had the opposite effect in response to Shigella infection. Although further work is needed to understand the role of Naip5, one possibility is that there is competition between Naip2 and Naip5 protein complexes and inhibition of Naip5 enhances the Naip2-Nlrc4 inflammasome pathway. Nlrc4 and caspase-1 contain CARD domains and they could interact directly via homotypic CARD-CARD interactions. However, the adaptor Asc is essential for the activation of caspase-1 in response to Salmonella and Shigella [18,33]. These results suggest that Asc is somehow required for the interaction between Nlrc4 and caspase-1 or that Asc is critical for another step which is important for inflammasome activation.

Materials and Methods

Ethic statement

All animal experiments were conducted according to the U.S.A. Public Health Service Policy on Humane Care and Use of Laboratory Animals. Animals were maintained in an AAALAC approved facility and all animal studies followed protocol 09716-2 that was approved by the Animal Care and Use Committee of the University of Michigan (Ann Arbor, MI).

Mice

Mice deficient in Nlrc4, Nlrp3, Asc and caspase-1/11 have been previously described [4,34,35]. All mice were crossed at least 5 times on a C57BL/6 background. Bone marrow samples from Pkdcl−/− mice in C57BL/6 background were provided by Hee-Jeong Im Sampen (Rush University Medical Center, Chicago, IL).

Bacterial strains and plasmids

Shigella flexneri strain YSH6000 [36] was used as the WT strain, and S325 (mxiA::Tn5) [37] was used as the T3SS-deficient control. The WT S. enterica serovar Typhimurium SR-11 y3181 and the isogenic flaA::Tn10 were provided by H. Matsui (Kitasato Institute for Life Science, Tokyo, Japan) [38]. AflA Salmonella mutant is impaired in the expression of flagellin [18]. CDNAs encoding mouse Naip2, Naip5, Nlrc4, Asc, caspase-1, and bacterial MxiI were amplified by PCR and cloned into the pCMV based mammalian expression vector or the MSCV-IRES-GFP retroviral expression vector (Addgene). Human pro-IL-1β clone (RDB6666) was provided by RIKEN BRC which is participating in the National Bio-Resource Project of the MEXT, Japan.

Cell culture and reagents, antibodies

BMDMs were prepared from the femurs and tibias of mice and cultured for 3–7 days in 10% FCS IMDM (Gibco) supplemented with 30% L-cell supernatant, non-essential amino acids, sodium pyruvate and antibiotics (Penicillin/Streptomycin). 293T cells were cultured on Dulbecco’s Modified Eagle’s medium (Sigma) containing 10% FCS and antibiotics (Penicillin/Streptomycin). The rabbit anti mouse caspase-1 p20 and anti-mouse Nlrc4 antibodies were produced in our laboratory by immunizing rabbits with mouse caspase-1 p20 subunit and mouse Nlrc4 (amino acids 1–152) recombinant proteins [39]. Anti–IL-1β p17 (#2021) and anti–Pkcδ (#2058) antibodies were from Cell Signaling. Mouse monoclonal anti–β-actin antibody was from Sigma. HRP-conjugated goat anti–rabbit (Jackson Laboratories) or anti–mouse IgG (Sigma) or anti–rat (Jackson Laboratories), or AP-conjugated goat anti-rabbit (Santa Cruz Biotechnology Inc.) or anti-mouse IgG (Santa Cruz Biotechnology Inc.) antibodies were used as secondary antibodies for immunoblotting.

Bacterial infection

Macrophages were seeded in 24-well plates at a density of 5 × 10^5 cells per well. Cells were stimulated with or without 0.1 μg/ml LPS (from E. coli O55:B5, Sigma) for 6 h and then infected with Shigella or Salmonella. Bacterial strains were pre-cultured overnight in Mueller-Hinton broth (Difco) at 30°C, then were inoculated into brain heart infusion broth (Difco) and incubated for 2 h at 37°C prior to infection. The cells were infected with Shigella at a bacteria/macrophage ratio of 10:1, or with Salmonella at a bacteria/macrophage ratio of 1:1 unless otherwise stated. The plates were centrifuged at 700 g for 5 min to synchronize the infection, and gentamicin (100 μg/ml) and kanamycin (60 μg/ml) were added after 20 min. At the indicated times after infection, cytokines were measured in culture supernatants by enzyme-linked immunoabsorbent assay (ELISA) kits (R and D Systems). RNA was isolated with E.Z.N.A. TM total RNA kit (Omega BioTec) according to the manufacturer’s instructions. RNA was reverse transcribed using the High Capacity RNA-to cDNA kit (Applied Biosystem) and cDNA was then used for RT-PCR. For immunofluorescence studies, the infected cells were fixed and immunostained, and then analyzed with a confocal laser-scanning microscope (LSM510; Carl Zeiss) or fluorescence microscopy (Olympus). Carboxyfluorescin FLICA (Immunochemistry Technologies, LLC) was added 1 hr before bacterial infection. Apoptosis was measured by the AnnexinV (Roche) and TUNEL (Promega) assays using fluorometric protocols according to the manufacturer’s recommendations. For the caspase-3 inhibitor studies, the cells were treated with 200 μM z-DEVD-fmk (Calbiochem) for 1 h before bacterial infection.

Reconstitution of the NLRC4 inflammasome in 293T cells

293T cells were seeded in 6-well plates at a density of 5 × 10^5 cells per well and incubated overnight. Then, the cells were transfected with or without 1 μg T7-tagged Nlrc4, 1 μg T7-tagged Asc, 0.4 μg HA-tagged caspase-1, and 0.4 μg FLAG-tagged proIL-1β [40], and 1 μg HA-tagged Naip2 or Naip5, using FuGENE 6 (Roche). Cells were infected one day after infection. Intensities of casp1 p20 or IL-1β p17 bands were quantified by densitometry, the values normalized to the β-actin protein levels and results were analyzed with ImageJ software.

Cell survival assays

The Shigella MxiI gene was cloned into the MSCV-ires-GFP retrovirus vector, which contains an IRES-GFP element to track retroviral infection. WT or Nlrc4−/− BMDMs were immobilized using the J2 virus to increase nucleofection efficiency [41]. Then, cells were nucleofected with MSCV-ires-GFP or MSCV-ires-GFP encoding Shigella MxiI using an Amaxa nucleofector system (Nucleofector kit V and the D-032 program). After 20 hrs, cell survival in the GFP-positive cell population was analyzed by fluorescence microscopy. The LDH activity in the culture supernatants of infected cells was measured using the CytoTox 96 assay kit (Promega) according to the manufacturer’s protocol. Assays were performed in triplicate for each independent experiment.

Bacterial invasion assay

The invasion efficiency of Shigella strains was evaluated using a gentamicin/kanamycin protection assay. Briefly, cells were infected
for 20 min and then incubated for 20 min at 37°C in medium containing gentamicin (100 μg/ml) and kanamycin (60 μg/ml) to kill extracellular bacteria. The infected cells were then washed in PBS, lysed in 0.5% TritonX-100/PBS, and serial dilutions were plated on LB agar plates to determine the number of intracellular bacteria.

DNA and siRNA transfection

DNA and siRNAs specific for Naip2 and Naip5 were transfected into macrophages using an Ammax nucleofector system (Y-001 program for primary macrophages or D-032 program for cell lines) according to the manufacturers’ instructions. siRNA pools for mouse Naip2 (17948; D-04131-01-04) and Naip5 (17951; D-04141-01-04) and non-targeting siRNAs were purchased from Dharmacon or synthesized by Sigma and targeting the sequences CTTACATGAAATACAGA (naip2) or GTGCCCTTTTATGCTCTTGT (naip5). Primer sets for RT-PCR were naip2-forward (AGGCCATGACATCCTGAGCACA), naip2-reverse (AACACATACTCCAGGCA), naip5-forward (TGCCCAAACCTTAAAGAGCTGTA), naip5-reverse (CAAGGCTTTTAGACTGGG-GATG), actin-forward (CATGTACGTTGCATCCAGGC) and actin-reverse (CTCTTAAATGTCCACGAGT). To compare caspase-1 p20 levels in immunoblotting experiments, the bands were quantified by densitometry, analyzed with ImageJ software, and normalized to the β-actin protein levels.

Immunoprecipitation

Cell were lysed in IP buffer (Celllytic M Cell Lysis Reagent (Sigma), 0.1 mM PMSF, and a complete protease inhibitor cocktail-EDTA (Roche)) and forced onto a 21-gauge needle 10 times. The cell lysates were centrifuged for 1 hr at 4°C with gela
tine rotation in IP buffer. Beads were washed with PBS, mixed with SDS-sample buffer and subjected to immunoblot analysis.

Immunostaining of Asc pyroptosomes

Cells were fixed with 4% paraformaldehyde and 0.1% NP40, washed and stained with anti-Asc antibody and FITC-conjugated anti-rat antibody (Sigma) as a secondary antibody. Imaging analysis was performed using fluorescence microscopy (Olympus), and percentage of cells containing Asc pyroptosomes was determined by counting at least 300 cells in 5 separate fields.

Asc dimerization assay

The Asc dimerization assay was previously described [25–27]. Briefly, cells were lysed (20 mM HEPES-KOH, pH 7.5, 150 mM KCl, 1% NP-40, 0.1 mM PMSF, and Complete protease inhibitor cocktail (Roche)) and forced onto a 21-gauge needle 10 times. The cell lysates were centrifuged at 6000 rpm for 10 min at 4°C to isolate the insoluble fraction in the pellet. The pellets were washed twice with PBS, resuspended in 500 μl of PBS and cross-linked with fresh 2 mM disuccinimidyl suberate (DTT, Sigma) for 30 min. The cross-linked pellets were isolated by centrifugation at 13000 rpm for 10 min and resuspended in 20 μl of SDS sample buffer for immunoblotting with anti-mouse Asc antibody.

Measurements of cytokines

Mouse cytokines in culture supernatants were measured by ELISA kits (R&D Systems). Assays were performed in triplicate for each independent experiment.

Statistical analyses

Statistical analyses were performed using the Mann–Whitney U test. Differences were considered significant at p<0.05.

Supporting Information

Figure S1 Naip2 promotes the processing of IL-1β in an inflammasome reconstitution system. A–B. 293T cells were transfected with vector producing Nlr4, Asc, caspase-1, pro-IL-1β, and HA-tagged Naip2 (A) or Naip5 (B). After 24 hrs, cells were infected with Shigella WT or S325 mutant for additional 3 hrs at a bacteria/cell ratio of 10:1. Cell lysates were analyzed by immunoblotting with the indicated antibodies or with anti-HA for Naip2/5 detection. The level of IL-1β was further quantified by densitometry. (A and B) Results are representative of three independent experiments. *p<0.05.

Figure S2 Inflammasome activation induced by Shigella infection is Asc- and Nlrc4- dependent, but Nlrp3-independent. (A–C) WT, Asc−/−, Nlrc4−/− or Nlrp3−/− BMDM were generated after differentiation for 4 days and infected with Shigella WT or S325 mutant at a bacteria/macrophase ratio of 10:1 for 30 min or the indicated time points. The activation of caspase-1 was evaluated by detecting cleaved caspase-1 (p20) by immunoblotting (A), the production of IL-1β in cell free supernatants was analyzed by ELISA (B), * p<0.0001. Results represent mean ± SD. Results are representative of three independent experiments.

Figure S3 WT and mutant Shigella uptake by BMDMs. BMDM were infected with Shigella for 20 min at a bacteria/macrophase of 10:1 and then incubated for 20 min at 37°C in medium containing gentamicin (100 μg/ml) and kanamycin (60 μg/ml) to kill extracellular bacteria. Cells were then washed in PBS, lysed in 0.5% TritonX-100/PBS and the number of intracellular bacteria evaluated by serial dilution on agar plates. The results represent mean ± SD and representative of at least three independent experiments.

Figure S4 The role of Asc in pyroptosis is influenced by the time of differentiation of bone marrow-derived macrophages in culture. BMDMs from WT and Asc−/− mice were differentiated in culture for 3, 4 and 5 days, infected with Shigella WT or S325, and subjected to the LDH assay 2 hr post-infection. *p<0.01. NS, not significant. The results represent mean and representative of at least three independent experiments.

Figure S5 Pkcα is essential for apoptosis and inhibition of apoptosis results in up-regulation of inflammasome. (A) Activation of MAPK and NF-kB pathways in WT and Pkcd−/− BMDMs were assessed by western blotting using Erk1/2, MEK1/2, and IκBα phospho-antibodies at the indicated time after Shigella infection. (B and D) Pkcd−/− regulates apoptosis induced by Shigella. BMDMs from WT and Pkcd−/− mice were infected with WT Shigella and after 2 hrs (B) or 5 hrs (C) post-infection, cells were subjected to phosphatidyserine staining with AnnexinV (B) or TUNEL assay (C) to assess the percent of apoptotic cells in infected macrophages. * p<0.0001. Results are based on the analysis of at least 300 hundred cells in 10 microscope fields. (D) z-DEVd-fink treatment enhances inflammasome activation. WT BMDMs were treated with caspase-3 inhibitor z-DEVd-fink for 1 hr, then infected with WT Shigella. IL-1β levels in the supernatants were measured by ELISA at 1 hr post-infection. * p<0.0022. Results represent mean ± SD. Results are representative of at least three experiments.
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