An AXL/LRP-1/RANBP9 complex mediates DC efferocytosis and antigen cross-presentation in vivo

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The phagocytosis of apoptotic cells (ACs), or efferocytosis, by DCs is critical for self-tolerance and host defense. Although many efferocytosis-associated receptors have been described in vitro, the functionality of these receptors in vivo has not been explored in depth. Using a spleen efferocytosis assay and targeted genetic deletion in mice, we identified a multiprotein complex — composed of the receptor tyrosine kinase AXL, LDL receptor–related protein–1 (LRP-1), and RAN-binding protein 9 (RANBP9) — that mediates DC efferocytosis and antigen cross-presentation. We found that AXL bound ACs, but required LRP-1 to trigger internalization, in murine CD8α+ DCs and human-derived DCs. AXL and LRP-1 did not interact directly, but relied on RANBP9, which bound both AXL and LRP-1, to form the complex. In a coculture model of antigen presentation, the AXL/LRP-1/RANBP9 complex was used by DCs to cross-present AC-associated antigens to T cells. Furthermore, a murine model of herpes simplex virus–1 infection, mice lacking DC-specific LRP-1, AXL, or RANBP9 had increased AC accumulation, defective viral antigen-specific CD8+ T cell activation, enhanced viral load, and decreased survival. The discovery of this multiprotein complex that mediates functionally important DC efferocytosis in vivo may have implications for future studies related to host defense and DC-based vaccines.

Introduction

Efferocytosis is the process by which apoptotic cells (ACs) are recognized and cleared by phagocytic cells (1, 2). The crucial role of this process is exemplified by the development of spontaneous autoimmune secondary to breakdown of self-tolerance in mice lacking certain efferocytosis receptors or their bridging molecules (3–6). Whereas macrophages are highly efficient at AC uptake (7) and are usually the focus of efferocytosis studies, efferocytosis by DCs can mediate the presentation of AC-associated antigens on MHC class I to naive and memory CD8+ T cells in a process called cross-presentation. This process has been proposed to induce either antigenic tolerance (8) or T cell priming against infectious agents (9), depending on the specific context. Mostly in vitro studies have identified a number of distinct efferocytosis receptors that can be used by DCs (7, 10, 11), suggestive of possible redundancy or tissue- or context-specific usage. However, the identity and mechanisms of receptors utilized by DCs to perform functionally important efferocytosis and cross-presentation in vivo constitutes a major gap in this important area of research.

Previous reports showed that the efferocytosis receptors CD36, αvβ5, and αvβ3 are not involved in antigen cross-presentation by DCs (12, 13). Bone marrow–derived DCs (BMDCs) from the AXL/TYRO3 double-KO mouse show defective efferocytosis in vitro (11), but the role of AXL/TYRO3 in DC efferocytosis in vivo and in cross-presentation is not known. An important study showed that T cell Ig domain and mucin domain 3 (TIM-3) participates in AC uptake by DCs and cross-presents a model antigen OVA to OT-I transgenic T cells (10). However, the relative contribution of TIM-3 in the context of other DC receptors and in functionally relevant cross-presentation is not known, especially because TIM-3 is also a negative regulator of T cell activation (14). Recently, scavenger receptor class F, member 1 (SCARF-1), was reported to be important for efferocytosis by DCs, and SCARF-1–deficient mice developed a lupus-like autoimmune disease (15). However, whether SCARF-1 plays a role in antigen cross-presentation or is utilized by human DCs was not explored. Given the critical role of DC efferocytosis and cross-presentation in maintaining tissue homeostasis, peripheral tolerance, and elicitation of antiviral immunity upon infection and vaccination, identification of the mechanisms and functionality of DC efferocytosis receptors in vivo is of paramount importance.

To address this critical gap in DC biology, we first used an in vivo spleen efferocytosis assay in which mice with absent or inhibited candidate DC efferocytosis receptors were injected with labeled ACs, followed by analysis of AC uptake by splenic DCs. Based on these initial findings, we carried out molecular-cellular studies and then gene-targeted mouse experiments to interrogate mechanism and function. This investigation led to the identification of a multiprotein efferocytosis complex that mediates DC efferocytosis in vivo. The proteins in the complex, AXL, LDL receptor–related protein–1 (LRP-1; also known as CD91), and
RAN-binding protein 9 (RANBP9), were unable to effect effrocytosis on their own, but rather functionally interacted to coordinate AC binding and internalization. Most importantly, these molecules played a substantial, nonredundant role in the cross-presentation of AC-associated antigens in vivo, including in clear-ance of a viral infection. Human splenic and monocyte-derived DCs also used these molecules for efferocytosis. Knowledge of the presence and functionality of this multiprotein complex in DC efferocytosis may have implications for future studies related to host defense and DC-based vaccines.

**Results**

**Splenic DCs utilize AXL, LRP-1, and TIM-3 for clearance of ACs in vivo and in vitro.** As a first approach to interrogating DC effrocytosis in vivo, we injected PKH67-labeled ACs into mice and then quantified the uptake of those ACs by splenic DCs using flow cytometry (16). To help validate this assay, we took advantage of the fact that mature splenic DCs, which can be increased in mice by injection with the Toll-like receptor activator CpG-DOTAP, and CD8α-splenic DCs have low effrocytosis efficiency (7, 17). The assay successfully identified DCs from CpG-treated mice and CD8α− DCs...
Figure 2
Screening for DC efferocytosis receptors in vivo. (A) In vivo splenic DC efferocytosis assay conducted 3 hours after injection of ACs in control mice and mice with absent or inhibited candidate efferocytosis receptors \((n = 3 \text{ per group})\). The \(y\) axis represents the proportion of ACs engulfed by splenic CD8\(^{\alpha+}\) DCs. LRP-1 was inhibited by i.v. injection of GST-RAP (2 mg/mouse), and TIM-3 was inhibited by i.p. injection of an anti–TIM-3 nAb (RMT3-23; 100 \(\mu g/mouse\)). (B and C) Flow cytometric quantification of the percentage of total DCs (CD11c\(^{hi}\)) and CD8\(^{\alpha+}\) DCs in the spleens of the indicated mice \((n = 3 \text{ per group})\). (D–F) Flow cytometric analysis of expression of AXL (D), LRP-1 (E), and TIM-3 (F) in CD8\(^{\alpha+}\) and CD8\(^{\alpha-}\) splenic DCs of WT mice. \(^*P < 0.05\) vs. respective control.
DCs but intact expression in splenic macrophages (Supplemental Figure 1B). CD8α+ splenic DCs in these mice showed significantly decreased efferocytosis compared with control Lrp1+/+ mice (Figure 2A), further validating a role for LRP-1 in DC efferocytosis in vivo.

Further studies showed that CD8α+ DCs, which had greater efferocytic efficiency than CD8α− DCs (Figure 1C), expressed substantially higher levels of AXL and LRP-1 compared with CD8α− DCs (Figure 2, D and E); this was also the case for another DC efferocytosis molecule, TIM-3 (Figure 2F and see below). In this context, the decrease in efferocytosis observed in the Axl−/− context, the decrease in efferocytosis observed in the Cd11cCre+Lrp1fl/fl mice was not due to changes in the number of total or CD8α+ DCs (Figure 2, B and C) or in the efferocytosis efficiency of splenic macrophages (Supplemental Figure 2). Furthermore, the spatial distribution of DCs within the splenic marginal zone, where most of the injected ACs traffic (21), was not different among WT, Axl−/−, and Cd11cCre+Lrp1fl/fl mice (data not shown).

We next explored 2 other molecules previously implicated in DC efferocytosis. Consistent with a previous report (10), injection of anti–TIM-3 neutralizing antibody (nAb) decreased efferocytosis by DCs in vivo (Figure 2A). An nAb against SCARF-1, another DC efferocytosis receptor blocking antibodies (Figure 3B). To demonstrate that the anti–SCARF-1 nAb was functional in human splenic DCs, we showed that it markedly decreased the uptake of one of SCARF-1’s ligands, acetylated LDL (Ac-LDL; Supplemental Figure 3A). Interestingly, AXL and LRP-1 were highly expressed on human splenic DCs, we showed that it markedly decreased the uptake of one of SCARF-1’s ligands, acetylated LDL (Ac-LDL; Supplemental Figure 3B). Thus, similar to our observations with AXL/LRP-1 and TIM-3 in mice, human monocyte-derived DCs and splenic DCs used AXL/LRP-1, but not SCARF-1, in conjunction with other, parallel receptor pathways to carry out efferocytosis.

AXL and LRP-1 cooperate to mediate efferocytosis. In view of our finding that the decreased DC efferocytosis in the spleens of Axl−/−, Cd11cCre+Lrp1fl/fl, and TIM-3–neutralized mice was only partial, we next sought to determine whether DC efferocytosis can be blocked further by various combinations of receptor inhibition. When Axl−/− mice were treated with an anti–TIM-3 nAb, the inhibition of splenic DC efferocytosis was additive (Figure 4A). This was not the case for CD36 or αV integrins (Supplemental Figure 4A), further supporting the data in Figure 2A suggesting these molecules are not involved in efferocytosis by murine DCs. Importantly, treatment of Axl−/− mice with the LRP-1 inhibitor GST-RAP did not result in any additional decrease in efferocytosis compared with each manipulation alone (Figure 4B), which suggests that LRP-1 and AXL function in the same pathway. In this context, it is important to note that expression levels of LRP-1 in Axl−/− DCs and AXL in LRP-1-deficient DCs were comparable to those in WT DCs (Supplemental Figure 4B).

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**Figure 3** Human monocyte-derived DCs and splenic DCs utilize AXL and LRP-1 for efferocytosis. (A and B) Human monocyte-derived DCs (MoDCs; n = 5; A) and splenic DCs (n = 7; B) were incubated with IgG control or the indicated combinations of nAbs against AXL, CD36, αV, or SCARF-1 (0.5 µg/ml) (R&D Systems) or the LRP-1 blocker GST-RAP (1 µg/ml). Cells were then exposed to PKH67-labeled apoptotic Jurkat cells (1:4 ratio) for 2 hours and assayed for efferocytosis by microscopic analysis. (C) Representative flow cytometric analysis of expression levels of AXL and LRP-1 in human CD141+ and CD141− splenic DCs (n = 5). *P < 0.05 vs. respective control; **P < 0.05 vs. all other groups (except as indicated).
To further explore the concept that AXL and LRP-1 might function in the same pathway, we considered the possibility that LRP-1 and AXL might carry out different steps in efferocytosis, such as AC recognition versus engulfment (24). To assess the role of AXL and LRP-1 in AC binding, we used the actin polymerization inhibitor cytochalasin D to prevent AC internalization. Whereas Axl<sup>-/-</sup> BMDCs were cleared deficient in AC binding using this assay, inhibiting LRP-1 using GST-RAP did not affect this process (Figure 4C). In contrast, when the assay was carried out in the absence of cytochalasin D to enable engulfment, both AXL deficiency and GST-RAP were inhibitory (Figure 4D). Similar results were obtained when the binding experiment was conducted at 4°C to block internalization (data not shown). Notably, blocking LRP-1 in Axl<sup>-/-</sup> BMDCs did not further augment the defect in AC engulfment (Figure 4D), consistent with the in vivo data in Figure 4B. These findings suggest a model in which AXL is involved primarily in AC recognition by DCs, whereas LRP-1 functions in engulfment of the bound ACs, but not AC binding per se.

**RANBP9 facilitates the functional interaction of AXL and LRP-1 in efferocytosis.** We reasoned that if AXL and LRP-1 functioned in a binding-internalization cooperative manner, they might be in close proximity or even physically interact with each other. To test this hypothesis, we first conducted double immunofluorescence confocal microscopy on BMDCs. Within the resolution of this technique, the images were suggestive of AXL/LRP-1 colocalization on the cell surface (Figure 5A). We also found that LRP-1 was expressed at a higher level than AXL in DCs (data not shown), which likely explains our finding that percent colocalization of AXL with LRP-1 was higher than that of LRP-1 with AXL (78% vs. 41%). To test our hypothesis using a complementary method, we determined whether AXL and LRP-1 could be coimmunoprecipitated in BMDCs. co-IP experiments showed the presence of LRP-1 in AXL IPs and the presence of AXL in LRP-1 IPs (Figure 5B). Notably, AXL was not detected in the IPs of LRP-1 in Axl<sup>-/-</sup> DCs (Supplemental Figure 5A), confirming the specificity of the antibodies used in these experiments. To determine whether AXL and LRP-1 interact in splenic DCs in vivo, we conducted a fluorescence resonance energy transfer (FRET) experiment in which primary antibodies recognizing AXL and LRP-1 were conjugated to Alexa Fluor 546 or Alexa Fluor 647 to act as FRET donor and acceptor, respectively. This assay was based on the principle that photons emitted by donor fluorophore can excite acceptor fluorophore when they are within 100Å of each other. A positive FRET is indicated by both an increase in fluorescence of the acceptor (sensitized emission) and a concomitant decrease in fluorescence of the donor (donor fluorescence quenching). FACS analysis of splenic DCs showed substantial FRET, as measured by analysis of both sensitized emission and donor fluorescence quenching (Figure 5, C and D). Moreover, consistent with our confocal data demonstrating greater colocalization of AXL with LRP-1, donor fluorescence quenching was greater when antibody against AXL was used as a donor (Figure 5D). As a control for this experiment, we showed that FRET was not observed in Axl<sup>-/-</sup> and LRP-1<sup>-/-</sup> deficient splenic DCs (Supplemental Figure 5B), thereby validating the fluorophore-conjugated antibodies used in this assay. In addition, FRET between fluorophore-conjugated antibodies against heavy and light chains of MHC class I, which are known to interact on the cell surface, was used as a positive control (Supplemental Figure 5C).

In order to determine the functional significance of AXL/LRP-1 interaction, it was necessary to identify the mechanism responsible for their association. Protein interaction algorithms suggested that LRP-1 and AXL do not directly bind each other, so we hypothesized the presence of a scaffolding protein. Further database searching indicated that the protein RANBP9 interacts with AXL in cancer cells (25) and with LRP-1 in neuronal cells (26). Although RANBP9 has not previously been implicated in any functional interaction between AXL and LRP-1 in efferocytosis, we considered the hypothesis that it can act as an AXL/LRP-1 scaffolding protein in DCs to couple AXL-mediated AC binding with LRP-1-mediated AC internalization.

We first probed the LRP-1 and AXL IPs for RANBP9 and found a positive immunoblot signal in both cases (Figure 5B), consistent with our scaffolding hypothesis. To test the functional significance of this interaction, we created bone marrow chimera mice using Ranbp9<sup>+/+</sup> donor cells, because Ranbp9<sup>+/+</sup> mice die in the early postnatal period (27). Thus, fetal liver cells from E15 Ranbp9<sup>+/+</sup> and WT embryos were transplanted into lethally irradiated C57BL/6J mice to generate chimeric mice in which BMDCs were deficient in Ranbp9. As confirmation, we verified that splenic DCs in the
The chimeric mice lost expression of RANBP9 compared with WT DCs (Supplemental Figure 6A). However, neither the expression level of AXL and LRP-1 in DCs nor the total number of CD8α+ DCs was affected by RANBP9 deficiency (Supplemental Figure 6, B–D). Consistent with our hypothesis, RANBP9 deficiency in DCs (assessed using BMDCs from Ranbp9fl/fl mice transduced with adenovirus) resulted in loss of FRET signal between AXL and LRP-1 (Figure 6A). Moreover, AXL failed to coimmunoprecipitate with LRP-1 in Ranbp9–/– DCs (Figure 6B). These data demonstrated that RANBP9 deficiency disrupted AXL/LRP-1 interaction. Most importantly, efferocytosis by splenic DCs was defective in Ranbp9–/– chimeric mice (Figure 6C). AC internalization after recognition by specific efferocytosis receptors involves actin cytoskeletal reorganization mediated by the GTP-bound form of Rac1 (28). Previous reports indicated that LRP-1 mediates AC engulfment through its adapter protein, engulfment adaptor PTB domain containing 1 (GULP1) (29), which facilitates the formation of Rac-GTP (30). Consistent with our hypothesis that LRP-1/GULP1 is primarily involved in engulfment of AXL-bound ACs, silencing GULP1 using siRNA in WT BMDCs (knockdown efficiency, >80%; data not shown) led to a significant decrease in AC engulfment, but AC binding was unaffected (Figure 6E). Furthermore, silencing GULP1 in Axl–/– BMDCs did not result in an additional decrease in AC engulfment compared with Axl–/– BMDCs transfected with negative control scrambled RNA (Supplemental Figure 7), which suggests that LRP-1/GULP1 by itself is unable to mediate AC recognition and internalization. These data support the concept that AXL and LRP-1/GULP1 function as a complex in which AXL binds to ACs, while LRP-1/GULP1 mediate subsequent engulfment.

Figure 5
AXL and LRP-1 interact in DCs in vitro and in vivo. (A) Representative immunofluorescent confocal microscopy of BMDCs stained at 4°C with anti–LRP-1 antibody followed by Alexa Fluor 488–labeled secondary antibody (green), and with anti-AXL antibody followed by Alexa Fluor 568–labeled secondary antibody (red). Points of AXL and LRP-1 colocalization are indicated by yellow staining in the merged image as well as in the mask of colocalization. Scale bar: 5 μm. (B) Endogenous AXL or LRP-1 was subjected to IP from whole-cell lysates of BMDCs, then probed by Western blotting for AXL, LRP-1, or RANBP9. Rabbit or rat IgG in the IP reaction was used as control. (C) Flow cytometric analysis of FRET (sensitized emission) in splenic DCs (CD11c+ gate) immunostained with fluorophore-conjugated antibody against AXL as donor and LRP-1 as acceptor (left) or vice versa (right). Other cells were stained singly with fluorophore-conjugated antibody against AXL (left) or LRP-1 (right) as donor. Cells stained with IgG conjugated to Alexa Fluor 546 were used as controls. (D) Donor fluorescence measurements in cells as in C. Data are representative of 3 independent experiments.
ACs and LRP-1/GULP1 mediates AC engulfment through actin cytoskeletal rearrangement.

AXL- and LRP-1–deficient DCs do not have defects in macrophagocytosis and phagocytosis. As a test of specificity for the role of the AXL/LRP-1/RANBP9 complex in DC efferocytosis, we next determined whether these molecules also play roles in the related processes of macropinocytosis, endocytosis, and phagocytosis. BMDCs obtained from WT, Axl–/–, or Cd11cCre+Lrp1fl/fl mice did not demonstrate any difference in the uptake of FITC-labeled dextran, a measure of macropinocytosis (Figure 7A). There was also no difference in the uptake of Dil-labeled LDL, a measure of receptor-mediated endocytosis (Figure 7B). Similarly, there was no difference in the ability of these DCs to phagocytose fluorescently labeled 1-μm latex beads (Figure 7C). Thus, AXL- and LRP-1–deficient DCs have a specific defect in efferocytosis.

The AXL/LRP-1/RANBP9 effero-

cytosis receptor complex is critical for cross-presentation of AC-derived antigens, both in vitro and in vivo. As mentioned above, one of the functional implications of DC efferocytosis is its ability to utilize AC-derived antigens for cross-presentation. We therefore tested whether ACs internalized via the AXL/LRP-1/RANBP9 complex could provide antigens for cross-presentation. We first used an in vitro model in which DCs from WT, Axl–/–, or Cd11cCre+Lrp1fl/fl mice were incubated with OVA-loaded ACs and then cocultured with CFSE-labeled CD8+ OT-I transgenic T cells. Both Axl–/– and Cd11cCre+Lrp1fl/fl DCs demonstrated decreased activation and proliferation of OT-I transgenic T cells (Figure 8A), indicative of defective antigen cross-presentation in the mutant DCs. Importantly, the Axl–/– and Cd11cCre+Lrp1fl/fl DCs were not intrinsically defective at cross-presentation, since they were robust in cross-presenting soluble as well as bead-conjugated OVA (Figure 8, B, and C).

The OVA assays were conducted in vitro using a model antigen, which is not directly relevant in vivo under physiologic or pathologic conditions. We therefore tested whether efferocytosis via the AXL/LRP-1/RANBP9 complex has relevance in antigen cross-presentation in a natural virus infection model. To achieve this, we used an experimental model of systemic herpes simplex virus–1 (HSV-1) infection in mice, because clearance of HSV-1 in vivo is dependent on successful cross-presentation and elicitation of a protective CD8+ T cell response (31). To monitor cross-presentation and activation of antigen-specific CD8+ T cells, we used mice that were adoptively transferred with CFSE-labeled CD8+ OT-I transgenic T cells. Both Axl–/– and Cd11cCre+Lrp1fl/fl DCs demonstrated decreased activation and proliferation of OT-I transgenic T cells (Figure 8A), indicative of defective antigen cross-presentation in the mutant DCs. Importantly, the Axl–/– and Cd11cCre+Lrp1fl/fl DCs were not intrinsically defective at cross-presentation, since they were robust in cross-presenting soluble as well as bead-conjugated OVA (Figure 8, B, and C).

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of nonproliferating gB498-505–specific CD8⁺ T cells in HSV-1–infected spleens (Figure 9A and Supplemental Figure 8A). Most importantly, in light of the role of CD8⁺ T cells in anti–HSV-1 immunity, all 3 KO models showed enhanced virus-induced pathogenesis, as demonstrated by adrenomegaly (Supplemental Figure 8B), massive adrenal infiltration of monocytes and neutrophils (data not shown), and viral titers more than 3-fold higher than in control mice (Figure 9B). Moreover, the adrenal glands of the mutant mice showed a significant increase in the number of TUNEL⁺ cells (Figure 9C), consistent with both increased HSV-1–induced apoptosis (33) and decreased efferocytic clearance of ACs. In theory, these data could be influenced by increased permissiveness of the KO cells to HSV-1 infection; however, HSV-1 infectivity was not increased in BMDCs lacking AXL or LRP-1 or in primary adrenal cells lacking AXL (Supplemental Figure 9). Furthermore, to directly relate these data to efferocytosis of HSV-1–killed cells, we rendered primary adrenal cells apoptotic by ex vivo infection with HSV-1 and then injected these ACs into WT and mutant mice. Consistent with the interpretation of the HSV-1 infection data, Axl⁻/- and Cd11cCre+Lrp1fl/fl mice demonstrated defective DC efferocytosis and decreased cross-presentation to gBT-I.1 CD8⁺ T cells (Supplemental Figure 10, A and B). Finally, to confirm and extend the biological significance of these findings, we compared the survival of WT and Cd11cCre+Lrp1fl/fl mice infected with a lethal dose of HSV-1 and found significantly decreased median survival in Cd11cCre+Lrp1fl/fl compared with WT mice (Cd11cCre+Lrp1fl/fl, 5.5 days; WT, 8.5 days; Figure 9D). Thus, despite the presence of multiple DC efferocytosis receptors, the AXL/LRP-1/RANBP9 efferocytosis complex was functionally important in antigen acquisition for cross-presentation and elicitation of a protective CD8⁺ cytotoxic T cell response in the setting of a natural virus infection.

A recent in vitro study demonstrated that enveloped viruses utilize DC TYRO3/AXL/MERTK (TAM) family receptors, including AXL, to promote viral replication via suppression of type I IFN responses (34). However, our in vivo data demonstrated that Axl⁻/- mice were more susceptible to HSV-1 (an enveloped virus) compared with WT mice on days 4, 7, and 10 after infection (Figure 9B and Supplemental Figure 11A). Moreover, the levels of IFN-α in serum were not significantly different between WT and Axl⁻/- mice (Supplemental Figure 11B). These data raise the interesting possibility that the pathogen and host may compete for the same cellular machinery. For example, during initial phases of infection, viruses may utilize TAM receptors on DCs to suppress an early IFN-α response and promote viral replication, while during later stages of infection, particularly when virus-induced ACs arise, DCs utilize the TAM receptor AXL to recognize ACs, including virus-infected ACs, triggering activation of CD8⁺ cytotoxic T cell response and clearance of virus.

Discussion

Efferocytosis by DCs serves several important functions, including maintenance of tolerance to self-antigens; initiation of immune response against virally infected cells; and, in theory, prevention of secondary necrosis of uncleared ACs (8). Numerous receptors have been implicated through in vitro and occasionally in vivo studies as being able to mediate AC uptake by a variety of phagocytic cell types (35). While these studies suggest that different types of phagocytes preferentially utilize specific sets of receptors to carry out efferocytosis, the role of AXL and LRP-1 in this process is less clear. Our data suggest that the AXL/LRP-1/RANBP9 efferocytosis complex is functionally important in antigen acquisition for cross-presentation and elicitation of a protective CD8⁺ cytotoxic T cell response in the setting of a natural virus infection.
out efferocytosis, there has been considerable uncertainty as to the receptors used by DCs for functionally important efferocytosis in vivo. The findings of the present study help clarify this critical area, with implications for understanding the roles of DC efferocytosis in physiology, pathophysiology, and possibly therapy, such as in DC-vaccine design (36).

AXL interacts with ACs through the bridging molecule GAS6, which binds both AXL and phosphatidylserine exposed on the surface of ACs (37). However, the mechanism by which AXL mediates AC engulfment was not previously elucidated. Here, we demonstrated that in murine DCs, AXL primarily acted as an AC recognition receptor that bound ACs, but did not internalize them. Engulfment of the bound ACs required RANBP9-mediated interaction of AXL with the LRP-1/GULP1 module. This mode of action raises the possibility of the tether-and-tickle hypothesis proposed by Henson and colleagues, in which macrophages were shown to bind ACs through one of a number of receptors, but internalization required another molecule, referred to as PSR (24). In DCs, AXL is an AC tether, and it is possible that this process somehow “ticks” the LRP-1/GULP complex to mediate cytoskeletal rearrangement and AC internalization. However, further studies will be needed to define if and how AC binding trig-

Figure 9
RANBP9, AXL, and LRP-1 play important roles in cross-presentation in an HSV-1 infection model. (A) Mice were infected i.v. with HSV-1 strain KOS (4 × 10⁵ pfu/mouse). All mice also received 10⁶ CFSE-labeled CD8⁺ T cells purified from spleens of gBT-1 transgenic mice. 4 days after infection, spleens of infected mice were analyzed for transgenic T cell proliferation by flow cytometric analysis of CFSE⁺CD8⁺ gated T cells. Data show percent undivided CFSE-labeled gBT-1 transgenic T cells. (B) HSV-1 titers in mouse adrenal glands 4 days after infection with HSV-1. (C) Representative microscopic images and quantification (n = 4 mice per group) of TUNEL staining in mouse adrenal glands 4 days after infection with HSV-1. Scale bars: 500 μm. (D) Kaplan-Meier survival curve analysis of WT Lrp1fl/fl mice and Cd11cCre+Lrp1fl/fl mice infected with a lethal dose of HSV-1 (2 × 10⁶ pfu/mouse). Median survival of WT mice was 8.5 days, compared with 5.5 days for Cd11cCre⁺Lrp1fl/+ mice (P < 0.05; n = 10 per group). *P < 0.05 vs. respective control.
Efferocytosis prevents secondary necrosis of ACs, and defects in efferocytosis have been linked to pathological tissue necrosis, such as that in the necrotic cores of advanced, clinically dangerous atherosclerotic lesions (47). For example, deletion of the macrophage efferocytosis receptors MERTK and LRP-1 have been demonstrated to exacerbate necrotic core formation in mouse models of advanced atherosclerosis (19, 48, 49). Interestingly, observational studies have shown that the necrotic cores of clinically dangerous human atheromata are often surrounded by mature DCs, which are relatively inefficient effecocytes that can constitute up to 30% of the phagocyte population of atherosclerotic lesions (7, 50, 51). Thus, pathologic accumulation of pools of efferocytosis-inefficient mature DCs may promote focal necrosis. However, testing of this hypothesis in atherosclerosis and in other pathological settings, such as in tumor immunosurveillance, has been hampered by lack of knowledge of DC-specific efferocytosis receptors (2, 52). Here, we found that neither TIM-3 neutralization nor RANBP9 deficiency blocked efferocytosis by macrophages in vivo (Figure 2 and Supplemental Figure 6E); thus, targeting these molecules may provide a strategy by which to interrogate the role of DC-specific efferocytosis in these settings. Moreover, given the critical role of efferocytosis and antigen cross-presentation in suppressing autoimmune responses, future investigation of the role of the AXL/LRP-1/RANBP9 complex in this area is warranted.

In summary, we characterized a new multiprotein complex utilized by DCs for efferocytosis and antigen cross-presentation in vivo and demonstrated a novel role for the scaffolding protein RANBP9 in facilitating DC efferocytosis. In addition to providing new insight and tools with which to study the physiologic and pathologic roles of DC efferocytosis, these findings may have implications for future DC-based immunotherapeutic strategies. For example, proteins or peptides packaged in AC-mimetic nanoparticles targeted to the AXL/LRP-1/RANBP9 complex may enhance efficiency of antigen cross-presentation for the purpose of eliciting a desired immune response against cancer or pathogens or a tolerogenic response in the setting of autoimmunity.

Methods

Animals and animal maintenance. Cd11b−/− and OT-1 mice were obtained from Jackson Laboratories (stock nos. 003991 and 003831, respectively). Axl−/−, Tyro3−/−, Axl−/−, and Merk−− mice were generated as described previously (53, 54). Lrp1fl/fl mice (strain name B6.129S7-Lrp1tm2Her/J; JAX stock no. 012604) were generated as described previously (55) and then bred with Cd11cCre−/− mice (Jackson Laboratories; stock no. 008068) to generate Cd11cCre/Lrp1fl/fl mice. Ranbp9−/− embryos were harvested from pregnant mice obtained from Ranbp9+/− mating (27). Ranbp9fl/fl mice, with exon 3 flanked by loxP sites, were generated using a targeting vector consisting of an FRT site–flanked β-neo cassette with loxP sites inserted upstream and downstream of exon 3 of the Ranbp9 gene. This construct was electrophoresed into JMRI A3 F1-derived ES cells. An ES cell clone (HEPD0592.3.D06) that had successfully undergone FLP-mediated recombination and no longer retained the cassette, but did retain the loxP-flanked exon 3, was obtained from The European Conditional Mouse Mutagenesis Program and injected in to C57BL/6 blastocysts. Male chimeric mice were then bred to C57BL/6 female mice for germline transmission. Male and female heterozygous Ranbp9+/− mice were bred with each other to generate homozygous Ranbp9−/− mice, which were used as a source of BMDCs for this study. gBT-I.1 mice were provided by D. Carr (University of Oklahoma Health Sciences Center, Oklahoma City, Oklahoma, USA).
**Primary culture of BMDCs.** Murine bone marrow cells were isolated from the femurs of mice, cultured for 7 days in petri dishes in DMEM containing 10% FBS, and supplemented daily with GM-CSF (10 ng/ml) and IL-4 (10 ng/ml). On day 3, half the medium was replaced to replenish nutrients. The nonadherent cells were harvested on day 7, and experiments were performed by plating these cells on non-tissue culture–treated 24-well plates. To delete RANBP9 in BMDCs generated from the bone marrow of Ranbp9−/− mice, cells were transduced with an adenoviral vector encoding Cre or LacZ control at an MOI of 200:1 (virus/DC), and experiments were conducted 48 hours later. Adeno-Cre and adeno-LacZ were obtained from Viraquest.

**Generation of human peripheral blood monocyte-derived DCs.** Buffy coats were obtained from New York Blood Center. The mononuclear cell layer was isolated by gradient centrifugation using Ficoll-Hypaque, then plated and incubated in tissue culture dishes. After 1 hour, the nonadherent cells were washed away. The medium (DMEM containing 10% FBS) was supplemented with 500 IU recombinant human GM-CSF and IL-4 on days 0 and 3 of culture. The nonadherent cell population on day 7 was used as DCs.

**Human splenic DC isolation.** Human splenocyte samples were maintained in cold saline and brought to the laboratory within 6 hours of organ procurement. In order to ensure high live cell yield, samples were rapidly processed through both enzymatic and mechanical digestion methods. Spleen was cleaned of fat, chopped into small pieces, and placed into 50-mL conical tubes. Enzymatic digestion media (25 mL) was added (RPMI containing 10% FBS, 1-glutamate, sodium pyruvate, nonessential amino acids, penicillin-streptomycin, 1 mg/mL collagenase D, 1 mg/mL trypsin inhibitor, and 0.1 mg/mL DNase I) and incubated with mechanical shaking at 37°C for 1 hour. Digested tissue was then disrupted using the gentleMACS tissue dissociator (Milenyi Biotec), and the resulting suspension was passed through a stainless steel tissue sieve (mesh size, 10–150), followed by centrifugation. Red blood cell lysis was performed by addition of 5 mL ACK lysing solution (BioWhittaker) directly to the cell pellet and incubation with periodic agitation for 5 minutes. To remove dead cells and debris, cell pellets were resuspended in 20 mL of 30% percoll (GE Healthcare Life Sciences) and centrifuged at 50 g for 12 minutes. The resulting top layer and bottom pellet were discarded. Cells were washed with complete RPMI to remove percoll, and the resulting cell suspension was passed through a 40-μm filter. DCs were obtained from this cell suspension by negative selection using human myeloid DC isolation kit (Milenyi Biotec).

**Generation of fluorescently labeled ACs.** Mouse splenocytes (in vivo experiments) or Jurkat cells (in vitro experiments) were labeled with PKH67-GL (2 μM; Sigma-Aldrich) per the manufacturer’s instructions. Cells were then exposed to ultraviolet light (254 nm, UVP) for 5 minutes for induction of apoptosis. After UV exposure, cells were incubated in a 37°C incubator exposed to ultraviolet light (254 nm, UVP) for 5 minutes for induction of apoptosis. The nonadherent cells were harvested on day 7, and experiments were performed by plating these cells on non–tissue culture–treated 24-well plates. To delete RANBP9 in BMDCs generated from the bone marrow of Ranbp9−/− mice, cells were transduced with an adenoviral vector encoding Cre or LacZ control at an MOI of 200:1 (virus/DC), and experiments were conducted 48 hours later. Adeno-Cre and adeno-LacZ were obtained from Viraquest.

**FRET assay.** Antibodies against AXL and LRP-1 (R&D Systems and Sigma-Aldrich, respectively) were conjugated to Alexa Fluor 546 or Alexa Fluor 647 using a commercially available antibody labeling kit (Invitrogen). The labeling ratio was approximately 3, calculated according to the manufacturer’s instructions. Antibody labeling of cells was performed at 4°C for 30 minutes. FRET analysis was conducted by flow cytometry on CD11c+ gated cells using a BD FACS Calibur equipped with 488- and 635-nm lasers. Data were analyzed using FlowJo software.

**IP and Western blotting.** IP was performed by lysing in vitro–cultured DCs using an IP buffer containing 20 mM Tris-HCl (pH 7.5), 150 mM NaCl, 1 mM MgCl2, 1 mM CaCl2, 1% Triton X-100, and 1 tablet of Roche EDTA-free protease inhibitor per 10 ml of buffer. Lysates were then incubated with protein A–sepharose beads (GE Healthcare) and 100 μg anti-AXL (Cell Signaling Technology) or anti–LRP-1 (Sigma-Aldrich) antibody overnight at 4°C on a rotor-shaker. Samples were centrifuged at 90 g for 5 minutes, and beads were washed and resuspended 3 times with IP buffer. Laemmli buffer was then added to the beads; after boiling, the suspensions were centrifuged at 9,000 g. The supernatant fractions were loaded onto 4%–20% Tris-glycine SDS-PAGE gels. After transfer onto nitrocellulose membranes, blots were probed for AXL (R&D Systems), LRP-1 (Sigma-Aldrich), and RANBP9 (Epicomix) using specific antibodies, and the immunoreactive bands were visualized by enhanced chemiluminescence (Thermo Scientific).

**siRNA transfection.** BMDCs (5 × 10^6 cells/well of 24-well non–tissue culture–treated plates) were resuspended in serum-free DMEM. siRNA transfection complex was prepared by incubating 20 pmol Gulp1 siRNA (Qia-gen) with Lipofectamine 2000 (Invitrogen) (1:1 ratio) in serum-free DMEM for 20 minutes at room temperature. BMDCs were incubated with 20 pmol siRNA transfection complex under serum-free conditions for 6 hours at 37°C in a 5% CO2 incubator, after which the culture medium was supplemented with serum to a final concentration of 5%. Cells were harvested 24 hours after transfection; knockdown efficiency was analyzed by measuring Gulp1 mRNA by RT-qPCR.

**Fetal liver transplantation.** E15 embryos from pregnant mice of Ranbp9+/− matings were harvested. Portions of the tail and head were preserved for genotyping by PCR and Western blotting to identify Ranbp9−/− embryos. Under sterile conditions, the fetal livers were harvested, and cell suspensions were prepared by passing the liver tissue sequentially 3 times through a 20-gauge needle and 1 time through a 25-gauge needle. Cells were centrifuged at 130 g for 5 minutes, resuspended in serum-free DMEM, and counted using a hemocytometer. 2 × 10^6 cells were injected i.v. into lethally irradiated (10 Gy; Gammmacell40 unit) C57BL/6J recipient mice. Recipient mice were fed antibiotic water (100 μg/ml neomycin and 10 μg/ml polymyxin B) for 2 weeks and then acidified water. Chimeric mice were used 4 weeks after fetal liver transplantation.

**In vitro antigen cross-presentation assay.** Jurkat cells were incubated with 1 mg/ml OVA, and the OVA-loaded cells were exposed to UV light to induce apoptosis. BMDCs from appropriate groups of mice were incubated with OVA-loaded ACs for 2 hours to facilitate efferocytosis. FACS-sorted CD8+ T cells from OT-1 transgenic mice, which express TCR specific to OVA257–264, were labeled with 5 μM CFSE as described previously (56). These CFSE-labeled OT-1 T cells (5 × 10^4/well) were then cultured with BMDCs...
exposed to OVA-loaded ACs (10%/well) for 72 hours. Cells were stained with anti-CD8α (BD Biosciences) to specifically detect OT-1 T cells. CFSE dye dilution as a readout of T cell proliferation was analyzed by flow cytometry on CD8α− gated cells. 

HSV-1 culture, mouse infection, and virus titering. HSV-1 strain KOS was purchased from ATCC and propagated by infecting Vero cells (ATCC) as described previously (57). Mouse infection was conducted by i.v. injection of 4 × 10^6 pfu/mouse (self-limiting dose) or 2 × 10^6 pfu/mouse (lethal dose) of HSV-1 in an ABSL2 facility. For experiments assaying antigen cross-presentation, mice were also infected with 1 × 10^6 CFSE-labeled gBTL1 transgenic CD8+ T cells. Antigen cross-presentation was analyzed by measuring CFSE dye dilution on CD8+ CFSE−-gated cells on a flow cytometer. To estimate the virus burden in tissues of infected mice, homogenized tissue samples were sonicated, and the resultant supernatant fraction was used to conduct virus titer analysis. Briefly, supernatants at several dilutions were added to confluent Vero cells grown in 24-cm2 tissue culture flasks and incubated for 2 hours. After washing away free virus, 7.5 μg/ml pooled human Ig (Sigma-Aldrich) was added to prevent reinitiation of extracellular virus particles. The flasks were incubated at 37°C for 2 days, and then plates were visualized by Giemsa staining and counted.

Primary adrenal cell culture and induction of apoptosis by HSV-1 infection. Primary adrenal cells were cultured as described previously (58). Briefly, mice were euthanized, and both adrenal glands were isolated, minced, and digested in HBSS containing 2 mg/ml collagenase I, 0.05 mg/ml DNase I, and 5 mg/ml bovine serum albumin. The isolated cells were cultured on tissue culture dishes in KO-DMEM (Invitrogen) supplemented with 15% KO serum replacement (Invitrogen) for 3 days. To induce adrenal cell apoptosis, cells were infected with HSV-1 strain KOS (MOI, 10 pfu/cell) for 2 hours. Cells were further cultured for 24 hours at 37°C (after which annexin V labeling revealed >65% of cells to be apoptotic), then UV-irradiated to inactivate the virus prior to injection in mice. 

Statistics. All data are expressed as mean ± SEM. Analysis of statistical significance was performed using 2-tailed Student’s t test; P value less than 0.05 was considered statistically significant.


41. Ishimoto Y, Ohashi K, Mizuno K, Nakano T. Prom-otion of the uptake of PS liposomes and apop-totic cells by a product of growth arrest-specific


