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**Graphical abstract**

Langerhans cells: Autophagy links antigen presentation and antimicrobial activity

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Autophagy links antimicrobial activity with antigen presentation in Langerhans cells

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Introduction

Langerhans cells (LC) are members of the DC family of professional antigen presenting cells that are distinguished by their residence in the epidermis of skin and mucosa, high expression of CD1a (1), and unique expression of langerin (CD207). As part of the host innate immune system at the interface with the external environment, LC are equipped with endocytic receptors for phagocytosis (2, 3) and have a key role in antigen presentation to T cells (4). In response to cutaneous fungal infection, LC are required for T cell differentiation (5–7).

CD1a is part of the human cluster of differentiation 1 (CD1) gene family consisting of a small number of genes that are structurally related to the MHC class I genes (8); however, unlike MHC class I molecules, CD1 proteins are nonpolymorphic. The CD1a, -b, and -c molecules are more closely homologous to one another than to CD1d (8) and are absent in mice. In skin, CD1a and -c isoforms are present on the surface of LC; whereas, CD1a, -b, and -c are expressed on dermal DC (9–11). CD1a-restricted T cell responses include autoreactive (12, 13) as well as against mycobacterial antigens, including host-derived sulfatides (14), “headless antigens” (13), pollen phospholipids (15), and dideoxymycobactin (16). The crystal structure of CD1a has been resolved with both sulfatide (17) and dideoxymycobactin (18), indicating the presence of 2 hydrophobic binding pockets A’ and F’.

CD1a autoreactive T cells home to skin (19), recognizing CD1a loaded with skin-derived lipids (13), which allow direct interaction between the T cell receptor and CD1a (21).
Leprosy, an infection of skin caused by the intracellular pathogen *Mycobacterium leprae* (*M. leprae*), provides a model to study mechanisms of host defense against an intracellular bacterial pathogen at the site of infection. The disease presents as a spectrum, in which the clinical manifestations correlate with the ability of the immune response to contain the pathogen. There are increased numbers of CD1a+ LC, both in the epidermis and dermal granulomas of the skin lesions—from the self-limiting tuberculoid (T-lep) form vs. the progressive lepromatous (L-lep) form of leprosy (9, 22, 23). LC have been shown to present *M. leprae* antigens to T cell clones derived from skin lesions of T-lep patients in a CD1a-restricted manner (24). Of relevance, the earliest lesions of leprosy are thought to arise in the epidermis (25), and LC have been shown to be infected by *M. leprae* (26).

There are several mechanisms by which CD1-restricted T cells contribute to host defense against mycobacterial infection (27–30). Some mycobacteria-specific group I CD1-restricted T cells secrete the Th1 pattern of cytokines (27), are cytolytic against infected targets (27, 31), and trigger antimicrobial activity (29, 32). These antimicrobial T cells express perforin, granzyme B, and the antimicrobial protein granulysin in intracellular granules (29). Furthermore, the ability of these T cells to release IFN-γ potentially leads to induction of autophagy as well as the vitamin D–dependent antimicrobial response in monocytes and macrophages, including upregulation of the antimicrobial peptides CAMP and DEFB4 (encoding cathelicidin [Cath]/LL-37 and human β-defensin-2, respectively) (33–35). Autophagy is an evolutionarily conserved process in which eukaryotic cells break down cytoplasmic contents by usage of the lysosomes during times of low nutrients or starvation, often as a result of an infection (36, 37). While it has been reported that suppression of Cath during *M. tuberculosis* infection resulted in reduced levels of autophagy in macrophages (38), the role of Cath is still unclear in DC (39).

Although LC have been shown to mediate an antiviral activity (40–42), there is little evidence that LC contribute to antibacterial immunity, rather the opposite that LC contribute to progressive infection (3). Yet in order to fulfill their function as antigen presenting cells, it is reasonable to expect that DC such as LC can mount an antibacterial response in order to facilitate processing of microbial antigens for presentation to T cells during active infection. Therefore, this work was undertaken to learn, through the study of leprosy, whether LC were capable of exerting an antimicrobial response with the ability to process bacterial-derived antigen for presentation to T cells.

**Results**

*IFN-γ induced antimicrobial activity in LCDC and epidermal LC.* Given the increased numbers of LC in the epidermis of T-lep vs. L-lep patients (9, 22, 23), we sought to determine whether LC could encounter IFN-γ in this location. Previously, IFN-γ mRNA was found to be more highly expressed in T-lep vs. L-lep lesions (43–45), yet it was unclear if IFN-γ protein was present in the epidermis in these lesions. Here, we detected higher levels of IFN-γ protein by immunoperoxidase and immunofluorescence in the epidermis of T-lep vs. L-lep lesions, colocalizing with CD1a+ LC (Figure 1, A–C, and Supplemental Figure 1; supplemental material available online with this article; https://doi.org/10.1172/jci.insight.126955DS1).

IFN-γ is known to activate antimicrobial pathways to kill intracellular pathogens. We sought to determine whether this cytokine could induce an antimicrobial activity in *M. leprae*-infected LC. LC-like DC (LCDC) were derived from CD34+ hematopoietic precursor cells, which have been used as surrogates for LC (Supplemental Figure 2A) (24, 41, 46). Electron microscopy of LCDC confirmed the presence of Birbeck granules, which are specific for LC (Supplemental Figure 2B). In addition, epidermal CD1a+ LC were also isolated from healthy skin (Supplemental Figure 2C) (6, 47). To establish whether *M. leprae* is able to infect LC in vitro, as this has not been demonstrated, LCDC and epidermal LC were cultured with live *M. leprae* at increasing multiplicities of infection (MOI) of 5, 10, and 20 per cell. A MOI of 10 yielded approximately 50% of the cells infected with *M. leprae*, with approximately 3–5 bacilli per cell. Both LCDC and LC maintained cellular morphology (Supplemental Figure 3, A and B) and retained the expression of the LC-specific marker CD207/langerin (Supplemental Figure 3C).

To measure the antimicrobial response of LC against intracellular *M. leprae*, bacterial viability was measured according to the ratio of bacterial 16S rRNA to repetitive element DNA using a PCR-based method (45, 48, 49), as the bacteria cannot be grown in culture. LCDC were infected with *M. leprae*, cultured with 10% human vitamin D–sufficient serum, and stimulated with either IFN-γ or media control. IFN-γ treatment of infected LCDC resulted in the induction of significant antimicrobial activity vs. media control (83% ± 5%, P < 0.001; Figure 1D). Similarly, IFN-γ–treated CD1a+ epidermal LC induced a significant antimicrobial response against *M. leprae* (77% ± 4%, P < 0.001; Figure 1E). Conversely, IFN-γ treatment of the CD1a+ epi-
dermal cells, which contained <10% LC, induced an approximately 6-fold lower antimicrobial response (12% ± 2%, P < 0.03 of CD1a− vs. CD1-negative LC; Figure 1F), suggesting that the CD1a+ LC mount a more robust antimicrobial response. As a control, we treated M. leprae–infected LCDC with IL-4, a cytokine more highly expressed in L-lep lesions. This did not induce an antimicrobial response (Supplemental Figure 4).

IFN-γ induces autophagy and phagolysosomal fusion in M. leprae–infected LC. The ability of M. leprae to block phagolysosomal fusion in macrophages prevents the delivery of antimicrobial effector molecules from lysosomes into the phagosomes, which contain the invading bacteria (50–55). This blockade can be overcome by the cellular process of autophagy, through the creation of autophagosomes and their subsequent fusion with lysosomes (34, 35, 56–60). Because there is little information on the induction of autophagy and its biological role in LC, we stimulated LCDC and epidermal LC with IFN-γ and the induction of autophagy was assessed by LC3 aggregation. IFN-γ induced an increase LC3 aggregation in both LCDC and epidermal CD1a+ LC as compared with cells cultured with media control (Figure 2, A–D). LCDC had 18.6 ± 2 puncta per cell for IFN-γ treatment vs. 0.9 ± 0.3 LC3 puncta per cell for media treatment (P < 0.01; Figure 2B). Similarly, epidermal LC expressed 22.0 ± 1.9 puncta per cell for IFN-γ treatment as compared with 2.5 ± 0.7 LC3 puncta per cell for media treatment (P < 0.01; Figure 2D). The ability of IFN-γ to induce autophagy was also evidenced by an increase in the percentage of cells with LC3+ vesicles (Supplemental Figure 5, A and B), as well as the accumulation of LC3-II protein resulting from the cleavage of LC3-I precursor (Supplemental Figure 5C) (61). We also utilized acridine orange, a lysosomotropic dye, to quantify the IFN-γ–induced formation of acidic vesicular organelles (AVO) (Supplemental Figure 6A) (62, 63). Analogous to the number of LC3 puncta, IFN-γ induced more AVO puncta per cell than media treated LCDC, 19.0 ± 0.9 vs. 5.2 ± 1.0 AVO per cell (P < 0.01; Supplemental Figure 6B). Electron micrograph revealed that IFN-γ stimulation of LCDC induced a significant increase in the number of electron-dense endolysosomes compared with media control: 22.1 ± 1.2 vs. 7.7 ± 0.7 endolysosomes per cell (P < 0.01; Supplemental Figure 6, C and D). Utilizing the principles of point counting (64), we quantified the cellular volume of single-membrane and double-membrane compartments, the latter being a characteristic of autophagosomes. LCDC treated with IFN-γ had higher double-membrane volume than media control, 0.07 ± 0.01 μm³ vs. 0.02 ± 0.01 μm³ (P < 0.03), while the inverse was observed between the conditions with single-membrane volume (Supplemental Figure 6E).

We next investigated the effects IFN-γ–mediated phagosome maturation and phagolysosomal fusion during M. leprae infection. As compared with cells treated with media alone, IFN-γ treatment of M. leprae–infected LCDC induced autophagy as well as colocalization of intracellular bacteria to the autophagosome, identified by LC3 aggregation and colocalization with the labeled M. leprae bacilli and further delineated by 3-dimensional modeling (Figure 2E). IFN-γ treatment of LCDC also resulted in colocalization of bacilli and LAMP1, a lysosome marker, indicating that the majority of M. leprae bacilli were localized to the lysosome compartment (Figure 2F). Furthermore, IFN-γ stimulation of M. leprae–infected LCDC triggered autophagolysosomal fusion, as evidenced by the colocalization of the bacilli with LC3 and LAMP1 (Figure 2G). By electron microscopy (EM), we confirmed that, in IFN-γ–treated LCDC, the M. leprae bacilli were contained within compartments surrounded by autophagosomes, characterized by double-membrane vesicles. We also detected evidence by EM of IFN-γ–induced fusion between autophagosomes, characterized by double membranes and containing bacilli, with lysosomes. In contrast, the majority of the intracellular bacilli in the control LCDC treated with media were contained in phagosomes with a single membrane (>90%; Figure 2H and Supplemental Figure 6F). Additionally, the mycobacterial cell walls were mostly intact in the media control–treated LCDC, while in IFN-γ–treated LCDC, the cells walls were often degraded and the bacilli lysed. Together, these data demonstrate that M. leprae blocks the inhibition of autophagolysosomal fusion is overcome by IFN-γ activation, facilitating an antimicrobial response.

IFN-γ induces antimicrobial peptides in LC. IFN-γ can trigger an antimicrobial response against intracellular pathogens by a variety of mechanisms, including the vitamin D–dependent generation of antimicrobial peptides CAMP and DEF4 in monocytes and macrophages (33, 34). The addition of IFN-γ was determined to be sufficient to induce the gene expression of the antimicrobial peptides CAMP and DEF4 in both LCDC (Figure 3A) and epidermal CD1a+ LC (Figure 3B). IFN-γ was also sufficient to induce Cath protein in LCDC as detected by confocal laser microscopy (Figure 3C and Supplemental Figure 7A). The percentage of Cath+ cells was significantly greater in IFN-γ vs. media-treated LCDC (74% ± 2% vs. 4.9% ± 2.7% in media control, P < 0.002, Figure 3D). In contrast, β-defensin-2 protein expression was detected in CD1a+ LCDC treated with media and was not further induced by IFN-γ stimulation, suggesting that it is constitutively expressed (Supplemental Figure 7, B and C), consistent with its constitutive expression in the skin (65).
IFN-γ induction of Cath in infected LCDC resulted in the colocalization of Cath with intracellular *M. leprae*. In IFN-γ–treated LCDC, Cath was detected surrounding intracellular bacilli and localized to the LAMP1+ lysosome (Figure 3E). In contrast, media treatment of LCDC did not result in detectable Cath. Therefore, to determine the role of Cath in LC-mediated antimicrobial activity against *M. leprae*, transfection of siRNAs targeting Cath mRNA was used. Approximately 50% specific knockdown of Cath mRNA was achieved, affecting cell viability by <10% (Supplemental Figure 8A). Vitamin D receptor (VDR) mRNA levels were unaffected by siRNA targeting Cath (Supplemental Figure 8B). In these experiments, the knockdown of Cath reduced IFN-γ–induced antimicrobial activity from 77% ± 2% to 29% ± 5% (*P* < 0.02), an approximately 60% reduction compared with IFN-γ paired with the small interfering control (siCTRL) (Figure 3F). During *M. leprae* infection, Cath peptide was undetectable in media control–treated LCDC but was found to colocalize with *M. leprae* inside LAMP1+ vesicles when stim-
Figure 2. IFN-γ induces autophagy in Langerhans cells. (A) Human LCDC were cultured with rIFN-γ, rapamycin, or medium overnight in 10% vitamin D–sufficient human serum, and they were fixed and immunolabeled with anti-LC3 antibody (green) and anti-CD207/langerin antibody (red). Nuclei were stained with DAPI (blue). Representative shown of 4 independent experiments. (B) The number of LC3 puncta per cell were quantified. Data are represented as mean puncta per cell ± SEM, n ≥ 30 cells. (C) Human primary CD1a+ LC were cultured and labeled as in A. Representative immunofluorescence images of 3 independent experiments are shown. (D) The number of LC3 puncta per cell were quantified. Data are represented as mean puncta per cell ± SEM, n ≥ 25 cells. (E-H) Human LCDC were stimulated with rIFN-γ for 4 hours, washed and infected with PKH26-M. leprae (red) at a MOI of 10 overnight, and washed and stimulated with rIFN-γ for an additional 4 hours. (E) Infected cells were fixed and immunolabeled with anti-LC3 antibody (green). Representative shown of 4 independent experiments. (F) Infected cells were fixed and immunolabeled with anti-LAMP1 antibody (green). Representative shown of 4 independent experiments. (G) Infected cells were fixed and immunolabeled with anti-LC3 antibody (green) and anti-LAMP1 antibody (cyan). Representative shown of 3 independent experiments. (A, C, F, and G) Images captured on a 63× lens, with 6× zoom. (H) Infected cells were fixed and processed for electron microscopy. N, nucleus; L, lysosome. Black boxes labeled 1–4 correspond with magnifications shown. White arrows indicate single-membrane phagosomes. Yellow arrows indicate double-membrane autophagosomes. Blue arrows indicate digested bacteria. Yellow asterisks indicate fusion of autophagosome with lysosome. Scale bars: 1 μm. Representative images of 3 independent experiments are shown. **P < 0.01. Repeated measures 1-way ANOVA.
Cath is known to be secreted (66). These data corroborate that skin-resident CD1a+ LC are in proximity to distinguish if the colocalization is due to cellular production or uptake from extracellular sources, given that γCath also required the presence of adequate levels of 25-hydroxyvitamin D (25D), as treatment of IFN-γLC (Supplemental Figure 13A) and LCDC (Supplemental Figure 13C). The addition of VDR antagonist ZK γdroxyvitamin D3 (1,25D3) to vitamin D–insufficient sera (Supplemental Figure 12, A–C). IFN-γ–induced autophagy was significantly greater in the presence of vitamin D–sufficient sera or the addition of 1,25-dihydroxyvitamin D3 (1,25D3) to vitamin D–insufficient sera (Supplemental Figure 14A) but could be rescued with the addition of 25D or bioactive 1,25D3 (Supplemental Figure 14C). The induction of autophagy in LC was relevant at the site of infection in leprosy, we utilized confocal microscopy. LC3 puncta were detected in CD1a+ LC in T-lep but not L-lep lesions, indicating the presence of autophagic responses (Figure 4, A and B, and Supplemental Figure 9, A and B). Cath protein was ubiquitously expressed in the epidermis of T-lep lesions and colocalized with CD1a+ LC. In contrast, in L-lep lesions, the expression of Cath was reduced (Figure 4, C–E, and Supplemental Figure 9, C and D). The isotype controls matching all monoclonal antibodies used for labeling were consistently negative (Supplemental Figure 9). Even though the confocal imaging demonstrated colocalization of Cath with CD1a+ LC in T-lep lesions, we cannot distinguish if the colocalization is due to cellular production or uptake from extracellular sources, given that Cath is known to be secreted (66). These data corroborate that skin-resident CD1a+ LC are in proximity to IFN-γ (Figure 1, A and B) and colocalize with Cath and autophagosomes, indicating the presence of key components of this antimicrobial pathway in LC at the site of infection.

As part of the DC family of professional antigen presenting cells, LC are characterized by an enhanced ability to activate T cell responses. In skin, there is a resident T cell population that responds to microbial antigens (67, 68). To study the resident T cell response against M. leprae, a CD1a-restricted T cell line, LCD4.G — derived from a T-lep skin lesion — was studied. LCD4.G is CD4+ expressing CD69 and CD45RO but not CD103 or CD45RA, consistent with the T-resident memory (T RM) phenotype (Supplemental Figure 10). Mycobacteria are known producers of a variety of lipids, which can form stable complexes with CD1 molecules and stimulate T cells (69, 70). To validate whether antigen presentation to LCD4.G T cells is CD1a dependent, we added CD1a blocking antibody to LCDC prior to the coculture with T cells. Anti-CD1a blocked the T cell response by about 80%, as compared with the addition of isotype control (Figure 5A). Supernatants, collected from LCD4.G after activation using anti-CD3/CD28 antibodies, contained high levels of IFN-γ (Figure 5B). When added to LCDC, the LCD4.G supernatants induced Cath and DEFB4 mRNAs. Neutralization of IFN-γ by neutralizing mAbs significantly blocked the induction of Cath and DEFB4 mRNA levels as compared with the isotype control (Figure 5C and Supplemental Figure 11A). Therefore, T RM cells, via the secretion of IFN-γ, triggered the induction of anti-microbial peptides in LCDC. We also determined that the CD1a-restricted T cell clone LCD4.G triggered the IFN-γ dependent upregulation of CYP27b1, VDR and CYP24A1 in LCDC (Supplemental Figure 11, B–D), components of the vitamin D antimicrobial pathway.

In studying the role of vitamin D in the LC response, we observed that the ability of IFN-γ to induce autophagy was significantly greater in the presence of vitamin D–sufficient sera or the addition of 1,25-dihydroxyvitamin D3 (1,25D3) to vitamin D–insufficient sera (Supplemental Figure 12, A–C). IFN-γ induction of Cath, DEFB4, and CYP24A1 mRNAs was dependent on activation of the VDR in both primary epidermal LC (Supplemental Figure 13A) and LCDC (Supplemental Figure 13C). The addition of VDR antagonist ZK 159 222 (VAZ) significantly inhibited the induction of these vitamin D–dependent genes. The induction of Cath also required the presence of adequate levels of 25-hydroxyvitamin D (25D), as treatment of IFN-γ in 10% FCS, which had low 25D concentrations and was unable to induce gene expression (Supplemental Figure 14A) but could be rescued with the addition of 25D or bioactive 1,25D3 (Supplemental Figure 14C). The levels of vitamin D did not affect IFN-γ induction of the vitamin D–independent gene CD64 (Supplemental Figure 13, B and D, and Supplemental Figure 14B). Conversely, the IFN-γ–induced antimicrobial response was vitamin D dependent, such that even in 25D-insufficient sera, the antimicrobial activity could be restored by in vitro supplementation with 25D (Supplemental Figure 15, A and B). The data indicate that IFN-γ induction of an antimicrobial response in LC is dependent on 25D levels.

Role of autophagy in LC function. In human macrophages, autophagy can be triggered by a vitamin D–dependent pathway (34, 71, 72), contributing to an antimicrobial response. In contrast, in DC, autophagy is recognized to contribute to effective antigen presentation to T cells by delivering antigen into antigen processing compartments (73, 74). This led us to hypothesize that autophagy may be a conserved mechanism by which DC, such as LC, link antimicrobial activity to efficient presentation of microbial antigens to T cells. To investigate the role of autophagy in LC function, we used the phosphatidylinositol 3-kinase (PI3K) inhibitor wortmannin to block IFN-γ–induced autophagy (Figure 6A). The addition of wortmannin prior to IFN-γ stimulation significantly
reduced the number of accumulating LC3 puncta, from 19.3 ± 2.9 puncta to 5.3 ± 1.0 puncta (P < 0.01; Figure 6B). Similarly, the percentage of LC3+ cells was also reduced in the presence of wortmannin (Supplemental Figure 16). The IFN-γ–induced antimicrobial activity in M. leprae–infected LCDC was dramatically inhibited by approximately 75% by the addition of wortmannin (Figure 6C), and by about 50% by the knockdown of ATG5 (Figure 6D), indicating that induction of autophagy is required for optimal antimicrobial activity. LC, by their proximity to the external environment, deal with microbial pathogens as they invade via the skin. LC express C-type lectins and other endocytic receptors that mediate the phagocytosis of various microbes (2, 3) subsequently instruct the adaptive T cell response (5–7). There is evidence that LC mediate an antiviral response (40, 41); however, little is known about the ability of LC, or DC in general, to kill such bacteria and fungi as part of the innate immune response. Given that Staphylococcus aureus, Streptococcus pyogenes,
and *C. albicans* are frequent skin pathogens, we infected LCDC and/or LC with these organisms and determined whether IFN-γ induced an antimicrobial response. IFN-γ reduced the viability in *S. aureus*–infected primary LC by about 60% (Figure 6E). Furthermore, we found that the induction of autophagy by IFN-γ was crucial in limiting the growth of *S. aureus*, *S. pyogenes*, and *C. albicans* in infected LCDC (Figure 6, F and G).

Because autophagy has been shown to enhance MHC class I– and MHC class II–restricted antigen presentation (53, 54), we investigated whether IFN-γ–induced autophagy in LC, which reduces bacterial viability, would augment CD1a-restricted antigen presentation of live bacilli. We verified that the treatment of LCDC with IFN-γ did not have a substantial effect on the expression of CD1a, regardless of the concentration of 25D or 1,25D in the culture (Supplemental Figure 17, A–F). In addition, the expression of CD1a and HLA-DR were not substantially affected by infection of LCDC with *M. leprae* (Supplemental Figure 17, G and H). Next, LCDC were infected with live *M. leprae* and cocultured with the CD1a-restricted T cell clone LCD4.G using a suboptimal number of antigen presenting cells in order to measure augmentation. IL-2 was used as a positive control to demonstrate proliferative response by LCD4.G (Supplemental Figure 18). IFN-γ stimulation enhanced antigen presentation by primary human epidermal LC and LCDC, resulting in over a 3-fold increase in T cell proliferation compared with antigen alone. The inhibition of IFN-γ–induced autophagy by wortmannin resulted in an approximately 80% decrease of the T cell response (Figure 7, A and B).

In comparison, monocyte-derived macrophages (MDM), which did not express CD1a (Supplemental Figure 19) and lack a consensus matching MHC class II allele (Supplemental Table 1), did not present live *M. leprae* to the CD1a-restricted LCD4.G T cells (Figure 7C). These data indicate that autophagy is required for the ability of LC to efficiently present antigen from live *M. leprae* to T cells. Together, these findings indicate that resident LC in skin are essential in promoting adaptive TRM responses during bacterial infection.
Discussion

In primitive eukaryotic organisms such as yeast and amoebas, autophagy has an essential role in cell survival by managing the flow of nutrients and by clearing damaged organelles (75, 76). It has become increasingly clear that more complex organisms have evolved to utilize autophagy for other physiologic pathways, including host defense, by bringing phagocytized microbial pathogens to lysosomes where they are destroyed. However, it is not known whether the ability of autophagy to facilitate antigen presentation by DC is linked to an antimicrobial response. Here, we provide evidence that, in LC, which are resident DC at the interface of the skin with the outside environment, autophagy delivers bacterial pathogens to lysosomes, where they are killed and subsequently processed into cognate antigens facilitating T cell recognition. These data suggest that autophagy is a critical mechanism allowing DC to survive invasion by a microbial pathogen via an antimicrobial response, which contributes to efficient antigen presentation to T cells.

A major function of DC is to bridge the innate and adaptive immune response through the efficient presentation of antigen to T cells. The ability of DC to optimally present antigen via MHC class II is enhanced by the induction of autophagy, as shown in murine models of infection including HSV-1 (77, 78), influenza (79), and mycobacteria (73, 80). Induction of autophagy in human DC, albeit derived from monocytes differentiated with GM-CSF and IL-4, enhanced MHC class II presentation of HIV-1 (81) and the C-fragment of tetanus toxin expressed by salmonella (74). Here, we found that IFN-γ, by inducing autophagy in M. leprae-infected LC, enhanced CD1a-restricted antigen presentation of live M. leprae to a CD4+ T cell line, LCD4.G, derived from the TRM population in a leprosy skin lesion. Furthermore, LC are highly specialized in their function; while IFN-γ induced autophagy and antimicrobial activity against mycobacteria in both human LC and macrophages (34, 45), only LC were equipped to present M. leprae antigen to the CD1a-restricted T cells. LC are not only involved in the presentation of microbial antigens to T cells but also present self-antigens to skin-homing T cells via CD1a (19), including skin-derived lipids (13) and free fatty acids (20). Further studies are needed to define the role of autophagy in the different pathways by which LC present antigen to T cells.

Mycobacteria are known to inhibit phagolysosomal fusion (50–55) in macrophages, which is overcome by autophagy (34, 35, 56–60). In infected human LC, as in macrophages, IFN-γ induction of autophagy and subsequent phagolysosomal fusion is required for an antimicrobial activity (34). IFN-γ also induces the vitamin D–dependent production of Cath in LC, which is delivered to phagolysosomal compartments containing M. leprae and is required for the antimicrobial response. Autophagy components may also contribute to host defense via other pathways (82). There are a few studies indicating that DC mount an antimicrobial response against an intracellular pathogen (40, 83), with one report indicating that the induction of autophagy enhances antimicrobial function (84). It was recently reported that HIV infection induces auto-
Phagy in LC, resulting in the trafficking of the virions into the autophagosome for degradation (42). One major limitation of our study is that we are unable to assess the in vivo role of autophagy in the immune response to *M. leprae*, as there is no animal model that mimics the disease spectrum of human leprosy. Also, it is not clear whether vitamin D is required for IFN-γ–induced autophagy in mouse macrophages and DC. Although IFN-γ induction of autophagy is vitamin D dependent, it is not clear if this is true in mouse macrophages. However, our data provide insight into the role of autophagy-induced antimicrobial activity in DC function, suggesting that this host defense pathway also results in the breakdown of the pathogen into cognate antigen for optimal activation of T cell responses.

We speculate that the ability of DC, such as LC, to mount this antimicrobial response not only contributes to enhanced antigen presentation, but may also represent a multifaceted survival mechanism. We found that IFN-γ–induced autophagy-dependent antimicrobial activity in LC against other cutaneous pathogens, including *S. aureus*, *S. pyogenes*, and *C. albicans*. Whereas in macrophages, autophagy only contributes to the destruction of pathogens in the endolysosomal compartment, autophagy in LC enhances the bactericidal effects of IFN-γ. This may provide a selection pressure for the development of virulence traits in pathogens that escape autophagy-induced killing in LC.
of such microbes (85–88), our data show that autophagy contributes both to an antimicrobial response in LC, which then allows DC to process and present pathogen-derived antigens to T cells, as well as to instruct the nature of the adaptive T cell response (5–7). The present findings are relevant to host defense at the site of infection, as the LC in leprosy lesions colocalized with IFN-γ exhibited an autophagic response and contained Cath protein. Finally, skin-derived LC were able to process and present live M. leprae to a TRM cell line derived from a leprosy lesion, resulting in the release of IFN-γ, providing an amplification loop to further augment autophagy, antimicrobial activity, and antigen presentation essential for effective immunity at sites of infection. In summary, these results provide evidence that LC contribute to host defense against cutaneous pathogens.

Methods

Reagents. Recombinant human IFN-γ was purchased from R&D Systems and used at a concentration of 10 ng/ml. Both 25(OH)D3 and 1,25D3 were purchased from BioMol, dissolved in ethanol, and used at a concentration of 1 × 10⁻⁷ M. Wortmannin, dissolved in DMSO and used at 500 nM; rapamycin, dissolved in DMSO and used at 300 nM; poly-L-lysine; and saponin were purchased from MilliporeSigma. RPMI cell culture media, IMDM cell culture media, Hyclone FCS, CD3/CD28 Dynabeads, DAPI antifade with prolong Gold, Lipofectamine RNAiMAX, and TRIzol were purchased from Invitrogen. CD3/CD28 DynaBeads were used at bead/cell ratio of 1:1. Monoclonal antibodies and their corresponding isotype controls used were the following: anti-IFN-γ (clone B27, IgG1), APC anti–human CD1a (clone HI149, IgG1), PE anti–human CD80 (clone L307.4, IgG1), PE anti–human CD86 (clone IT2.2, IgG2b), and PE anti–human HLA-DR (clone Tu39, IgG2a) were purchased from BD Biosciences. Purified anti–human CD1a (clone SK9, IgG2b), purified anti–human CD207/langerin (clone 4C7, IgG2a), and Alex647 anti–CD107/LAMP1 (clone H4A3, IgG1) were obtained from BioLegend. PE anti-CD207/langerin (clone DCGM4, IgG1) was purchased from Beckman Coulter. Purified anti–human LC3 (clone 4E12, IgG1) was obtained from MBL International. Anti-Cath (clone OSX12, IgG1) and anti-β-defensin-2 (rabbit polyclonal EPR20469) were purchased from Abcam. CD1a magnetic beads were purchased from Miltenyi Biotec. Recombinant IL-2, used at 1 nM, were from Chiron Diagnostics. Recombinant GM-CSF was obtained from Genzyme Corporation, a Sanofi company.

Generation of LCDC. CD34⁺ cells were cultured in RPMI in the presence of SCF (25 ng/ml), GM-CSF (100 ng/ml), and TNF-α (500 ng/ml). At day 9, CD14⁺ cells were depleted from the culture. The remaining CD14⁺ cells were replated in the presence of GM-CSF (100 ng/ml) and TGF-β1 (1 ng/ml) to increase CD1a expression. LCDC were harvested at day 13–15 and enriched for CD1a by using anti-CD1a magnetic beads.
beads. Of the CD1a+ LCDC, 80%-90% were found to also be CD207+, by surface staining.

**Preparation of human LC.** Epidermal cell suspensions were prepared from deidentified normal human skin taken from patients undergoing abdominal plastic surgery at UCLA Division of Plastic and Reconstructive Surgery with the approval of UCLA Tissue Procurement Core Laboratory, as previously described (6, 47). The attached adipose and connective tissues were removed, and the skin was cut in approximately into 0.5 cm² pieces, which were rinse in HBSS (Thermo Fisher Scientific) without calcium, magnesium, or phenol red. Skin pieces were then incubated in 0.2% dispase HBSS. The epidermis was separated from the dermis, mechanically separated into a single cell suspension, and filtered through 40-μm cell strainers (BD Biosciences). CD1a+ LC were isolated from the epidermal cell suspension by using anti-CD1a magnetic beads. Of the CD1a+ LC, 50% were found to also be CD207+ by surface staining. Epidermal LC were cultured in IMDM.

**Human serum collection and 25D quantification.** Blood was collected from healthy donors in the absence of anticoagulants and allowed to clot for 2 hours. The serum layer was collected and passed through a 0.22-μm filter and frozen in small aliquots at –80°C for future use. The concentration of 25D was determined by radioimmunoassay, as previously described (33, 89). Serum was pooled from several donors to obtain vitamin D–sufficient and vitamin D–insufficient batches. The 25D level in the vitamin D–sufficient pooled serum batches was approximately 100 nM (40 ng/ml), and in the vitamin D–insufficient serum, batches less than 35 nM (14 ng/ml). The 25D level in the FCS was less than 18 nM or 7 ng/ml.

**LCDC and epidermal LC culture for PCR.** Following CD1a selection, cells were cultured in 10% FCS or 10% human serum, with or without 10% final volume of T cell supernatants, as indicated. IFN-γ-induced gene expression of CYP27B1, VDR, CYP24, Cath, and DEFB4 mRNAs were detected after 2 hours.

**PCr.** mRNA was isolated from cells using TRIzol according to the manufacturer-recommended protocol. cDNA was prepared and gene expression levels were measured by quantitative PCR (qPCR) and calculated by the 2^ΔΔCt method. Primer sequences for human Cath, DEFB4, CYP27B1, VDR, CYP24A1, human 36B4, CD64, and 16S rRNA and *M. leprae*-specific repetitive element (RLEP) of *M. leprae* were previously reported (33, 34, 48).

**siRNA transfection of LCDC.** Cath siRNA, ATG5 siRNA, and siCTRL, obtained from GE Dharmacon, were incubated with Lipofectamine RNAiMAX (Invitrogen) for 30 minutes to allow complexing. Then, siRNA was added to the LCDC culture at 200 nM, per transfection, 30 minutes and overnight prior to the addition of IFN-γ for Cath siRNA and ATG5 siRNA, respectively.

**Infection of epidermal LC and LCDC, treatment with IFN-γ, and quantification of antimicrobial activity.** *M. leprae* was grown in the footpad of nu/nu mice, as described previously (90), and was provided by the National Hansen’s Disease Program (Baton Rouge, Louisiana, USA). LCDC were pretreated with 10 ng/ml of IFN-γ or IL-4 in 10% human serum for 4 hours or overnight. The cells were washed prior to the overnight infection with *M. leprae* at a multiplicity of infection (MOI) of 10 in 10% FCS. Extracellular bacteria were removed by vigorous washing. The infected cells were then treated again with the same concentration of IFN-γ or IL-4 in 10% human serum and harvested after 4 days. Epidermal LC were treated with IFN-γ and infected with mLEP similarly as above. RNA and DNA were isolated from infected cells following TRIzol protocol. The viability of intracellular *M. leprae* was determined by qPCR and quantified as previously described (45, 48, 49). Briefly, cDNA was synthesized from the total RNA as described for both human and bacterial mRNAs. The *M. leprae* 16S rRNA and genomic element DNA (RLEP) levels were then assessed by real-time PCR. In order to normalize for the total number of LC present in culture, 36B4 was also evaluated. Comparison of the bacterial DNA to the mammalian 36B4 levels was used to monitor infectivity between all the conditions in the assay, as well as PCR quality. The 16S rRNA and genomic DNA values were calculated using the ΔACT analysis, with the bacterial DNA value serving as the housekeeping gene. The efficiency of infection was determined by confocal microscopy and FACS. Approximately 45%-60% of cells were infected. Cells viability following washes and infection was 81%-87% of the total cells.

**Intracellular killing of S. aureus and S. pyogenes by LCDC and epidermal LC.** *S. aureus* strain DU5938 (Hla Hila Hlb Hlg) (91) or *S. pyogenes* (ATCC 700294) (92) were grown to the mid-log phase at 37°C with shaking (150 rpm) in brain-heart infusion (BHI) medium, collected by centrifugation for 10 minutes at 5,000 g. For cellular infection, the bacterial suspension was diluted with sterile 1× PBS. The number of viable bacteria was determined by serial dilution and plating onto BHI agar plates (93). LCDC or epidermal LC (5 × 10⁶ cells) were plated in 24-well plate at 1 ml and infected with *S. aureus* at MOI of 3 for 3 hours in 10% FCS antibiotic-free media. Next, 100 μg/ml of gentamycin was added to the wells for 20 minutes to kill extracellular bacteria. Cells were then washed to further remove any remaining extracellular bacteria. Infected cells were cultured.
overnight in 10% human serum with or without IFN-γ at 37°C in a 4% CO₂ incubator (Thermo Scientific Midi 40 Small Capacity CO2 Incubator). To measure intracellular killing of bacteria, cells were pelleted and lysed with 100 μl of 0.2% saponin in 1× PBS on ice for 20 minutes. PBS 1×, 900 μl, was added to bring the cell lysate to 1 ml final volume, of which 1 μl was plated overnight and CFU was determined.

Antifungal activity. *Candida albicans* (ATCC 18804) were grown by agitation overnight at 28°C in yeast mold (YM) media. The cells were then washed twice with 1× PBS before use as live yeasts (94). LCDC (5 × 10⁶) were plated in 24-well plates at 1 ml and infected with *C. albicans* at MOI of 3 for 3 hours in 10% FCS antibiotic-free medium at 37°C. Amphotericin B (1 μg/ml) was added to wells for 20 minutes to kill extracellular *C. albicans*. Cells were then washed to further remove any remaining extracellular *C. albicans*. Infected cells were cultured overnight in 10% human serum with or without IFN-γ at 37°C in a 4% CO₂ incubator (Thermo Scientific Midi 40 Small Capacity CO2 Incubator). To measure antifungal activity, cells were pelleted and lysed with 100 μl of 0.2% saponin in 1× PBS on ice for 20 minutes. PBS 1×, 900 μl, was added to bring the cell lysate to 1 ml final volume, of which 1 μl was plated for CFU at room temperature. Colonies were counted after 2 days.

Patients and clinical specimens. Patients with leprosy were classified according to the criteria of Ridley and Jopling (95). The designation of T-lep included patients who were classified clinically as borderline tuberculoid (BT), and the designation of L-lep only included patients classified as “LL” (lepromatous leprosy). All T-lep and L-lep skin biopsy specimens were taken at the time of diagnosis, prior to initiating treatment. Specimens were embedded in OCT medium (Ames), snap frozen in liquid nitrogen, and stored at −80°C.

Tissue immunoperoxidase labeling. Frozen tissue sections were blocked with normal horse serum before incubation with mAbs for LC3, Cath, and IFN-γ for 60 minutes, followed by incubation with biotinylated horse anti–mouse IgG (see Reagents above) for 30 minutes. Slides were counterstained with hematoxylin and mounted in crystal mounting medium (Biomeda) and were visualized using the ABC Elite system (Vector Laboratories). Skin sections were examined using a Leica microscope (Leica). Ratios were calculated by ImmunoRatio online software (Jorma Isola & Vilppu Tuominen – Institute of Biomedical Technology, University of Tampere, Tampere, Finland) (96), an automated image analysis application that calculates the percent diaminobenzidine-stained (DAB-stained) nuclear area per total area.

Cell culture immunofluorescence labeling. Cells were treated with IFN-γ or left untreated in 10% human serum or 10% FCS overnight. Then, cells were washed and infected with live *M. leprae* (mLEP) at a MOI of 10 overnight and further stimulated with IFN-γ in 10% human serum for 3 hours. Following stimulation or infection, cells were adhered to poly-L-lysine–coated slides for 1 hour. Cells were then washed and fixed for 30 minutes with 4% PFA before being washed again. Next, cells were permeabilized with 0.25% saponin for 20 minutes, blocked with serum for 30 minutes, and immunolabeled with primary antibodies for LC3, CD1a, CD207, Cath, or DEFB4 for 2 hours. Following washing, cells were stained with secondary antibodies (see Reagents above) for 90 minutes, washed, and mounted with DAPI. For the quantification of autophagy, the number of puncta per cell and the percentages of LC3 punctated cells, as defined the presence of >5 puncta per cell, were evaluated using fluorescence microscopy and quantified by Imaris and Image J (NIH). Approximately 50–100 cells, over 6 different random fields of view, were scored for each condition of each experiment. Rapamycin was used as a positive control for autophagy induction, and wortmannin was used to inhibit autophagy. DMSO used as vehicle control in all media conditions.

Immunofluorescence of cell cultures was examined using a Leica-TCS-SP MP inverted single confocal laser-scanning and a 2-photon laser microscope (Leica) at the Advanced Microscopy/Spectroscopy Laboratory Macro-Scale Imaging Laboratory (California NanoSystems Institute, UCLA). Three-dimensional modeling of microscopy images were generated by Imaris software.

Tissue immunofluorescence labeling. Immunofluorescence was performed by serially incubating cryostat tissue sections with anti-human mAbs of different isotypes for 2 hours and washed 3 times with 1× PBS, followed by incubation with isotype-specific, fluorochrome-labeled (A488, A568, A647) goat anti–mouse immunoglobulin antibodies (Molecular Probes) for 90 minutes. Controls included staining with isotype-matched antibodies. Nuclei were stained with DAPI. Immunofluorescence of skin sections was examined using a Leica-TCS-SP MP inverted single confocal laser-scanning and a 2-photon laser microscope (Leica) at the Advanced Microscopy/Spectroscopy Laboratory Macro-Scale Imaging Laboratory (California NanoSystems Institute, UCLA).

Transmission electron microscopy. Human LCDC were fixed with 2% paraformaldehyde and 2.5% glutaraldehyde (Electron Microscopy Sciences) in 1× PBS (Gibco), pH 7.35, for 1 hour at room temperature and stored at 4°C overnight. The cell pellets were washed with PBS, embedded in 4% low-melt agarose, cut into small pieces, and post-fixed with 1% OsO₄ in ddH₂O. Following washing with ddH₂O, the cells were then washed with 50% ethanol, and post-fixed with 100% ethanol. Then, the samples were dehydrated with increasing concentrations of ethanol, infiltrated with Polybed 812 (Polysciences), and embedded in epoxy resin. Ultrathin sections were cut on a Leica ultracut-U ultramicrotome and placed on 200-mesh copper grids coated with 4% uncoated carbon. The sections were stained with 2% uranyl acetate and lead citrate and examined with a Jeol 1010 transmission electron microscope (Jeol). Bacteria were identified as intracellular based on the presence of the cell wall and the absence of a host envelope.
were stained en bloc with 2% uranyl acetate, washed, dehydrated through a graded series of ethanol, treated with propylene oxide, and embedded in Eponate 12 (Ted Pella Inc.). Silver to gold interference color sections were cut on a TMC ultramicrotome and picked up on formvar-coated copper grids. The sections were stained with Reynolds lead citrate and examined on a JEOL 100CX electron microscope at 60 kV. Images were collected on type 4489 EM film, and the negatives were digitized at 1200 dpi. Adobe Photoshop was used to quantify the pixel area of cellular compartments. The volume fractions of images were calculated by dividing the sum of the area occupied by of single- or double-membrane autophagic compartments with the area of the cell (64).

**Immunoblotting analysis.** LCDC, 1 × 10⁶ per condition, were collected in NP-40 lysis buffer (50 mM Tris [pH 7.4], 50 mM NaCl, 5 mM EDTA, 50 mM NaF, 1 mM Na₃VO₄, 1% Nonidet P40 [NP40], 0.02% NaN₃ and 1 mM PMSF) containing complete protease inhibitors (Roche Applied Science). Total protein from cell lysates were quantified using Bradford Assay. Lysates were separated on a 7.5% nondenaturing gel, followed by immunoblot analysis with anti–β-actin and anti-LC3 (see Reagents above) and detected by enhanced chemiluminescence (Pierce Biotechnology).

**CD1a-restricted T cell clone.** An M. leprae–reactive CD4⁺ T cell line (LCD4.G) was derived from a biopsy of a skin lesion from an untreated patient with T-lep as described (24). To obtain supernatants, 1 × 10⁶ T cells were cultured in 1 ml of media containing 10% FCS with or without the addition of CD3/CD28 Dynabeads, with a bead/cell ratio of 1:1. Supernatants were collected after 24 hour stimulation and stored at –80°C. IFN-γ levels of T cell supernatants were measured by sandwich ELISA.

**Antigen presentation and proliferation assay.** LCD4.G is a CD4⁺ T cell line derived from a tuberculoid lesion as described (24). T cells were maintained by serial antigenic stimulation using LCDC in medium supplemented with recombinant IL-2. Antigen presentation by LC was measured using a suboptimal ratio of 0.1 LC to 1.0 T cell, instead of 1:1, in order to measure both inhibition and augmentation of the T cell response. Epidermal LC and LCDC were pretreated with 10 ng/ml of IFN-γ in 10% human serum overnight. The cells were washed prior to the overnight infection with M. leprae at a MOI of 10 in 10% FCS. Extracellular bacteria were removed by vigorous washing. The infected cells were then treated again with the same concentration of IFN-γ for 2 hours, and they were washed again to remove IFN-γ prior to coculture with T cells. For measurement of T cell proliferation, 1 × 10⁴ T cells were cultured with 1 × 10⁴ uninfected-LCDC (MED) or mLEP-LC (MOI 10) in 10% human serum. T cells were also cultured with 100 U/ml of IL-2 as positive control. Following 4 days of culture in 96-well plates in triplicate at 37°C in a 4% CO₂ incubator, cells were pulsed with ³H-thymidine (1 μCi/well; ICN Biomedicals Inc.) and harvested 4 hours later for liquid scintillation counting. To validate the CD1a restriction of the T cell lines, neutralizing CD1a monoclonal antibodies (20 μg/ml) were added 30 minutes before the addition of T cells. Alternately, 1 × 10⁴ T cells were cultured with 1 × 10⁴ M. leprae–infected monocyte-derived-macrophage. Following 4 days of culture in 96-well plates in triplicate at 37°C in a 4% CO₂ incubator, cells were pulsed with ³H-thymidine (1 μCi/well; ICN Biomedicals Inc.) and harvested 4 hours later for liquid scintillation counting.

**Statistics.** Statistics reported are of entire series of experiments and described as mean ± SEM. GraphPad Prism 6 software was used for graphing and statistical analysis. For comparison between 3 or more groups, we utilized repeated measures 1-way ANOVA, with the Greenhouse-Geisser correction, along with Tukey’s multiple comparisons test, with individual variances computed for each comparison. The 2-tailed Student’s t test was used for all other 2-group analysis. A P value less than 0.05 was considered significant.

**Study approval.** This study was conducted according to the principles expressed in the Declaration of Helsinki. The study was approved by UCLA IRB (#13000047 and #11001274). All donors provided written informed consent for the collection of peripheral blood and subsequent analysis. All leprosy patients were recruited with approval from the IRB of University of Southern California School of Medicine and the Institutional Ethics Committee of Oswald Cruz Foundation, as well as UCLA.

**Author contributions**

RLM and ATD designed the experiments, interpreted the data, and drafted the manuscript. RMBT, PTL, MG, and BRB assisted with the conceptual framework and writing. ATD performed most of the experiments. RMBT, KP, and GC performed the autophagy immunoblotting experiment. RMBT quantified confocal imaging. AC and AL assisted with T cell cultures. ENS and MTO provided the clinical samples.
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