The dendritic cell receptor DNGR-1 controls endocytic handling of necrotic cell antigens to favor cross-priming of CTLs in virus-infected mice

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DNGR-1 (CLEC9A) is a receptor for necrotic cells required by DCs to cross-prime CTLs against dead cell antigens in mice. It is currently unknown how DNGR-1 couples dead cell recognition to cross-priming. Here we found that DNGR-1 did not mediate DC activation by dead cells but rather diverted necrotic cell cargo into a recycling endosomal compartment, favoring cross-presentation to CD8+ T cells. DNGR-1 regulated cross-priming in non-infectious settings such as immunization with antigen-bearing dead cells, as well as in highly immunogenic situations such as infection with herpes simplex virus type 1. Together, these results suggest that DNGR-1 is a dedicated receptor for cross-presentation of cell-associated antigens. Our work thus underscores the importance of cross-priming in immunity and indicates that antigenicity and adjuvanticity can be decoded by distinct innate immune receptors. The identification of specialized receptors that regulate antigenicity of virus-infected cells reveals determinants of antiviral immunity that might underlie the human response to infection and vaccination.

Introduction
DCs are central players in the establishment and maintenance of self-tolerance, as well as in the induction of immunity against invading microorganisms (1, 2). In mice and humans, distinct subsets of DCs can be variably defined by phenotype, ontogeny, and function (3–5). They include the CD8α+ DC subset found in mouse lymphoid organs and the related CD103+ DC subset in non-lymphoid tissues, both of which are selectively lost in animals lacking the transcription factor BATF3 (6, 7). Cells bearing a similar phenotype have recently been described in humans, humanized mice, and sheep, indicating cross-species conservation of the CD8α+ DC family (8–12). This extended family has distinct functional properties, most notably a remarkable efficiency at capturing material from dead or dying cells, as well as processing exogenous antigens for cross-presentation on MHC class I (8–11, 13–16). These two features allow CD8α+ DC to cross-present cell-associated antigens (17, 18) and trigger CTL responses against infectious agents or tumors (6, 19–24). In addition to priming CD8+ T cells, CD8α+ DCs have been implicated in the establishment of cross-tolerance to tissue-specific cell-associated antigens (18, 25–27).

The ability of CD8α+ DCs to either cross-prime or cross-tolerize CD8+ T cells against cell-associated antigens implies that they can decode the context in which they encounter dead cells. In the case of infection, priming may be favored by signals from pattern recognition receptors (PRRs) that recognize bacterial or viral signatures associated with the dying infected cell (28, 29). In the case of uninfected cells such as tumors or allografts, it has been argued that the type of cell death dictates immunogenicity versus tolerogenicity, with necrosis and some forms of apoptosis favoring the former (30–32). This notion implies that CD8α+ DCs possess receptors that couple innate sensing of necrotic cells to the induction of adaptive immunity. In this regard, we have identified DNGR-1, also known as CLEC9A, as a receptor for necrotic cells that favors cross-priming of CTLs to dead cell–associated antigens in mice (33). DNGR-1 is selectively expressed at high levels by mouse CD8α+ DCs (34, 35) and CD103+ DCs (36) and by their human equivalents (9, 37) and recognizes an intracellular ligand that is only exposed after cell death (33). Notably, loss-of-function experiments in the mouse showed that DNGR-1 is not required for dead cell uptake but plays a non-redundant role in priming CD8+ T cells against antigens contained within necrotic cells (33). However, it is currently unknown how DNGR-1 couples dead cell recognition to cross-priming and whether it only controls immunity to cell-associated antigens or is also involved in cross-tolerance.

DNGR-1 is a type II transmembrane C-type lectin receptor (CLR) bearing an extracellular C-type lectin-like domain (CTLD) and a cytoplasmic tail with a hemi–immunoreceptor tyrosine-based activation motif (hemiITAM) motif that promotes signaling via spleen tyrosine kinase (Syk) (34, 37). The related CLR, Dectin-1, contains a similar hemiITAM motif and functions as an activatory receptor in myeloid cells, promoting NF-κB activation and proinflammatory cytokine production in response to engagement by agonist ligands (38, 39). Reflecting the structural similarity between the two receptors, DNGR-1 has been proposed to also act as an activatory CLR (37). Therefore, it is plausible that interaction of DNGR-1 with necrotic cell cargo signals DNGR-1 to activate DCs and favor cross-priming. Here, we show that DNGR-1 is a receptor for necrotic cells required by DCs to cross-prime CTLs against death-induced cell antigens in mice.
with its ligand on dead cells leads to activation of CD8α+ DCs, thereby enhancing their ability to prime naïve CD8+ T cells. Consistent with this notion, the function of DNGR-1 in cross-priming to dead cell–associated antigens is dependent on the integrity of the DNGR-1 hemITAM motif and requires Syk kinase (33).

Here, we analyze the role of DNGR-1 in immunity to dead cells and the ability of the receptor to function as a DC activatory receptor. We find that DNGR-1 specifically controls cross-priming, but not cross-tolerance, to cell-associated antigens. However, contrary to expectations, we found that DNGR-1 does not mediate activation of DCs by dead cells. Rather, we show that DNGR-1 has a dedicated function in endocytic handling of necrotic cargo that results in diversion from a degradative fate and favors MHC class I cross-presentation to CD8+ T cells. Consistent with that notion, we show that DNGR-1 deficiency impairs CTL priming even in highly immunogenic situations such as upon infection with herpes simplex virus (HSV) type 1. Thus, DNGR-1 is a dedicated DC receptor for cross-presentation of dead cell–associated antigens that can impact CTL responses to virus infection in mice and may therefore regulate immunity in humans.

Results

DNGR-1–dependent cross-priming of CTLs against dead cell–associated antigens. The conclusion that DNGR-1 is required for efficient cross-priming of naïve CD8+ T cells against antigens borne by dead cells rests on the observation of decreased expansion of antigen-specific CD8+ T cells and impaired development of CTL activity when antigen-bearing dead cells are injected i.v. into DNGR-1–deficient (Clec9a<sup>gfp/gfp</sup>) mice (33). To confirm that this protocol results in immunity, we injected mice with UV-treated OVA-MEFs 1 day later. Spleens were analyzed 5 days after immunization for OT-I proliferation. CFSE dilution (left panel) and total number of divided cells (right panel) are shown; each dot represents 1 mouse. (D) DNGR-1–sufficient or DNGR-1–deficient RIP-mOVA mice received CFSE-labeled CD45.1+ OT-I cells. Renal LNs were analyzed 3 days after injection for OT-I proliferation quantified by CFSE dilution (left panel) and total number of divided cells (right panel); each dot represents 1 mouse. *P < 0.05, **P < 0.01, ***P < 0.001, unpaired Student's t test and 1-way ANOVA (Tukey's post-test).
cantly lower in DNGR-1−deficient mice (Figure 1B). Even when precursor frequencies of antigen-specific CD8+ T cells were artificially increased by adoptive transfer of TCR-transgenic CD8+ T cells (OT-I), i.e. injection of necrotic OVA-MEFs promoted more robust clonal expansion of OVA-specific T cells in WT mice than in DNGR-1−null mice as measured by CFSE dilution and by total cell counts (Figure 1C). Overall, these results confirm and extend our previous observations that immunization with necrotic cells carrying a foreign antigen results in bona fide cross-priming of CTLs, which is partially dependent on DNGR-1.

Cross-tolerance of CD8+ T cells against dead cell–associated antigens is DNGR-1 independent. To test whether DNGR-1 might additionally be involved in promoting cross-tolerance to cell-associated antigens, we crossed DNGR-1−deficient mice with RIP-mOVA expressing B3Z (B3Z.Syk) cells, a Syk-transduced T cell–derived cell line expressing or overexpressing DNGR-1. This response was absent in B3Z.Syk cells expressing a Y7F mutant DNGR-1 (33), indicating that it was hemITAM dependent (Supplemental Figure 4). These results demonstrate that recognition of dead cells by DNGR-1 can induce gene expression and cell activation via hemITAM signaling. However, this response was observed in lymphoid-derived cells ectopically expressing DNGR-1 but absent in myeloid cells constitutively expressing or overexpressing DNGR-1.

DNGR-1 does not act as a myeloid activatory CLR. The inability of DNGR-1 to trigger myeloid cell activation in response to dead cells could result from nullification by simultaneous engagement of inhibitory receptors by the corpses (45). To selectively trigger only DNGR-1, we tried receptor cross-linking with mAbs but failed to detect signs of CD8α+ DC activation (Supplemental Figure 5), as reported by others (46). Therefore, we adopted an approach used for human DNGR-1 (37) and generated a chimeric receptor comprising the cytoplasmic tail of mouse DNGR-1 fused to the transmembrane domain and extracellular domain of mouse Dectin-1 (Figure 2A). The chimera can be uniquely triggered using a pure β-glucan, curdlan (39), thereby circumventing the need for a specific DNGR-1 agonist and obviating the possibility of additional receptor engagement by dead cells.

In order to test for proper folding and signaling capacity of the chimeric receptor, we first expressed it in B3Z.Syk cells. Surface expression of the receptors was monitored with an anti–Dectin-1 mAb and was comparable in chimera- and Dectin-1–expressing cells (Supplemental Figure 6A). In addition, curdlan stimulation promoted NFAT activation and IL-2 and TNF-α secretion by B3Z.Syk cells transduced with retroviral vectors encoding either Dectin-1 or the chimeric receptor, but not by cells transduced with an empty vector or a vector encoding a chimeric receptor with a tyrosine to phenylalanine mutation (Y7F) in the hemITAM (Supplemental Figure 6, A and C). Similar results were obtained with LK cells (Supplemental Figure 6B), indicating that the chimeric receptor is able to recognize curdlan and signal for cytokine production with efficiency comparable to that of WT Dectin-1 in lymphoid cells.

We next compared Dectin-1−deficient GM-CSF–grown BMDCs retrovirally transduced to express either Dectin-1 or the chimeric receptor. The retroviral constructs also encoded GFP, which allowed us to sort the transduced cells based on different GFP levels and compare only those with matching levels of surface receptor expression (Figure 2B). Curdlan promoted robust production of TNF-α, IL-2, or IL-10 by DCs transduced with Dectin-1–encoding retrovirus but not by DCs transduced with empty control retrovirus encoding only GFP (Figure 2C). In stark contrast, curdlan-induced cytokine production was undetectable or greatly reduced in DCs expressing the chimeric receptor (Figure 2C). A control stimulus, CpG, induced comparable levels of cytokine production by all 3 populations of transduced DCs (Supplemental Figure 7).

Similar results were obtained when we used a different chimeric receptor consisting of both the cytoplasmic tail and the transmembrane domain of mouse DNGR-1 fused to the extracellular domain of Dectin-1 (Supplemental Figure 8). These results suggest that, compared with that of Dectin-1, the tail of DNGR-1 is markedly deficient in its ability to signal for cytokine production in myeloid cells. This conclusion appears to be in contrast to that in a previous report in which RAW264.7 macrophages transduced with an analogous chimeric receptor of human DNGR-1 and mouse
Dectin-1 secreted significantly more TNF-α than control cells upon stimulation with zymosan particles (37). Importantly, curdlan specifically triggers Dectin-1, while zymosan can trigger other PRRs such as TLR2 and Dectin-2. It is therefore conceivable that binding of the particles to the chimeric receptor facilitated their recognition by other zymosan-specific receptors, which in turn signaled for TNF-α secretion. To address these possibilities, we procured the cells used in that study (37) and compared their response to zymosan versus curdlan. All cells expressed comparable levels of surface receptor and bound zymosan particles with similar efficiency (Supplemental Figure 9). As reported (37), zymosan induced increased TNF-α production in cells overexpressing either the chimera or Dectin-1 when compared with control cells transduced with an empty vector (Supplemental Figure 9). However, curdlan led to enhanced TNF-α production only in Dectin-1–transduced cells and not in cells expressing the chimera (Supplemental Figure 9). These results are concordant with our earlier conclusion that signaling by the cytoplasmic tail of DNGR-1 by itself is markedly inefficient at promoting activation of myeloid cells.

Examination of the data revealed a very low level of curdlan-induced cytokines in DCs expressing the Dectin-1/DNGR-1 chimera but not in cells transduced with empty vector (Figure 2C and Supplemental Figure 8). This appeared to depend on weak signal-
The receptor, as it was not seen with the Y7F mutant (Figure 2C), despite equivalent surface expression of all chimeras (Figure 2B). A DEDG sequence immediately upstream of the hemITAM tyrosine and shared by Dectin-1 and CLEC-2 but not by DNGR-1 has been shown to contribute to signaling by CLEC-2 (47). In DNGR-1 from all species, the glycine was replaced by an isoleucine and, in mouse but not human DNGR-1, it was preceded by two rather than three acidic residues (Figure 2A). We mutated these residues in the Dectin-1/DNGR-1 chimeras to determine whether restoration of a Dectin-1–like DEDG sequence could potentiate the weak signaling by DNGR-1. All chimeras were expressed at similar levels in Dectin-1–deficient BMDCs and signaled to NFAT with comparable efficiency when expressed in B3Z.Syk cells (Figure 2B and Supplemental Figure 6C). However, restoration of the tri-acidic motif had no effect on DC activation by curdlan (Figure 2C). We conclude that the DNGR-1 hemITAM can potentially signal to induce myeloid cell activation but this ability is constrained by the presence of an isoleucine residue immediately upstream of the tyrosine. DNGR-1 deficiency does not impact the adjuvanticity of dead cells. Despite the failure to lead to overt DC activation, weak DNGR-1 signaling could still mediate a subtle change in CD8α+ DCs that would lead to an improved ability to stimulate T cells in response to dead cells and account for the role of the receptor in cross-priming. To test for such an effect, we pulsed CD8α+–like Flt3L-derived BMDCs with a limiting concentration of OVA peptide (SIINFEKL) and subsequently allowed them to interact with necrotic MEFs (lacking OVA) before adding naive CFSE-labeled OT-I cells and measuring T cell proliferation. Notably, the expansion of OT-I cells was clearly improved as a function of the ratio of dead cells to DCs (Figure 3A), confirming that dead cell–derived signals improve the stimulatory ability of CD8α+ DCs for naive CD8+ T cells. However, this “adjuvant” effect of dead cells was independent of DNGR-1, as it was seen equally with WT and DNGR-1–deficient DCs (Figure 3A). These results suggest that DNGR-1 does not play a role in DC activation by dead cells even when the latter is measured with a T cell assay that presumably integrates known and unknown parameters of DC activation.

**Figure 3**

DNGR-1 regulates MHC class I cross-presentation but not MHC class II presentation of dead cell–associated antigens. (A) The adjuvanticity of dead cells is independent of DNGR-1. Purified CD8α+–like Flt3L-BMDCs from WT or DNGR-1–deficient (Clec9agfp/gfp) mice were pulsed with 1 pM SIINFEKL peptide, washed, and cultured with UV-treated H-2bm1 MEFs and CFSE-labeled OT-I T cells. OT-I proliferation was quantified after 3 days. (B and C) DNGR-1 regulates cross-presentation of dead cell–associated antigens. Purified CD8α+–like Flt3L-BMDCs from WT or DNGR-1–deficient mice were cultured with UV-treated OVA-MEFs or latex beads coated with OVA (OVA beads). Proliferation of CFSE-labeled OT-I (B) or OT-II (C) T cells was quantified on day 3 or 5 (respectively) of coculture with DCs. (D) Purified CD8α+–like Flt3L-BMDCs from WT, DNGR-1–deficient, or Tap1+/– mice were cultured with UV-treated OVA-MEFs and B3Z cells. NFAT reporter activity in B3Z cells was measured using a colorimetric assay after 24 hours of coculture. Results are mean ± SD and representative of at least 3 independent experiments. *P < 0.05, **P < 0.01, unpaired Student’s t test.
DNGR-1 diverts cargo to the early/recycling endosomal pool. (A) Purified splenic CD8α+ DCs from WT mice were cultured for 30 minutes (EEA-1) or for 90 minutes (Rab5a and Rab27a) with Alexa Fluor 488–coupled anti–DNGR-1 Ab. Cells were fixed and stained for confocal microscopy. Distribution of internalized anti–DNGR-1 Ab+ vesicles relative to EEA-1, Rab5a+, or Rab27a+ vesicles was quantified as percentage of vesicles positive for anti–DNGR-1 antibody also positive for endosomal markers in 15 cells per condition. (B and C) Purified CD8α+–like FcγRII-BMDCs from WT or DNGR-1–deficient mice were cultured with UV-treated Alexa Fluor 647–labeled H-2Kb-MEFs. Cells were fixed after 60 or 240 minutes and stained for confocal microscopy. Distribution of cargo derived from dead cells relative to Rab5a (B) and Rab11 (C) was quantified as percentage of total vesicles positive for endosomal markers containing dead cell cargo in 15 cells per condition. Scale bars: 5 μm. Results are representative of 2 independent experiments. *P < 0.05, **P < 0.01, unpaired Student’s t test.

DNGR-1 diverts phagocytosed dead cell cargo to a non-degradative recycling endosomal compartment. To extend these findings to the context of real virus infection in vivo, we tested co-localization with Rab11, a marker for recycling endosomes, which have also been implicated in antigen cross-presentation (32). At 60 minutes after uptake, neither WT nor DNGR-1–deficient DCs contained more than 10% of cargo-positive vesicles that were also Rab11+ (Figure 4C). However, at 240 minutes, approximately 50% of cargo-containing vesicles contained Rab11 in WT DCs. This co-localization with the recycling endosomal marker was significantly reduced in DCs lacking DNGR-1 (Figure 4C). Together, these data suggest that a proportion of phagocytosed dead cell–derived material associates with DNGR-1 and is diverted to the recycling endosomal route, thereby facilitating cross-presentation.

DNGR-1 is non-redundant for CTL responses to virus infection. To first test this notion in vitro, we used an OVA-expressing “suicide” cytopathic Semliki Forest virus (SFV) that infects cells and replicates to produce double-stranded RNA but no progeny viruses (28, 33). Phagocytosis of dead or dying SFV-OVA–infected cells by CD8α+ DCs leads to TLR3-dependent activation of the cells and drives robust cross-priming against OVA (28). Indeed, co-culture of SFV-OVA–infected H-2Kb–MEFs with H-2Kb CD8α+–like DCs led to marked CD40 and CD86 upregulation, as well as IL-6 and TNF-α production, all of which were markedly reduced in the absence of TLR3 (Figure 5A). Consistent with this defect in activation and/or an additional effect of TLR3 on cross-presentation, TLR3–/– CD8α+ DCs were also impaired in their ability to drive OT-I T cell expansion in response to SFV-OVA–infected cells (Figure 5B). Notably, neither upregulation of DC maturation markers nor the level of cytokines produced was affected by DNGR-1 deficiency (Figure 5A). However, the expansion of OT-I T cells primed by DNGR-1–deficient DCs was still significantly decreased (Figure 5B).

To extend these findings to the context of real virus infection in vivo, we tested co-localization with Rab11, a marker for recycling endosomes, which have also been implicated in antigen cross-presentation (32). At 60 minutes after uptake, neither WT nor DNGR-1–deficient DCs contained more than 10% of cargo-positive vesicles that were also Rab11+ (Figure 4C). However, at 240 minutes, approximately 50% of cargo-containing vesicles contained Rab11 in WT DCs. This co-localization with the recycling endosomal marker was significantly reduced in DCs lacking DNGR-1 (Figure 4C). Together, these data suggest that a proportion of phagocytosed dead cell–derived material associates with DNGR-1 and is diverted to the recycling endosomal route, thereby facilitating cross-presentation.

DNGR-1 diverts phagocytosed dead cell cargo to a non-degradative recycling endosomal compartment. To extend the above results, we tested whether DNGR-1 specifically functions by diverting necrotic cargo to a subcellular compartment that favors cross-presentation, as suggested for another CLR, the mannose receptor (50). We first asked whether DNGR-1 is directed to early endosomal compartments associated with antigen cross-presentation. Consistent with that hypothesis, anti–DNGR-1 mAb bound to the surface of CD8α+ DCs was selectively delivered to vesicles positive for EEA-1, Rab27a, or Rab5a, which mark such compartments (50, 51). Indeed, 75% of DNGR-1+ endosomes colocalized with EEA-1 at 30 minutes and with Rab27a or Rab5a at 60–90 minutes after internalization (Figure 4A). We then examined the fate of necrotic cargo taken up by CD8α+–like DCs. We found that DNGR-1+ vesicles containing dead cell–derived cargo colocalized with Rab5a. Indeed, at 60 minutes of coculture of WT DCs with Alexa Fluor 647–labeled necrotic MEFs, we detected Rab5a in approximately 60% of vesicles with dead cell–derived material (Figure 4B). This percentage decreased to approximately 20% at 240 minutes of coculture, indicating that the nature of DNGR-1+ vesicles changes at later time points after phagocytosis. Interestingly, co-localization of cargo with Rab5a was significantly reduced in DNGR-1–deficient DCs (Figure 4B), suggesting that DNGR-1 may actively determine the intracellular route along which the cargo is directed.
(gB) epitope in C57BL/6 mice were markedly reduced in Batf3−/− mice, indicating a major contribution of the CD8α+ DC family to CD8+ T cell priming (Figure 6A). Remarkably, a reduction of around 30%–80% in the CTL response to gB was also observed in DNGR-1−deficient animals, when measured as number or frequency of gB-specific CTLs (Figure 6A) or by examination of gB peptide-specific IFN-γ production by CD8+ T cells (Figure 6B). In contrast, there was no defect in anti–HSV-1 antibody titers in DNGR-1−deficient mice (Supplemental Figure 11), indicating that DNGR-1 selectively regulates CTL priming. Importantly, no reduction in HSV-1–specific CTL numbers or frequency was seen when DNGR-1−deficient mice were infected via the i.v. route (Supplemental Figure 12). This is
DNGR-1 contributes to the CTL response to HSV-1 lung infection. WT, DNGR-1–deficient, or Batf3–/– mice were infected with HSV-1 i.n. Mediastinal LNs were harvested 7 days later, and CTL responses were measured directly ex vivo or after in vitro restimulation. (A) Left panel: H-2Kb-gB pentamer+ cells as a percentage of CD8+CD44+ T lymphocytes. Right panel: Number of H-2Kb-gB pentamer+CD8+CD44+ T cells. (B) IFN-γ+ cells as a percentage of CD8+CD44+ T lymphocytes after restimulation with gB496-505 peptide for 6 hours. Data were pooled from 3 independent experiments, with each dot representing 1 mouse; naive WT, n = 5; WT, n = 13; DNGR-1–deficient, n = 13, Batf3–/–, n = 8 mice. *P < 0.05, **P < 0.01, ***P < 0.001, 1-way ANOVA (Tukey’s post-test).

Discussion

Antigens expressed by dead or dying cells can be made visible to the adaptive immune system, inducing either activation or inactivation of existing lymphocyte repertoires in many experimental models. However, the mechanisms by which innate recognition of cell death is translated into adaptive immunity or tolerance are not fully understood. CD8α+ DCs and the related CD103+ lineage are thought to be the major DC subsets mediating cross-priming and cross-tolerance to cell-associated antigens (3, 54, 55). We have shown previously that CD8α+ DCs utilize DNGR-1 to recognize a preformed signal exposed in necrotic cells and that ablation or blockade of the receptor reduces their ability to stimulate CTL responses against dead cell–associated antigens in vitro and in vivo (33). Those data and the previously reported function of DNGR-1 as a myeloid activatory CLR (36) suggested that DNGR-1 promotes cross-priming by activating DCs in response to contact with dead cells. Surprisingly, here we failed to find a role for DNGR-1 in promoting DC activation, which is probably regulated by other receptors (32). Rather, we suggest that the receptor acts to regulate the intracellular trafficking of internalized dead cell material, diverting it from a lysosomal proteolytic fate to a non-degradative recycling endosomal compartment that favors antigen extraction for TAP-dependent cross-presentation on MHC class I. We further show that the dedicated function of DNGR-1 in cross-presentation of dead cell–derived antigens contributes in a non-redundant manner to the overall CTL response to HSV-1 lung infection, indicating that it can be a key component of the immune response to cytopathic viruses.

The primacy of the CD8α+ DC family in cross-priming to dead cell–associated antigens has been put in question by a recent report suggesting that CD11c+CD169+ subcapsular sinus macrophages are the primary LN APC cross-presenting necrotic cell–derived antigens upon subcutaneous immunization with dead cells (56). Contrary to that report, we found that the CTL response to dead cell–associated antigens remains dependent on DNGR-1 and BATF3 even after subcutaneous immunization (S. Zelenay, B.U. Schraml, and C. Reis e Sousa, unpublished observations). Thus, cross-priming to dead cell–associated antigens remains dependent on DNGR-1 expressed by CD8α+ DC family members irrespective of administration route.

Coupling of recognition of dead cells to production of pro-inflammatory cytokines was previously reported for macrophage-inducible C-type lectin (Mincle) (57). This ITAM-coupled receptor drives production of TNF-α and MIP-2 and thereby mediates recruitment of neutrophils into the damaged tissue. Mincle and other innate immune receptors, which signal for cellular activation upon recognition of microbial or endogenous ligands, are widely expressed among immune cells. In contrast, DNGR-1 is selectively expressed at high levels by the CD8α+ DC subset, with lower levels of expression on plasmacytoid DCs (34). Moreover, the expression of DNGR-1 on CD8α+ DCs is not increased in vivo upon LPS administration in contrast to that of many CLRs with activating functions in myeloid cells (58). Therefore, DNGR-1 exhibits several properties that appear to be unique within the family of C-type lectins that function as innate immune receptors. One additional unique feature underscored in the present study is the apparent separation of cell death sensing from proinflammatory signaling.
Our conclusion that DNGR-1 does not act by promoting CD8α+ DC activation in response to cell corpses is supported by multiple lines of investigation. First, coculture with dead cells fails to promote DNGR-1-dependent cytokine production or DC maturation. This is the case when testing CD8α+ FLt3L BMDCs constitutively expressing DNGR-1 or GM-CSF–derived BMDCs and BM-derived macrophages ectopically expressing high levels of DNGR-1. Second, using DNGR-1/Dectin-1 receptor chimeras, we found that, unlike Dectin-1, DNGR-1 signaling only weakly induces production of proinflammatory cytokines by DCs. The chimera approach allows us to stimulate cells with a pure Dectin-1 agonist, avoiding the potential co-engagement of multiple agonistic and antagonistic receptors when using dead cells as a stimulus. Notably, the inability of DNGR-1 to signal for cytokine expression can be attributed to specific sequence features of its intracellular tail. Thus, a chimeric receptor consisting of Dectin-1 with the cytoplasmic tail of DNGR-1 was unable to signal for cytokine production, but conversion of a single isoleucine at position 6 into glycine markedly improved stimulatory potential. Third, DC activation triggered by phagocytosis of virus-infected cells was greatly diminished by loss of TLR3 but comparable in WT and DNGR-1–deficient DCs. This finding suggests that, independent of any ability to activate DCs autonomously, DNGR-1 also does not act synergistically with TLR to promote DC activation and further invalidates a model in which DNGR-1 acts by delivering infected cell cargo to TLR3-accessible endosomal compartments. Fourth, loss of DNGR-1 did not affect the ability of CD8α+ DCs to stimulate CD4+ T cells against dead cell–associated antigens or to stimulate CD8+ T cells in response to preprocessed antigen plus necrotic cells as “adjuvants.” Interestingly, DNGR-1 could drive activation in T and B cell–derived cell lines, but not in myeloid cells, such as Flt3L BMDCs, GM-CSF BMDCs, BM-derived macrophages, or RAW264.7 macrophages. GM-CSF BMDCs, in particular, are especially amenable to activation by the Dectin-1/Syk pathway, in part because of high expression of CARD9, which permits coupling to NF-kB (59). These data suggest that signaling downstream of Syk-coupled CLR, in particular to NF-kB, is not only cell context dependent (59) but is also controlled in a CLR-specific manner. Further studies will be required to determine the molecular basis of this regulation, which leaves open the possibility that DNGR-1 may still trigger a specific type of CD8α+ DC activation in some circumstances that are not mimicked by our present analyses. Nevertheless, collectively, our results do not support the notion that DNGR-1 is a receptor whose primary function is to activate DCs in response to contact with necrotic cells.

Surprisingly, our data highlight an underappreciated aspect of CLR function in regulating endocytic traffic and the retrieval and cross-presentation of dead cell–associated antigens. We have previously demonstrated that DNGR-1+ vesicles in CD8α+ DCs do not colocalize with a tracer that accumulates in lysosomes (33). We now find that the receptor and associated dead cell–derived cargo localize to vesicles bearing markers of early endosomes and the recycling endocytic route, indicating that DNGR-1 is confined to the recycling endosome pool. Mannose receptor–mediated uptake and transport of soluble OVA to Rab5a-positive vesicles has been previously reported to favor cross-presentation of OVA-derived peptides (50), and the recycling endosomal compartment is the most favorable point in the endosomal pathway for cross-presentation (60). This may be because recycling endosomes are poorly degradative relative to other endosomal compartments and/or because the recycling pathway intersects with a specialized intracellular compartment where acidification is limited by production of NOX2-dependent reactive oxygen species (61). In either case, the limited acidity and proteolytic activity allowed for preservation of intact antigens and translocation into the cytosol for TAP-dependent cross-presentation (61–64). In contrast, generation of peptides for presentation on MHC class II molecules does not require sequestration from degradation, and the presentation of dead cell–associated antigens to CD4+ T cells was not impaired in the absence of DNGR-1. DNGR-1 seems, therefore, to specifically direct dead cell–derived cargo toward a compartment where antigens can be preserved and retrieved for cross-presentation. As such, DNGR-1 may be a useful subcellular marker for identifying and characterizing the elusive “cross-presentation compartment” in CD8α+ DCs. Of course, our data do not exclude the possibility that DNGR-1 may additionally control the process of cross-presentation, for example, by promoting fusion with the ER–Golgi intermediate compartment (65) or by facilitating translocation of ligand to the cytosol, as recently shown for another myeloid CLR, the mannose receptor (66). Consistent with that notion, the Y7F hemITAM mutant DNGR-1 still localized to the recycling endosomal compartment even though it was impaired in its ability to promote cross-presentation of dead cell–associated antigens (unpublished observation). Further dissection of the role of DNGR-1 in the regulation of cross-presentation will be greatly facilitated by the identification of the DNGR-1 ligand (67).

Our findings indicate that DNGR-1 regulates cross-priming to dead cell–associated antigens by controlling the antigenicity rather than the adjuvanticity of dead cells. Similarly, it has previously been argued that recognition of HMGB-1 by TLR4 might primarily serve to regulate the ability of DCs to cross-present dead cell antigens (68). If DNGR-1 is involved in regulating cross-presentation rather than DC activation, it might seem odd that its function is apparent only in cross-priming conditions and not in a model of cross-tolerance to cell-associated antigens. This could simply reflect that, during physiological cell death in tissues such as pancreas, the dying cells are cleared before loss of membrane integrity occurs. As the ligands for DNGR-1 are intracellular (33), normal clearance of apoptotic cells during homeostatic conditions may not allow for engagement of the receptor. Even if apoptotic cells are sampled by CD8α+ DCs, digestion of the corpse with consequent ligand exposure is likely to occur only in late, degradative endocytic compartments from which, as shown here, DNGR-1 is excluded. Thus, we believe that DNGR-1 involvement is only likely to become apparent in a pathological setting rather than during steady-state physiological cell turnover.

To test for such involvement, we resorted to a model of cytopathic lung infection with HSV-1. Lung contains a prominent population of CD103+ DCs (69) that expresses DNGR-1 at high levels (36) and corresponds to the migratory DCs previously shown to ferry antigen to draining LNs following virus infection (21, 70). In addition, DNGR-1–expressing CD8α+ DCs in the draining mediastinal LNs can also be prominent APCs for virus-specific CD8+ T cells following lung infection (21). Therefore, the CD8α+ DC family is likely to be important in inducing CTL responses to respiratory viruses. Consistent with that notion, we found that the CTL response to HSV-1 antigens was markedly reduced in Batf3-deficient mice, which lack CD103+ and CD8α+ DCs. Importantly, this response was also reduced by an average of 50% in DNGR-1–deficient mice. HSV-1 lung infection is cleared independently of lymphocytes (71), and therefore, we did not see any increase in virus accumulation in...
the lungs of DNGR-1– or Batf3-deficient mice (data not shown). However, in a different model of vaccinia virus infection, DNGR-1 deficiency clearly impairs virus clearance (72). While we do not know whether the DNGR-1–dependent component reflects cross-priming by the CD8ε+ DC family or the activity of other cells (e.g., plasmacytoid DCs, which express low levels of the receptor), these findings nevertheless indicate that DNGR-1 can play a key role in the CTL response even under highly immunogenic situations such as virus infection. Our observations therefore suggest a novel and non-redundant point of control in immunity to infection, in which some DC receptors mark dead cells as cargo that contains antigens worthy of cross-presentation, whereas other receptors detect signs of infection in the corpses. As shown here, the former are necessary for cell-associated antigen processing and presentation. In contrast, the latter correspond to typical PRRs, which contribute to effective CTLs (LK cells) (74) were from ATCC (HB-98). OVA-expressing H-2 bm1 MEFs (bm1 MEFs) were previously described (37) and were provided by Gordon Brown. H-2 KααA3D mutant, and 5′-TTTCCCGTCGACATGCATGCGGAAGAAATATTATACCTCT-3′ for the Y7F-mutant. All constructs were verified by sequencing. BM cells were additionally supplemented with non-essential amino acids, HEPES, and sodium pyruvate (Invitrogen). Mouse BMDCs were generated using GM-CSF (GM-CSF–BMDCs) as described previously (73) or by culturing BM cells in the presence of 100–150 ng/ml Flt3L (R&D Systems) for 10 days to generate Flt3L–BMDCs (44). B3Z cells containing a reporter for costimulatory and other factors that promote effector T cell development. Manipulation of the pathways involved in dead cell sensing and antigen extraction may have wide implications for the design of vaccines designed to elicit CTLs in infectious disease or cancer immunotherapy. In addition, the identification of specialized receptors that regulate antigenicity of virus-infected cells reveals previously unappreciated determinants of antiviral immunity that might underlie the human response to infection and vaccination.

Methods
Mice. C57BL/6, Clec9aγf/f (DNGR-1–deficient), Clec7α−/− (Dectin-1–deficient; gift from Gordon Brown, University of Aberdeen, Aberdeen, United Kingdom), Batf3−/− (gift from Kenneth Murphy, Washington University, St. Louis, Missouri, USA), H2-Aα−/− (Aα−/−), Thrk−/−, Tap1−/−, B6.SJL, OT-II, OT-I x B6.SJL, OT-I x Rag1−/−, RIP-mOVA Clec9aγf/f, and RIP-mOVA Clec9aα−/− mice were bred at Cancer Research UK in specific pathogen–free conditions. Cells. Culture medium was RPMI 1640 supplemented with glutamine, penicillin, streptomycin, 2-mercaptoethanol (all from Invitrogen), and 10% heat-inactivated fetal calf serum (Source BioScience). For T cell cultures, medium was additionally supplemented with non-essential amino acids, heat-inactivated fetal calf serum (Source BioScience). For T cell cultures, medium was additionally supplemented with non-essential amino acids, HEPES, and sodium pyruvate (Invitrogen). Mouse BMDCs were generated using GM-CSF (GM-CSF–BMDCs) as described previously (73) or by culturing BM cells in the presence of 100–150 ng/ml Flt3L (R&D Systems) for 10 days to generate Flt3L–BMDCs (44). B3Z cells containing a reporter for NFAT coupled to LaZ activity have been previously described (49) and were from N. Shastri (University of California, Berkeley, California, USA). LK3.2 cells (LK cells) (74) were from ATCC (HB-98). OVA-expressing H-2Kαα MEFs (OVA-MEFs) have been described (33). RAW264.7 macrophages stably transduced with an empty retroviral vector or with a retroviral vector coding for the complete open reading frame of mouse Dectin-1 or for a chimeric human DNGR-1 mouse Dectin-1 were previously described (37) and were provided by Gordon Brown. H-2Kαα MEFs were infected with OVA-expressing SVF for 2–4 hours as previously described (28). UV-treated OVA-MEFs were generated by exposure to UV light (240 mJ/cm2), followed by overnight culture in complete medium to induce secondary necrosis, resulting in more than 85% dead cells. CD8ε+–like Flt3L–BMDCs were enriched by negative selection using PE-labeled anti–SIRPα antibody (Miltenyi Biotec). OT-I T cells were purified from OT-I x Rag1−/− or OT-I x B6.SJL mice and single-cell suspensions of LNs and spleens prepared by liberase/DNAse digestion, followed by negative selection using a cocktail of PE-conjugated antibodies (anti-CD11c, CD11b, B220, MHC II, CD4, NK1.1) and then anti-PE microbeads (Miltenyi Biotec). OT-II cells were similarly purified using an antibody cocktail containing anti-CD8 instead of anti-CD4 mAb.

Injections and infections. Cells were suspended in PBS and injected i.v. into the tail vein or subcutaneously into the flank (200 μl per mouse). For immunization with dead cells, mice received 5 × 106 to 7.5 × 106 UV-treated OVA-MEFs. For tumor challenge experiments, mice were untreated or immunized with OVA-expressing UV-treated OVA-MEFs. One week later, mice were given OVA-expressing B16 melanoma cells (2.5 × 105/mouse) i.v. Mice were sacrificed 18 days after tumor challenge, and tumor burden was assessed by counting lung foci. For virus infection, mice were anesthetized and infected i.n. or i.v. with 1 × 104 TCID50 units of HSV-1 (KOS VR-1493 from ATCC) as previously described (71).

Flow cytometry. Unconjugated anti-FcγRII/III, FITC–, PE– or APC-conjugated anti-CD11c, FITC– or PE–coupled anti-CD24, PECy5–coupled and biotinylated anti-CD44, PE-conjugated anti-CD172a (SIRPα), anti-CD11b, anti-NK1.1, anti-CD3, anti-CD45.1, anti-B220, FITC–, PE–, or APC-conjugated anti-CD4, PE-conjugated anti–IFN-γ and FITC–, PE–, PerCP–, or APC-conjugated anti-CD8 antibodies were all from BD. Alexa Fluor 647–coupled and biotinylated anti–Dectin-1 and isotype–matched control mAbs were from Serotec. Pacific Blue–coupled anti-CD8 was from BioLegend. Capture anti–IL-2 (BD), anti–IL-10 (BD), anti–IL-12p40 (BD), and anti–TNF-α (R&D) and detection biotinylated anti–IL-2 (BD), anti–IL-10 (BD), anti–IL-12p40 (BD), and anti–TNF-α (R&D) antibodies were used for ELISA. Mouse IL-10, IL-6, and TNF-α cytokometric bead array Flex sets were from BD. Biotinylated anti–DNGR-1 (1F6) was previously described (34). H-2Kα/OVA iTAG MHC tetramers (Beckman Coulter) and H-2Kβ– SSIEFARL pentamers (ProlImmune) were used for detection of OVA–specific and gB–specific CD8+ T cells, respectively. Cell suspensions from spleen, LNs (prepared by liberase/DNAse digestion), or blood were washed and stained in PBS containing 2% FCS, 2 mM EDTA, and 0.02% sodium azide. Cells were incubated with a saturating amount of anti-FcγRII/III before staining with the appropriate antibody cocktail. Propidium iodide was added to the final suspension to exclude dead cells before acquisition on a FACS Calibur or Fortessa (BD Biosciences) and analysis performed with FlowJo (Tree Star) software. Live lymphocyte counts were calculated from the acquisition of a fixed number of 10-μm latex beads (Beckman Coulter) mixed with a known volume of unstained cell suspension. For intracellular IFN-γ staining, cells from mediastinal LNs were restimulated with 1 μM gBpep.52–60 peptide (synthesized and purified by HPLC at the London Research Institute) for 6 hours. Brefeldin A (5 μg/ml, Sigma-Aldrich) was added for the last 5 hours of stimulation. Cells were then stained with anti-CD8–Pacific Blue and anti-CD4–PECy5, fixed and permeabilized, and then stained with PE–coupled anti–IFN-γ mAb.

Chimeric receptors. The complete open reading frame of mouse DNGR-1 or mouse Dectin-1 was cloned upstream of an IRES-GFP sequence into the retroviral vector pFB and pMSCV. The DNGR-1/Dectin chimeric receptor consisting of the cytoplasmic tail of DNGR-1 fused to the transmembrane, stalk region, and CTLD of Dectin in pFB-IRES-GFP was generated using overlap extension PCR with primers 5′-CCATTTGACGTGGTTAGGG-3′ and 5′-CTAAAACCATCCGATGTTGCTCCTGAAATTTGTCATTAG-3′. For the DNGR-1/Dectin chimeric receptor consisting of the cytoplasmic tail and transmembrane region of DNGR-1 fused to the stalk region and CTLD of Dectin in pMSCV, the primers were 5′-TGCTCTTTCTAATGGATCAT-3′ and 5′-AGATCCCAAATAGGAAGGCACCTG-CAGTCATACCTGTAAT-3′. Mutagenesis of DNGR-1- or chimeric receptors was carried out by PCR with primers 5′-TTTTCCCGTGCACATGCGATTCGGCAGCAGG-3′ and 5′-CTGCGCGCGCGCGCGCCAGG-3′. For the I6G mutant chimera, 5′-TTTTCCCGTGCACATGCGATTCGGCAGCAGG-3′, 5′-TTTCCCGTGCACATGCGATTCGGCAGCAGG-3′, and 5′-TTTCCCGTGCACATGCGATTCGGCAGCAGG-3′ for the Y7F mutant. All constructs were verified by sequencing. Receptors were introduced into Syk-expressing B3Z cells, LK cells, and Dectin-1–deficient BMDCs by retroviral transduction.
Cell culture. For DC activation assays in vitro, cells were plated at 2.5 x 10^4 to 5 x 10^5 cells/ml in 96-well plates at 37 °C in the presence of different stimuli. After overnight culture, cytokine concentration in the supernatant was determined by ELISA or by cytometric bead array using standard procedures. For cross-priming assays, CD8α-like Flt3L-BMDCs were cultured at 2.5 x 10^5 cells/ml with purified and CFSE-labeled OVA-specific OT-I or OT-II T cells in 96-well plates for 3 or 5 days, respectively. As a source of antigen, we used pre-treated OVA-MEFs, S-FV-ova-infected MEFs, SIINFEKL peptide (synthesized and purified by HPLC at the London Research Institute), or low-endotoxin OVA protein (Calbiochem) coated on 3-μm polystyrene beads (Sigma-Aldrich). For NFAT reporter assays in B3Z cells, cells were washed in PBS after culture and LacZ activity measured by lysis in CPGR-containing (Roche) buffer. Two to four hours later, OD 595 nm was measured using OD 655 nm as a reference.

Confocal laser scanning microscopy. For uptake of anti-DNGR-1 mAb, purified splenic CD8α+ DCs from WT mice were cultured in the presence of Alexa Fluor 488–coupled anti-DNGR-1 mAb (7H11, Biolegend). Samples were washed in PBS containing 1 mM MgCl2, fixed in 3.7% paraformaldehyde/PBS for 10 minutes, and permeabilized with 0.1% Triton X-100/PBS for 3 minutes. After washing with PBS, cells were blocked with PBS containing 1% bovine serum albumin (Sigma-Aldrich), 5% goat serum (Sigma-Aldrich), and anti-FcγRII/III receptor. DNGR-1 was detected using 7H11 mAb conjugated to Alexa Fluor 546. DNGR-1 was detected using 7H11 mAb conjugated to Alexa Fluor 546. Samples were mounted in Fluoromount-G (SouthernBiotech) and analyzed using an Axiosport 100 laser scanning confocal microscope (Zeiss). Image analysis was performed with LSM 510 software (Zeiss) and ImageJ software.

Microarray data. Microarray data have been deposited at the GEO data repository (GEO GSE36435; NCBI tracking system #16501876).

Statistics. Statistical significance was determined using 1-way ANOVA (Tukey’s post-test) and unpaired 2-tailed Student’s t test. A P value less than 0.05 was considered significant.

Study approval. All animal experiments were performed in accordance with national and institutional guidelines for animal care and were approved by the London Research Institute Animal Ethics Committee and by the Home Office, UK.

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