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Critical Roles of ASC Inflammasomes in Caspase-1 Activation and Host Innate Resistance to Streptococcus pneumoniae Infection

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Streptococcus pneumoniae is a Gram-positive, extracellular bacterium that is responsible for significant mortality and morbidity worldwide. Pneumolysin (PLY), a cytolysin produced by all clinical isolates of the pneumococcus, is one of the most important virulence factors of this pathogen. We have previously reported that PLY is an essential factor for activation of caspase-1 and consequent secretion of IL-1β and IL-18 in macrophages infected with S. pneumoniae. However, the host molecular factors involved in caspase-1 activation are still unclear. To further elucidate the mechanism of caspase-1 activation in macrophages infected with S. pneumoniae, we examined the involvement of inflammasomes in inducing this cellular response. Our study revealed that apoptosis-associated speckle protein containing a caspase recruitment domain (ASC), an adaptor protein for inflammasome receptors such as nucleotide-binding oligomerization domain-like receptor family, pyrin domain containing 3 (NLRC3) and absent in melanoma 2 (AIM2), is essentially required for the induction of caspase-1 activation by S. pneumoniae. Caspase-1 activation was partially impaired in NLRP3−/− macrophages, whereas knockdown and knockout of AIM2 resulted in a clear decrease in caspase-1 activation in response to S. pneumoniae. These results suggest that ASC inflammasomes, including AIM2 and NLRP3, are critical for caspase-1 activation induced by S. pneumoniae. Furthermore, ASC−/− mice were more susceptible than wild-type mice to S. pneumoniae, with impaired secretion of IL-1β and IL-18 into the bronchoalveolar lavage fluid after intranasal infection, suggesting that ASC inflammasomes contribute to the protection of host from infection with PLY-producing S. pneumoniae. The Journal of Immunology, 2011, 187: 4890–4899.
inflammatory cytokines and their roles in vivo have been established; however, little is known about the mechanism by which these cytokine responses are induced in a PLY-dependent manner. Caspase-1 is a cysteine protease that processes proforms of IL-1β and IL-18 into mature forms and is important for secretion of these cytokines. We have also shown that caspase-1 activation is dependent on PLY in macrophages infected with *S. pneumoniae* (10).

Recent studies have demonstrated that the inflammasome, a multiprotein complex that typically consists of a nucleotide-binding oligomerization domain-like receptor (NLR) molecule and procaspase-1, mediates the activation of caspase-1 in response to a wide variety of stimuli (19–38). Several inflammasomes have been identified, and each has been designated according to the receptor within the protein complex. Of these, the NLR family, pyrin domain containing 3 (NLRC4) inflammasome has been shown to be activated by endogenous stimuli such as urate crystals (20), extracellular ATP (21), cholesterol crystals (22), and also exogenous stimuli such as bacterial and viral pathogens, pore-forming toxins, asbestos, silica, and so on (23–30). It has been reported that the NLR family, caspase recruitment domain containing 4 (NLRC4) inflammasome is activated by flagellin, basal body rod proteins of the type III secretion systems, and pilin, whereas cytosolic DNA activates the absent in melanoma 2 (AIM2) inflammasome (31–38). Apoptosis-associated speck-like protein containing a caspase recruitment domain (ASC) plays a critical role in the activation of inflammasomes as an adaptor protein that bridges procaspase-1 and inflammasome receptors like NLRC3 and AIM2 (19, 39). Recent reports have identified a distinct role for inflammasomes in host defense against several pathogens.

In this study, we sought to elucidate the induction mechanism of PLY-dependent activation of caspase-1 in macrophages infected with *S. pneumoniae* by examining the involvement of inflammasomes in the cellular response. Our results suggest that the activation of caspase-1 and the subsequent maturation of IL-1β and IL-18 in response to PLY-producing *S. pneumoniae* are critically dependent on ASC inflammasomes, including AIM2 and NLRC3. We also demonstrated a significant role of ASC inflammasomes in host defense against pulmonary pneumococcal infection. Thus, this study has revealed a pathophysiologic link between the virulence factor PLY and the inflammasome.

### Materials and Methods

#### Mice

Female C3H/HeN, C3H/HeJ, and C57BL/6 mice were purchased from Japan SLC (Hamamatsu, Japan). NLRC3−/− mice generated by Prof. Jürg Tschopp (University of Lausanne) (20), NLRC4−/− mice generated by Dr. Vishva Dixit (Genentech) (40), ASC−/− mice generated by Prof. Shun-ichiro Taniguchi (Shinshu University) (41), and caspase-1−/− mice generated by Dr. Keisuke Kuida (Millennium Pharmaceuticals) were kindly gifted. Mice were maintained in specific-pathogen-free conditions and used at 7–9 wk of age. All of the experimental procedures performed on mice were approved by the Animal Ethics and Research Committee of Kyoto University Graduate School of Medicine, Kyoto, Japan.

#### Bacterial strains

*Streptococcus pneumoniae* D39 (serotype 2) was purchased from the National Collection of Type Cultures (NCTC 7466; Central Public Health Laboratory, London, U.K.). A deletion mutant of *S. pneumoniae* D39 for the PLY gene (*ply*) was constructed by using a homologous recombination method as described previously (10). Bacteria were grown on tryptic soy agar (Difco Laboratories, Detroit, MI) with 5% (v/v) de-fibrated sheep blood (Nacalai Tesque, Kyoto, Japan) and in Todd–Hewitt broth (Difco Laboratories) supplemented with 0.5% yeast extract (THY) at 37°C and 5% CO₂ and subsequently stored at −80°C in THY plus 10% glycerol. For the preparation of bacterial stocks for macrophage stimulation, pneumococci were grown overnight on blood agar plates at 37°C and 5% CO₂. Colonies were inoculated into the THY medium, grown until midlogarithmic phase (OD at 600 nm [OD600] = 0.5), and centrifuged at 6000 × g for 15 min. The bacterial pellet was suspended in PBS and stored at −80°C. The concentration was determined by viable cell counting on blood agar plates.

**Collection of peritoneal macrophages and infection by *S. pneumoniae* in vitro**

Mice were injected i.p. with 5% thiglycollate broth (Eiken Chemical, Tokyo, Japan), and peritoneal exudate cells (PECs) were collected 3 d later. After washing with RPMI 1640 medium (Nacalai Tesque), PECs were suspended with RPMI 1640 supplemented with 10% FCS and incubated in 48-well microplates at a density of 2.5 × 10⁵ cells/well at 37°C plus 5% CO₂, and nonadherent cells were removed after 2 h. Adherent macrophages were infected with *S. pneumoniae* at a multiplicity of infection (MOI) of 10 for 8 h, then 100 µg/ml gentamicin (Wako Pure Chemical Industries, Osaka, Japan) was added to the cultures and incubated for an additional 16 h. To confirm the enrichment for macrophages in adherent PECs, the percentage of F4/80⁺ macrophages was determined by flow cytometry. Whole and adherent PECs collected by using Cell Dissociation Buffer Enzyme-Free PBS-based (Invitrogen, Carlsbad, CA) were incubated with anti-mouse CD16/32 (BioLegend, San Diego, CA) for 30 min at 4°C. Then the cells were stained with PE anti-mouse F4/80 (BioLegend) for 30 min at 4°C and analyzed on an FACSCalibur flow cytometer (BD Biosciences, Franklin Lakes, NJ). The proportions of F4/80⁺ cells in whole PECs from wild-type (WT), caspase-1−/−, ASC−/−, and NLRC3−/− mice were 94.8, 96.0, 95.6, and 95.0%, respectively, and were increased to 98.2, 98.5, 98.3, and 95.9% in adherent PECs, respectively. Thus, the adherent PECs were sufficiently enriched for F4/80⁺ macrophages.

**Macrophage cell lines**

Immortalized WT and AIM2−/− macrophage cell lines were generated by infecting primary bone marrow cells with J2 recombinant retrovirus (a kind gift from Dr. Howard Young, National Cancer Institute at Frederick, Frederick, MD) as described previously (42). The macrophage lines were infected with *S. pneumoniae* at an MOI of 10 for 8 h, and then 100 µg/ml gentamicin was added to the cultures and incubated for an additional 16 h.

**ELISA**

Levels of secreted cytokines in culture supernatants were determined by two-site sandwich ELISA. ELISA kits for IL-6 and IL-1β were purchased from eBioscience (San Diego, CA). For the titration of IL-18, biotin-labeled and unlabeled mAbs specific to IL-18 (Medical & Biological Laboratories, Nagoya, Japan) were used.

**Western blot analysis**

PECs were cultured on 12-well plates at a density of 5 × 10⁵ cells/well in RPMI 1640 plus FCS at 37°C for 2 h. Then medium was replaced with Opti-MEM (Invitrogen), and adherent cells were infected with *S. pneumoniae* at an MOI of 10. Supernatants were collected 24 h postinfection, and the cells were lysed with radioimmunoprecipitation assay buffer (Nacalai Tesque). The supernatants were concentrated 20-fold using 20% (w/v) trichloroacetic acid (Nacalai Tesque). The precipitates and cell lysates were subjected to SDS-PAGE and subsequently transferred to polyvinylidene difluoride membranes by electroblotting. The membranes were immunoblotted with anti–caspase-1 Ab (Santa Cruz Biotechnology, Santa Cruz, CA), anti–IL-1β Ab (R&D Systems, Minneapolis, MN), or anti–β-actin mAb (Sigma-Aldrich, St. Louis, MO) (43).

**Detection of lactate dehydrogenase release**

Lactate dehydrogenase (LDH) activity was measured using an LDH detection kit (TaKaRa BIO, Otsu, Japan). The percentage of LDH release was calculated by using the following formula: percentage of release = 100 × (experimental LDH release − spontaneous LDH release)/(maximal LDH release − spontaneous LDH release). To determine the maximal LDH release, cells were treated with 1% Triton X-100. For spontaneous LDH release, adherent PECs were left uninfected.

**RNA interference**

RNA interference experiments were done using the method employed in our previous study (43). Briefly, PECs were transfected with Stealth siRNA duplexes (Invitrogen) at a final concentration of 20 nM using siPORT Amine (Ambion, Austin, TX). The transfected cells were incubated at 37°C for 24 h posttransfection.
37°C for 48 h and used after washing with RPMI 1640 plus FCS. The sense small interfering RNA (siRNA) sequences were: AIM2-1, 5'-UUA UCU UCU GGA CUU UAA ACA GCC C-3' and AIM2-2, 5'-UAA AGU CAU UGU CAC UGC GGG UGG C-3'. Stealth RNAs Negative Control Medium GC Duplex #2 (Invitrogen) was used as a control siRNA.

**RT-PCR analysis**

Total cellular RNA of adherent PECs was extracted by using NucleoSpin RNA Clean-up (MACHEREY-NAGEL, Düren, Germany), and quantitative real time RT-PCR was carried out using the ABI PRISM 7000 (Applied Biosystems, Foster City, CA) and EXPRESS Two-Step SYBR GreenER (Invitrogen), according to the manufacturer’s instructions. Primers for quantitative real-time RT-PCR were as follows: IFN-β forward, 5'-CTG GAG CAG CTG AAT GGA AAG-3' and reverse, 5'-CTT GAA GTC CGC CCT GTA GG-3'; β-actin forward, 5'-GCC CTG ATG CCT TCT TCC AG-3' and reverse, 5'-TGC CAC AGG ATT CTA TAC CC-3'; and AIM2 forward, 5'-TTG TAT CTA GGC TGA TCC TGG GAC-3' and reverse, 5'-ACC TGC ACT TTG AAT CAG GTG TGG-3'.

**Extraction of *S. pneumoniae* genomic DNA**

*S. pneumoniae* D39 was inoculated into THY broth and culture for overnight, then the bacteria was centrifuged, and the pellet was suspended in the lysis buffer containing 10 mM Tris buffer, 50 mM EDTA, 20 mg/ml lysozyme (Nacalai Tesque), and 100 μg/ml RNaseA (Sigma-Aldrich) and then incubated at 37°C for 1 h. Next, the lysate was mixed with 5 M NaCl and 10% CTAB/0.7 M NaCl solution and incubated at 65°C for 10 min. Finally, the genomic DNA was extracted by chloroform/isoamyl alcohol (24:1) and phenol/chloroform/isoamyl alcohol (25:24:1). To stimulate macrophages with purified *S. pneumoniae* DNA or poly(deoxyadenylic-deoxythymidylidic) acid (poly dA:dT; Sigma-Aldrich), macrophages were transfected with 400 ng DNA by using Lipofectamine LTX (Invitrogen).

**Quantitation of macrophage-associated and phagocytosed *S. pneumoniae***

Adherent macrophages were infected with *S. pneumoniae* at an MOI of 10 for 2, 4, 6, or 8 h. To enumerate *S. pneumoniae* associated with macrophages, cells were washed with chilled PBS three times to remove non-associating bacteria and lysed in PBS containing 0.1% Triton X-100. The cell lysates were diluted with PBS and plated on blood agar plates. CFU were counted after overnight incubation. To enumerate the phagocytosed bacteria, macrophages were infected with *S. pneumoniae* as above and additionally cultured for 30 min in the presence of 100 μg/ml gentamicin. The cells were then washed, and the bacterial numbers were counted. To monitor the survival of *S. pneumoniae* inside macrophages, the numbers of intracellular bacteria were determined every 0.5 h until 3 h after gentamicin addition. To inhibit the phagocytosis of bacteria by macrophages, cytochalasin B was added to the macrophage cultures at a final concentration of 10 μg/ml. An equal volume of DMSO was added to macrophages as a solvent control.

**Intranasal infection of mice**

Mice were anesthetized with pentobarbital (Nacalai Tesque) and inoculated with 5 × 10^10^ bacteria in 25 μl PBS. Bronchoalveolar lavage fluid (BALF) was collected according to the method of Lauw et al. (17). BALF from noninfected mice was collected as negative control. Survival of infected mice was monitored every day until 2 wk postinfection and analyzed using the Kaplan–Meier method. Lungs were excised 48 h postinfection and homogenized in 1 ml PBS, serial diluted in PBS, and plated onto blood agar; CFUs were counted after overnight culture.

**Statistical analysis**

For comparisons between two groups, the Student t test was used. Multigroup comparisons of mean values were made according to the ANOVA and Bonferroni post hoc test. Statistical analysis for survival curves was performed using the log-rank test. Statistical significance was determined as p < 0.05.

**Results**

**Caspase-1 activation in macrophages induced by *S. pneumoniae***

We have previously reported that PLY plays an important role in caspase-1 activation and the subsequent maturation and secretion of caspase-1-dependent cytokines in macrophages infected with *S. pneumoniae* (10). However, the host receptors involved in this process still remain unclear. In our previous report, we also showed that secretion of IL-1β and IL-18 induced following the stimulation of macrophages with recombinant protein of PLY is critically dependent on TLR4. We therefore first tested whether PLY-producing *S. pneumoniae* activates caspase-1 through TLR4. Macrophages from C3H/HeN and C3H/HeJ (tlr4<sup>d4–d8</sup>) mice were infected with *S. pneumoniae* D39 at an MOI of 10 for 24 h, and then the culture supernatants were assayed for the p10 fragment of mature caspase-1 by Western blotting. There was no significant difference between the two mouse strains in caspase-1 activation in response to *S. pneumoniae* (Fig. 1A). Furthermore, HeN and HeJ macrophages secreted comparable levels of mature IL-1β postinfection with *S. pneumoniae* (Fig. 1A, 1B). The level of IL-1β in HeJ macrophages was almost the same as that in HeN macrophages following a treatment with PamC3SK4 (a TLR2 ligand) plus nigericin, whereas HeJ macrophages never responded to the treatment with LPS (a TLR4 ligand) plus nigericin (Fig. 1C, 1D). These results suggested that *S. pneumoniae* induces caspase-1 activation in a TLR4-independent manner.

**ASC plays a critical role in caspase-1 activation in macrophages infected with *S. pneumoniae***

Caspase-1 was essential for *S. pneumoniae*-induced secretion of IL-1β and IL-18, as macrophages from caspase-1<sup>−/−</sup> mice were incapable of secreting these cytokines postinfection with *S. pneumoniae* D39 (Fig. 2A, 2B). LDH release occurred following infection of normal macrophages with *S. pneumoniae* D39 but not the Δply strain (Fig. 2C), and D39-induced LDH release was not observed in caspase-1<sup>−/−</sup> macrophages. It appeared that this bacterium induces macrophage pyroptosis in a PLY-dependent manner. NLRP3 is a well-studied NLR receptor and has been shown to induce caspase-1 activation in response to cholesterol-dependent cytolysins (24–26). Therefore, we investigated whether NLRP3 and its adaptor protein ASC are involved in caspase-1 activation upon *S. pneumoniae* infection. Caspase-1 activation, secretion of IL-1β and IL-18, and LDH release induced postinfection with *S. pneumoniae* D39 were completely abolished in ASC<sup>−/−</sup> macrophages, whereas these cellular responses in NLRP3<sup>−/−</sup> macrophages were only partially reduced as compared with those in WT macrophages (Fig. 2A–C). Deficiency of NLRC4 did not affect or only slightly reduced the secretion of IL-1β and IL-18 and LDH release induced by *S. pneumoniae* D39 infection (Fig. 2B, 2C). Taken together, these results suggested that ASC is essential for the caspase-1 response to *S. pneumoniae* and additional receptor(s), other than NLRP3, are involved in ASC-dependent caspase-1 activation induced by this pathogen.

**AIM2 inflammasome is involved in caspase-1 activation induced by *S. pneumoniae***

AIM2 has been shown to recognize cytoplasmic DNA and to be responsible for ASC-dependent caspase-1 activation in response to DNA (35–38). We tested whether the AIM2 inflammasome is involved in caspase-1 activation in macrophages infected with *S. pneumoniae*. Knockdown of AIM2 was carried out in macrophages using two siRNAs, AIM2-1 and AIM2-2, which target independent sequences in the AIM2 transcript, as described previously (43). The efficiency of the knockdown was assessed by real-time RT-PCR, and the expression of AIM2 was lowered by both AIM2-targeting siRNAs (Fig. 3A). Knockdown of AIM2 resulted in a reduction of IL-18 secretion induced by poly dA:dT transfection (Fig. 3B) without affecting the secretion of IL-1β in response to LPS plus nigericin (Fig. 3C). A significant reduction in *S. pneumoniae*-induced caspase-1 activation was observed in

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AIM2 knockdown macrophages (Fig. 3D). Moreover, secretions of IL-1β, IL-18, and LDH release in response to *S. pneumoniae* were decreased by knockdown of AIM2, whereas IL-6 response was not affected (Fig. 3D–F). These findings suggest that the AIM2 inflammasome plays an important role in caspase-1 activation induced by *S. pneumoniae*. To confirm the involvement of

**FIGURE 2.** ASC is essentially required for caspase-1 activation in *S. pneumoniae*-infected macrophages. Adherent PECs from C57BL/6 WT, caspase-1−/−, ASC−/−, NLRP3−/−, or NLRC4−/− mice were left uninfected or infected with *S. pneumoniae* D39 or the Δply strain at an MOI of 10 for 24 h as described in Fig. 1, and culture supernatants and cell lysates were then collected. A, Culture supernatants and cell lysates were subjected to Western blot analysis as described in Fig. 1. Filled and open triangles indicate molecular mass markers of 47 and 9 kDa, respectively. B, Levels of IL-1β in the culture supernatants were determined by ELISA. C and D, Adherent PECs were left unstimulated or stimulated with Pam3CSK4 (100 ng/ml) or LPS (10 mg/ml) for 3 h and sequentially with nigericin (5 μM). After 21 h, culture supernatants were collected to assess the levels of IL-1β by ELISA. All of the experiments were repeated three times. The ELISA data are presented as the mean and SD of triplicate assays. **FIGURE 1.** Caspase-1 activation in macrophages induced by *S. pneumoniae* is independent of TLR4. A and B, Adherent PECs from C3H/HeN or C3H/HeJ mice were left uninfected or infected with *S. pneumoniae* D39 at an MOI of 10; 8 h later, gentamicin was added to cultures (final concentration 100 μg/ml), and culture supernatants were collected after an additional 16 h incubation (24 h postinfection). A, Culture supernatants were concentrated by trichloroacetate precipitation. The precipitates and cell lysates were subjected to Western blot analysis using Abs specific for caspase-1, IL-1β, or β-actin. Filled and open triangles indicate molecular mass markers of 47 and 9 kDa, respectively. B, Levels of IL-1β in the culture supernatants were determined by ELISA. C and D, Adherent PECs were left unstimulated or stimulated with Pam3CSK4 (100 ng/ml) or LPS (10 mg/ml) for 3 h and sequentially with nigericin (5 μM). After 21 h, culture supernatants were collected to assess the levels of IL-1β by ELISA. All of the experiments were repeated more than three times. The ELISA data are presented as the mean and SD of triplicate assays. Statistical significance was determined by one-way ANOVA followed by the Bonferroni test. *p < 0.05.
FIGURE 3. AIM2 is involved in caspase-1 activation upon *S. pneumoniae* infection. A–F, Adherent PECs were transfected with siRNAs (AIM2-1, AIM2-2, or control siRNA) and cultivated for 2 d. G–K, Immortalized macrophages from WT or AIM2<sup>−/−</sup> mice were employed. A, AIM2 expression levels were analyzed by real-time RT-PCR. B, Cells were transfected or not with poly dA:dT and incubated for 24 h. Culture supernatants were then collected, and levels of IL-18 in the culture supernatants were determined by ELISA. C and K, Cells were left unstimulated or stimulated with LPS (10 ng/ml) for 3 h and sequentially with nigericin (5 μM). After 21 h, culture supernatants were collected, and levels of IL-1β in the culture supernatants were determined by ELISA. D–F and G–J, Cells were left uninfected or infected with *S. pneumoniae* D39 at an MOI of 10 for 24 h as described in Fig. 1, and culture supernatants and cell lysates were then collected. Levels of IL-1β (E, H), IL-18 (E, I), and IL-6 (E) in the culture supernatants were determined by ELISA. Culture supernatants and cell lysates were subjected to Western blot analysis as described in Fig. 1. Filled and open triangles indicate molecular mass markers of 47 and 9 kDa, respectively (D, G). LDH release from D39-infected macrophages was determined by the LDH activity in the culture supernatants, and the data are expressed as percent LDH release (F, J). L, Adherent PECs were infected with *S. pneumoniae* D39 or Δply at an MOI of 10 for 3, 6, or 9 h, and the expression of IFN-β was quantified by real-time RT-PCR. White bars, untreated macrophages; black bars, macrophages infected with *S. pneumoniae* D39; and gray bars, macrophages infected with the Δply strain. All of the experiments were repeated more than three times. The results are presented as the mean and SD of triplicate assays. Tests for statistical significance were performed by using one-way ANOVA followed by the Bonferroni test (A, B, E, F, H, I, L) or Student t test (J). *p < 0.05.
AIM2, similar experiments were conducted with immortalized macrophages established from WT or AIM2 \(-/-\) mice, and the results showed that D39-induced caspase-1 activation, secretion of caspase-1-dependent cytokines, and LDH release were significantly decreased in the absence of AIM2 (Fig. 3G–J), though the NLRP3 inflammasome was intact in both macrophage lines (Fig. 3K), further suggesting the involvement of AIM2 in caspase-1 activation upon \(S.\ pneumoniae\) infection. However, a question may be raised at this point as to how \(S.\ pneumoniae\), an extracellular bacterium, is recognized by AIM2, a sensor molecule of cytosolic DNA. Because PLY is important for caspase-1 activation in response to \(S.\ pneumoniae\), we hypothesized that PLY may play a role in the delivery of bacterial DNA into the macrophage cytoplasm. If this is the case, a type I IFN response could also be induced by \(S. pneumoniae\) in a PLY-dependent manner, as cytosolic DNA induces not only activation of AIM2 inflammasome but also expression of type I IFNs (44, 45). Consistent with this hypothesis, we found that the expression of IFN-\(\beta\) was induced in macrophages infected with \(S. pneumoniae\) D39 but not with the \(\Delta\)ply strain (Fig. 3L).

\(S. pneumoniae\) genomic DNA activates the AIM2 inflammasome when introduced into macrophages

We next checked whether \(S. pneumoniae\) genomic DNA can be a ligand for AIM2. We purified genomic DNA from liquid cultures of \(S. pneumoniae\), and the DNA sample was split into one sample treated with DNase I and another left untreated (Fig. 4A). The DNA was then transfected into macrophages, and the activation of caspase-1 and the secretion of IL-18 were determined. \(S. pneumoniae\) genomic DNA without treatment could induce the activation of caspase-1 and the secretion of IL-18, whereas DNase I–treated genomic DNA did not retain such an activity (Fig. 4B, 4C). Moreover, knockdown of AIM2 in macrophages resulted in a decrease of secretion of IL-18 induced by \(S. pneumoniae\) genomic DNA (Fig. 4D). IL-1\(\beta\) was not secreted from macrophages transfected with \(S. pneumoniae\) genomic DNA (data not shown), probably due to the absence of pro–IL-1\(\beta\) expression (Fig. 4B). Taken together, these results indicated that \(S. pneumoniae\) genomic DNA can be recognized by AIM2 and that this then induces caspase-1 activation and the secretion of a caspase-1–dependent cytokine.

**Phagocytosis of \(S. pneumoniae\) by macrophages is important for caspase-1–dependent cytokine production**

Though \(S. pneumoniae\) is an extracellular bacterium that is not capable of invading the macrophage cytoplasm, the present findings indicated the recognition of \(S. pneumoniae\) by AIM2 that functions in the cytosolic space. A possible interpretation is the recognition of bacterial DNA possibly released from the phagosome after phagocytosis of bacteria by macrophages. We therefore tested whether phagocytosis of bacteria is a prerequisite for caspase-1 activation induced by \(S. pneumoniae\). Before the examination, we determined the fate of \(S. pneumoniae\) D39 in macrophages postinfection, because the bacterial strain has a capsule that is known to affect phagocytic processes (1–4). The numbers of bacteria associated with macrophages or phagocytosed by macrophages were counted at different time points postinfection, showing that the numbers of both macrophage-associated bacteria and phagocytosed bacteria reached their peaks at 6 h postinfection (Fig. 5A, 5B). We observed a rapid decrease of intracellular bacteria after addition of gentamicin 4 h postinfection (Fig. 5C). These results suggested that \(S. pneumoniae\) D39, though the strain is encapsulated, is phagocytosed by macrophages for several hours postinfection, and the phagocy-
viability of phagocytosed bacteria was determined. WT macrophages were infected with D39 or the cultured for an additional 30 min in the presence of 100 μg/ml gentamicin. The numbers of bacteria in cell lysates were then counted. Non–cell-associated bacteria were removed by washing at 2, 4, 6, and 8 h postinfection, and macrophages were untreated controls. All of the experiments were repeated three times. Tests for statistical significance were performed using one-way ANOVA followed by the Bonferroni test. *

FIGURE 5. Phagocytosis of S. pneumoniae by macrophages is important for caspase-1 activation. Adherent PECs were infected with S. pneumoniae D39 (A–D) or Δply (E–F) at an MOI of 10. A, Bacteria associating with macrophages were counted. Infected cells were washed with chilled PBS three times and lysed at the indicated time points. The lysates were plated on blood agar plates after serial dilutions, and CFU were counted after overnight incubation. B, Phagocytosed bacteria were counted. Non–cell-associated bacteria were removed by washing at 2, 4, 6, and 8 h postinfection, and macrophages were cultured for an additional 30 min in the presence of 100 μg/ml gentamicin. The numbers of bacteria in cell lysates were then counted. C and D. The viability of phagocytosed bacteria was determined. WT macrophages were infected with D39 or the Δply mutant (C). WT, ASC−/−, or caspase-1 macrophages were infected with D39 (D). Gentamicin was added to the cultures 4 h postinfection, and the cells were additionally incubated for the indicated times. The numbers of bacteria in cell lysates were then counted. E, WT macrophages were left untreated or pretreated with cytochalasin B or DMSO and then infected with S. pneumoniae D39. The phagocytosed bacteria were counted 6.5 h postinfection as described in B. F. Cytochalasin-pretreated or untreated PECs were infected with S. pneumoniae for 24 h, and culture supernatants and cell lysates were subjected to Western blot analysis as described in Fig. 1. Cytochalasin-pretreated or untreated macrophages were transfected with DNA or infected by S. pneumoniae for 24 h, levels of IL-18 and IL-1β in the supernatants were determined by ELISA, and the data are expressed as percent IL-18 induction (G) and percent IL-1β induction (H) with respect to untreated controls. All of the experiments were repeated three times. Tests for statistical significance were performed using one-way ANOVA followed by the Bonferroni test. *p < 0.05. ND, not detected.

decreased (Fig. 5G). Also, IL-1β secretion from S. pneumoniae-infected macrophages was significantly decreased by cytochalasin B (Fig. 5H). Thus, it appears that phagocytosis is important for the caspase-1 activation in macrophages infected with S. pneumoniae.

ASC inflammasomes protect the host from S. pneumoniae infection in vivo

To determine whether ASC-dependent inflammasome formation plays a role in vivo, we have employed an intranasal infection of S. pneumoniae in WT, ASC−/−, and NLRP3−/− mice. Consistent with the in vitro findings using cultured macrophages, a significant level of IL-1β was detected in the BALF of WT mice infected with S. pneumoniae D39 but not the Δply strain (Fig. 6A). To assess whether ASC inflammasomes are required for the secretion of IL-1β and IL-18 upon infection with S. pneumoniae D39 in vivo, the levels of these cytokines in the BALF were compared among WT, ASC−/−, or NLRP3−/− mice. Compared with WT mice, ASC−/− mice secreted significantly lower levels of IL-1β and IL-18 after S. pneumoniae infection (Fig. 6B, 6C). In contrast, NLRP3 was partially required for secretion of these cytokines (Fig. 6B, 6C). IL-6 secretion in response to S. pneumoniae infection, which is not dependent on inflammasomes, was not affected at all in ASC−/− and NLRP3−/− mice (Fig. 6D). These results are consistent with the in vitro findings that multiple ASC inflammasomes are necessary for inducing caspase-1 activation and maturation of IL-1β and IL-18 in S. pneumoniae-infected macrophages. Furthermore, the protection of mice as determined by survival curves and bacterial numbers in the lungs was significantly impaired in ASC−/− and NLRP3−/− mice (Fig. 6E, 6F). The Δply mutant was cleared more rapidly from the lungs of WT mice than the D39 parental strain (n = 4 in each group, data not shown), and when WT or ASC−/− mice were infected with the mutant strain, the bacterial numbers were comparable in the lungs of both mice at 48 h postinfection (n = 3 in each group, data not shown). These results have clearly indicated that ASC inflammasomes are essential for the secretion of caspase-1–dependent cytokines and eventual protection in mice infected with the PLY-producing strain of S. pneumoniae.

4896 ROLE OF ASC INFLAMMASOMES IN S. PNEUMONIAE INFECTION

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Innate immune responses play important roles in host defense against S. pneumoniae infection. In this study, we elucidated the mechanism of caspase-1 activation in macrophages infected with PLY-producing S. pneumoniae. We found that ASC inflammasomes, including both AIM2 and NLRP3, but not TLR4, are indispensable for inducing the activation of caspase-1 and the maturation and secretion of IL-1β and IL-18 and for pyroptosis. In addition, our results clearly demonstrate the essential requirement for ASC for the secretion of IL-1β and IL-18 into the BALF in a mouse model of pneumococcal pneumonia. We also showed that the absence of ASC results in a significant increase in the susceptibility to S. pneumoniae infection in vivo. Thus, this study revealed a novel role of ASC inflammasomes in mediating host resistance to pneumococcal pneumonia, most likely through the induction of the protective cytokines IL-1β and IL-18.

In contrast to our results, PLY has been shown to inhibit the production of proinflammatory cytokines including IL-1β by human DCs infected with the nonencapsulated mutant of S. pneumoniae TIGR4 (46), suggesting that the role of PLY in inducing host inflammatory responses may vary in different situations. In contrast, findings similar to ours were presented in recent reports demonstrating that PLY expression and the NLRP3 inflammasome are involved in the secretion of IL-1β from mouse DCs or macrophages infected with S. pneumoniae D39 (11, 12). These studies also suggested that NLRP3 is important for host defense against S. pneumoniae. All of these results are in agreement with ours, as in our present study, NLRP3−/− macrophages exhibited a decreased secretion of IL-1β and IL-18 in response to S. pneumoniae compared with WT macrophages, and we found NLRP3−/− mice to be more susceptible to pneumococcal pneumonia than WT mice. However, our data are indicative of the involvement of other inflammasome receptors in addition to NLRP3, judging from the observations that ASC seemed to be more essential for the caspase-1 response and host resistance to S. pneumoniae than NLRP3. Indeed, we identified AIM2 as a receptor involved in the activation of caspase-1 in NLRP3−/− macrophages. Thus, we suggest that not only NLRP3 but also other receptors, such as AIM2, are responsible for ASC-dependent secretion of IL-1β and IL-18 and consequent early defense against pneumococcal pneumonia.

We and other groups have demonstrated the ability of PLY to activate the TLR4 signaling, and we have previously shown that TLR4−/− macrophages secrete a lower level of IL-18 post-infection with S. pneumoniae D39 than WT macrophages (8, 10, 47, 48). In addition, it has been shown that in some cases, LPS, a canonical TLR4 ligand, induces the activation of the NLRP3 inflammasome dependent on TLR4 and a TLR adapter protein, Toll/IL-1 receptor domain-containing adapter protein inducing IFN-β (49). For these reasons, we assumed that PLY activates the NLRP3 inflammasome through recognition by TLR4. However, there was no significant difference in caspase-1 activation in response to S. pneumoniae between HeN macrophages and HeJ macrophages (Fig. 1). Moreover, McNeela et al. (11) recently showed that secretion of IL-1β induced by rPLY plus other TLR stimuli, including heat-killed S. pneumoniae, does not require TLR4. Thus, it appears that TLR4 is dispensable for the caspase-1 response to PLY-producing S. pneumoniae and that the decrease in S. pneumoniae-induced secretion of IL-18 in the absence of TLR4 that we previously published might be due to reasons other than caspase-1 activation. As experiments demonstrating the ability of PLY to activate TLR4 signaling were usually performed using large amounts of PLY protein, the interaction between PLY and TLR4 may be not always be accurately reflected in innate immune responses to PLY-producing S. pneumoniae.

AIM2 is a cytoplasmic receptor that recognizes dsDNA in the cytoplasm, and we have verified that S. pneumoniae genomic DNA can be a ligand for AIM2. In the current study, knockdown or the deficiency of AIM2 caused significant decreases in caspase-1 activation, secretion of IL-1β and IL-18, and pyroptosis in...
response to *S. pneumoniae*. These results suggest that the AIM2 inflammasome is involved in caspase-1 activation in *S. pneumoniae*-infected macrophages. However, how AIM2 detects DNA upon infection with the extracellular pathogen remains unclear. In our study, inhibiting of bacterial uptake significantly reduced the activation of caspase-1 in *S. pneumoniae*-infected macrophages, suggesting that phagocytosis of bacteria by macrophages is an important process leading to the activation of AIM2 inflammasome. Although *S. pneumoniae* engulfed by macrophages was shown to undergo rapid death, which might cause the release of bacterial DNA from the bacterial cell, DNA in the phagosome would hardly be diffused across the phagosomal membrane, degraded by lysosomal hydrolases, such as DNase II, and therefore not be detected by AIM2. Importantly, PLY, a pore-forming toxin secreted upon the breakdown of the pneumococcal cell wall, is required for the activation of caspase-1 in response to *S. pneumoniae*. It is thus conceivable that PLY released from autolyzed or killed bacteria in the phagosome might cause destabilization of the phagosomal membrane, leading to the leakage of bacterial DNA into the cytoplasm. AIM2 has been reported to play a role in caspase-1 activation in infection with *Francisella tularensis*, *Listeria monocytogenes*, and DNA viruses that are all characterized as intracellular parasitic pathogens (50–54). By contrast, *S. pneumoniae* is generally regarded as an extracellular bacterium. Therefore, activation of the AIM2 inflammasome is not a host response restricted to infection by intracellular parasitic microbes, and other pathogens may also activate the AIM2 inflammasome regardless of their niches if they possess the ability to deliver DNA into the cytoplasm. Our data suggest that NLRP3 is also involved in the activation of caspase-1 in *S. pneumoniae*-infected macrophages. The NLRP3 inflammasome is activated by a wide variety of stimuli. In particular, pore-forming toxins, including PLY, have been reported to activate the NLRP3 inflammasome, owing to their pore-forming activities by causing K+ efflux (11, 21, 23–26). Because cell membrane permeation by PLY leads to K+ efflux (55), and PLY-induced activation of the NLRP3 inflammasome is inhibited by adding KCi to the cultures (11), it is conceivable that the NLRP3 inflammasome activation induced postinfection with *S. pneumoniae* depends on K+ efflux caused by PLY. Consistent with the assumption, addition of 20 mM KCi to cell cultures resulted in a significant decrease in the secretion of IL-1β and IL-18 from WT macrophages infected with *S. pneumoniae* (data not shown). However, it was difficult to judge whether the effect of KCi was due to specific inhibition of the NLRP3 inflammasome, as the same concentration of KCi also decreased the cytokine responses of NLRP3−/− macrophages to *S. pneumoniae* (data not shown). Because the presence of the cathepsin B inhibitor CA-074-Me reduces the secretion of IL-1β from PLY-stimulated DCs (11), cathepsin B may also participate in the activation of the NLRP3 inflammasome in response to *S. pneumoniae*. If it is assumed that PLY released from phagocytosed *S. pneumoniae* damages the phagosomal membrane, *S. pneumoniae*-containing phagosomes may provide cathepsin B for the NLRP3 pathway. Based on our experimental results, we have developed a mechanistic model of caspase-1 activation in macrophages infected by PLY-producing *S. pneumoniae*. PLY is an important virulence factor of *S. pneumoniae* that promotes inflammation, bacterial survival in the host, and ultimately, pathogenicity of the pathogen. In contrast, PLY is required for the activation of ASC inflammasomes that are critical for host defense against pneumococcal pneumonia by means of inducing the secretion of pyroptotic cytokines. The findings of this study have revealed a pathophysiological link between a virulence factor, PLY, and the inflammasome.

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**Disclosures**

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