Rapid Nutritional Remodeling of the Host Cell upon Attachment of Legionella pneumophila

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Upon entry of Legionella pneumophila into amoebas and macrophages, host-mediated farnesylation of the AnkB effector enables its anchoring to the Legionella-containing vacuole (LCV) membrane. On the LCV, AnkB triggers docking of K48-linked polyubiquitinated proteins that are degraded by the host proteasomes to elevate cellular levels of amino acids needed for intracellular proliferation. Interference with AnkB function triggers L. pneumophila to exhibit a starvation response and differentiate into the nonreplicative phase in response to the basal levels of cellular amino acids that are not sufficient to power intracellular proliferation of L. pneumophila. Therefore, we have determined whether the biological function of AnkB is temporally and spatially triggered upon bacterial attachment to the host cell to circumvent a counterproductive bacterial differentiation into the nonreplicative phase upon bacterial entry. Here, we show that upon attachment of L. pneumophila to human monocyte-derived macrophages (hMDMs), the host farnesylation and ubiquitination machineries are recruited by the Dot/Icm system to the plasma membrane exclusively beneath sites of bacterial attachment. Transcription and injection of ankB is triggered by attached extracellular bacteria followed by rapid farnesylation and anchoring of AnkB to the cytosolic side of the plasma membrane beneath bacterial attachment, where K48-linked polyubiquitinated proteins are assembled and degraded by the proteasomes, leading to a rapid rise in the cellular levels of amino acids. Our data represent a novel strategy by an intracellular pathogen that triggers rapid nutritional remodeling of the host cell upon attachment to the plasma membrane, and as a result, a gratuitous surplus of cellular amino acids is generated to support proliferation of the incoming pathogen.

The Legionnaires’ disease-causing bacterium, Legionella pneumophila, replicates within alveolar macrophages, causing pneumonia (1). The organism is transmitted to humans from the aquatic environment, where L. pneumophila replicates within amoebas and ciliates (for recent reviews, see references 2, 3, 4, and 5). Coevolution and adaptation of L. pneumophila to the intracellular lifestyle within amoebas in the aquatic environment is believed to have played a major role in its ability to exploit evolutionarily conserved eukaryotic processes, which enables its proliferation within human alveolar macrophages (2–4, 6, 7). Within both evolutionarily distant host cells, L. pneumophila evades endocytic fusion and intercepts endoplasmic reticulum (ER)-to-Golgi vesicle traffic to remodel its phagosome into an ER-derived vacuole, designated the Legionella-containing vacuole (LCV) (1, 3, 8, 9). During late stages of intracellular proliferation, the bacteria escape from the LCV into the cytosol, where they finish the last 1 or 2 rounds of proliferation, during which the bacteria exhibit a starvation response, presumably in response to the depletion of nutrients (10–13).

Modulation of various cellular processes by L. pneumophila is dependent on a functional Dot/Icm type IVB secretion system (14, 15). This system injects into the host cell a cadre of ~300 effectors to modulate a myriad of cellular processes to reprogram the host cell into a proliferation niche (1, 8, 9, 16). However, the roles of most of the Dot/Icm-translocated effectors in the intracellular infection remain unknown, and only few have a detectable role in intracellular replication (1). The AnkB effector, which is found in all genome-sequenced L. pneumophila strains and in 211/217 tested strains (17), is essential for proliferation of L. pneumophila within mammalian and protozoan cells and for intrapulmonary bacterial proliferation and manifestation of pulmonary disease in the mouse model (18–23). Most of the structure of AnkB is composed of eukaryotic domains or motifs, including an F-box domain involved in polyubiquitination, two Ankyrin protein-protein interaction domains (24), and a C-terminal “CaaX” farnesylation motif (3, 25). In contrast to what is seen in the AA100/130b strain and other genome-sequenced strains of L. pneumophila, the AnkB homologue of the Paris strain is missing the farnesylation motif due to truncation of the C terminus, which likely explains functional differences and the various levels of attenuation of the ankB mutant among the different strains (26).

Host-mediated prenylation by farnesylation of AnkB is essential for anchoring AnkB of strain AA100 to the LCV membrane, which is essential for biological function of the effector in macrophages and amoebas and for virulence in mice (22, 27). Prenylation is a eukaryotic posttranslational modification that covalently links a 15-carbon farnesyl or 20-carbon geranyl-geranyl lipid moiety to a conserved cysteine residue within the C terminus “CaaX” motif of a protein, which enables anchoring of hydrophilic proteins into the lipid bilayer of membranes (25). Farnesylation (15-carbon farnesyl addition) of AnkB allows its anchoring into the outer leaflet of the LCV membrane (22). Many other C terminus CaaX motif-containing effectors of L. pneumophila have been shown to be anchored to host membranes through host-mediated prenylation (28, 29). In silico genomic analyses have shown the presence of the “CaaX” motif in numerous proteins of unknown function or protein effectors of other pathogens (22, 25), suggest-
ing the potential of a general paradigm of exploiting host prenylation to anchor bacterial effectors into host membranes. The eukaryotic enzymes necessary for farnesylation, i.e., farnesyl transferase (FTase), Ras-converting enzyme 1 (RCE1), and isoprenyl cytochrome carboxyl methyltransferase (IcmT), are all recruited to the LCV in a Dot/Lcm-dependent fashion (22). Anchoring of AnkB into the LCV membrane via farnesylation is indispensable for its biological activity, since substitution of the cysteine residue within the CaaX motif, RNA interference (RNAi) knockdown, or chemical inhibition of the host FTase abolishes intracellular replication of *L. pneumophila* (22, 27).

The host SCF1 E3 ubiquitin ligase complex contains the RING domain protein RBX1 (RING-box 1), Cul1 (cullin 1), and Skp1 (S-phase-kinase associated protein 1), which bind to the F-box domain protein RBX1 (RING-box 1), Cul1 (cullin 1), and Skp1 anchored AnkB functions as a platform for the assembly of K48-linked polyubiquitinated proteins (21, 23, 27, 39) that are degraded intracellularly. The AnkB interaction with a protein-protein interaction domain such as leucine-rich repeat (LRR), WD40, or ankyrin domain. The AnkB effector of *L. pneumophila* is a bona fide F-box protein that interacts with the SCF1 ubiquitin ligase of amoebas and mammals (21, 23, 27, 31). Deletion of the F-box domain of AnkB or substitution of two of its conserved residues abolishes AnkB-Skp1 interaction (21). Furthermore, knockdown of Skp1 expression by RNAi restricts intracellular proliferation of *L. pneumophila* (21).

Based on nutrient availability, growth of *L. pneumophila* cycles between two phases, which are the motile nonreplicative and the nonmotile replicative phases (33). In response to nutrient depletion during late stages of infection within macrophages and amoebas, *L. pneumophila* exhibits a dramatic starvation response manifested by triggering expression of RelA and SpoT (32), both of which synthesize the alarmone ppGpp (34). The elevated ppGpp levels in *L. pneumophila* trigger a complex network of regulatory cascades that leads to differentiation into the nonreplicative phase associated with dramatic phenotypic modulation, such as flagellation (12, 33–38).

Within amoebas and mammalian cells, the LCV membrane-anchored AnkB functions as a platform for the assembly of K48-linked polyubiquitinated proteins (21, 23, 39) that are degraded by the host prosomal machinery (32), which is required for intracellular proliferation (32, 39). This generates short peptides (2 to 24 amino acids) that are rapidly degraded by host cytosolic oligo- and aminopeptidases into free amino acids. This increases the cellular levels of amino acids, particularly the limiting ones, such as Cys (40), which is semimissential in mammals and is essential for amoebas (32). Cysteine is an essential amino acid for *L. pneumophila*, which converts it to pyruvate and metabolizes it through the tricarboxylic acid (TCA) cycle as a major and metabolically preferable source of carbon and energy to power intracellular replication (32). Upon entry of the wild-type (WT) strain into proteasome-inhibited cells or entry of the ankB mutant into untreated cells, the bacteria exhibit a dramatic starvation response manifested by triggering expression of RelA and SpoT (32), both of which synthesize the alarmone ppGpp (34). However, the starvation response and the associated differentiation exhibited by the ankB mutant upon entry into untreated host cells or by the wild-type strain upon entry into proteasome-inhibited cells are totally circumvented by amino acid supplementation (32). Therefore, to prevent a starvation response and differentiation into a nonreplicative phase by *L. pneumophila*, we hypothesize that AnkB is likely to function rapidly upon contact to the host cell to generate a gratuitous surplus of cellular amino acids, which are the major sources of carbon and energy generation for *L. pneumophila*.

To test our hypothesis, we have determined the early temporal and spatial manipulation of the host cell by AnkB to circumvent such an early starvation response and differentiation into the nonreplicative phase by *L. pneumophila*. Here, we show that upon initial intimate contact with human monocyte-derived macrophages (hMDMs), *L. pneumophila* injects native AnkB, which becomes anchored into the cytosolic side of the plasma membrane exclusively beneath sites of bacterial attachment. Membrane anchoring of the injected AnkB effector is mediated by the host farnesylation machinery, which is rapidly recruited to the plasma membrane, in a Dot/Lcm-dependent manner, beneath sites of bacterial attachment. The host SCF1 ubiquitin ligase is also recruited to the plasma membrane beneath attached extracellular *L. pneumophila*, with subsequent AnkB-dependent rapid proteasomal degradation of K48-linked polyubiquitinated proteins leading to a rapid rise in the cellular levels of amino acids. Taken together, attached extracellular *L. pneumophila* utilizes the AnkB effector to trigger rapid nutritional preparation of the host cell through a rapid rise in the cellular levels of amino acids upon bacterial entry. This suppresses a potential bacterial starvation response, triggers differentiation into the replicative phase, and provides a gratuitous source of carbon and energy needed for robust intracellular proliferation.

**MATERIALS AND METHODS**

**Bacterial strains, cell cultures, and plasmids.** *L. pneumophila* strain AA100/130b (ATCC BAA-74) and the isogenic ankB and dotA mutants were grown on buffered charcoal yeast extract (BCYE) agar plates for 3 days at 37°C prior to use in infections as described previously (20). hMDMs and U937 cells were cultured using RPMI 1640 medium as we described previously (21). The plasmid PXDC61M, which contains the blaM gene, encoding the mature form of TEM-1 beta-lactamase, was obtained from Zhao-Qing Luo at Purdue University. The ankB gene was PCR amplified with restriction enzymes and cloned in frame with the bacterial entry. This-suppresses a potential bacterial starvation response, triggers differentiation into the replicative phase, and provides a gratuitous source of carbon and energy needed for robust intracellular proliferation.

**Real-time qPCR.** Quantitative real-time PCR (qPCR) on attached bacteria was performed as we described previously (11, 20). Briefly, hMDMs were plated at a density of 5 × 10^5 in 24-well plates and treated with 1 μM cytochalasin D for 30 min prior to infection. The hMDMs were then infected with WT bacteria at a multiplicity of infection (MOI) of 10 for 0, 7.5, or 15 min, and immature and synchronized attachment was achieved by centrifugation at 1,000 rpm for 3 min. To assess RNA expression levels of ankB, monosP, and 16S RNA gene in response to attachment to hMDMs, total RNA was extracted from infected cells at the indicated time points using the RNeasy Minikit (Qiagen, Valencia, CA) as recommended by the manufacturer. Total RNA was treated with DNase I (Invitrogen, CA) and random primers. Real-time qPCR was done in triplicate using the Power SYBR green PCR Master Mix kit in a 20-μl reaction volume, as recommended by the manufacturer (Applied Biosystems, CA), using specific primers. The
PCR conditions were 2 min at 94°C initially, followed by 10 s at 96°C, 20 s at 47°C, and 15 s at 72°C for 40 cycles. Changes in mRNA expression were determined by the comparative $C_T$ method (threshold cycle number at the cross point between amplification plot and threshold), and values were normalized to 16S RNA. Negative or positive values were considered downregulation or upregulation when there was a minimum of 2-fold difference of gene expression.

Preferential plasma membrane permeabilization and loading of the cytosol with antibodies. Human monocyte-derived macrophages were isolated and maintained as described previously (20). Monocytes were seeded in 24-well plates at $1 \times 10^6$ cells/well. Cells were treated with cytochalasin D (5 μg/ml), an actin polymerization inhibitor, prior to infection and throughout the experiment. hMDM plasma membranes were selectively permeabilized for 5 min in room temperature with an RPMI 1640 solution containing digitonin (50 μg/ml) as well as anti-AnkB antiserum, as we described previously (41, 42). Following permeabilization, cells were extensively washed with medium and infected with wild-type *L. pneumophila* as well as the ankB and dotA mutant strains at an MOI of 25, 50, or 75 for 15 min. The hMDMs were extensively washed with medium and incubated an additional 30 min for antibody-antigen interaction. Cells were then fixed with 3.7% formaldehyde for 15 min at room temperature. To ensure that cytochalasin D inhibited phagocytosis and bacteria remained extracellular, antibody labeling of *L. pneumophila* with specific rabbit polyclonal antisera was performed prior to permeabilization, followed by Alexa Fluor 488 donkey anti-rabbit secondary antibody (Invitrogen). Cells were then permeabilized with 0.1% Triton X-100 for 10 min at room temperature, followed by anti-AnkB antiserum detection by Alexa Fluor 555-conjugated donkey anti-rabbit IgG (Invitrogen).

**TEM transfection assay.** The U937 cells were grown in RPMI 1640 containing 10% fetal bovine serum (FBS), seeded in black clear-bottom 96-well plates at $1 \times 10^5$ cells/well, and treated with phorbol myristate acetate (PMA) for 24 h prior to infection. *L. pneumophila* strains containing the TEM-1 fusion proteins were grown for 3 days on BCYE containing chloramphenicol (5 μg/ml) and then streaked onto BCYE containing chloramphenicol and 0.5 mM IPTG to induce expression of the fusion proteins. Cell monolayers were loaded with the β-lactamase substrate CCF4 by adding 20 μl of 6× CCF4-AM solution (LiveBLAzer-FRET B/G loading kit; Invitrogen) containing 0.1 M probenecid. Cells were incubated with the solution for 2 h at room temperature. U937 cells were treated with the actin polymerization inhibitor cytochalasin D (5 μg/ml) for 10 min prior to infection and maintained throughout the infection using different MOIs. Plates were centrifuged (1,000 rpm, 5 min) to initiate bacterium-cell contact and incubated for 10 min. Fluorescence was quantified on a BioTek Synergy HT microplate reader with excitation at 405 nm, and emission was detected at 460 nm and 530 nm. Bacterial effector translocation was determined by the emission ratio 460 nm/530 nm to normalize the β-lactamase activity to noninfected substrate loaded cells.

**Recruitment of host farnesyls and ubiquitination machinery to sites of *L. pneumophila* attachment.** A total of $5 \times 10^5$ hMDMs on glass coverslips in 24-well plates were pretreated for 30 min with cytochalasin D (5 μg/ml) and then infected with wild-type *L. pneumophila* and the ankB and dotA mutants at an MOI of 10 for 15 min. Processing of infected cells for confocal microscopy was performed as described previously (21). Briefly, fixed and permeabilized cells were blocked for 1 h with 3% bovine serum albumin (BSA)–phosphate-buffered saline (PBS) and then mouse anti-L. pneumophila antisera (dilution, 1/1,000), and anti-Skp1, anti-Cul1, anti-FTα, anti-RCE1, and anti-IcmT antibodies (dilution, 1/200) (Abcam, Cambridge, MA) were added to 3% BSA–PBS and incubated at room temperature for 1 h. Following extensive washing with 3% BSA–PBS, bound antibodies were detected with Alexa Fluor 488- or 555-conjugated donkey anti-rabbit or mouse IgG antibodies (Invitrogen, Carlsbad, CA) for 1 h. Following this, the glass coverslips were mounted on glass slides using ProLong Gold antifade reagent (Invitrogen, Carlsbad, CA). The fixed cells were examined with an Olympus FV1000 laser scanning confocal microscope as we described previously (21). On average, 8 to 15 0.2-μM serial Z sections of each image were captured and stored for further analyses, using Adobe Photoshop CS5.

**GC-MS analyses of free amino acids.** The cellular levels of free amino acids were determined as part of the global metabolomics profile. The hMDMs were seeded in 6-well plates at $1 \times 10^5$ cells/well, and prior to infection, the cells were treated with 1 μM cytochalasin D for 30 min. The hMDMs were infected with WT or ankB mutant *L. pneumophila* at an MOI of 100 for 1 h, and the infected cells were lysed in aqueous 90% methanol. Lysates were stored at $-20^\circ$C for 1 h and then centrifuged (21,000 x g at 4°C) for 10 min. The resulting supernatants were dried using a Speed-Vac and prepared for gas chromatography–mass spectrometry (GC-MS).

All GC-MS analyses were performed at the University of Utah Metabolomic core facility using a Waters GCT Premier mass spectrometer fitted with an Agilent 6890 gas chromatograph and a Gerstel MPS2 autosampler, as we described previously (32). The dried samples were suspended in 40 μl of 40-mg/ml O-methoxylamine hydrochloride in pyridine and incubated for 1 h at 30°C. A 25-μl sample of this solution was transferred to autosampler vials followed by the addition of N-methyl-N-trimethylsilyl trifluoroacetamide and further incubated for 30 min at 37°C with shaking. A 1-μl sample was injected to the gas chromatograph inlet in the split mode set to a 10:1 ratio. Injector temperature was held at 250°C. The gas chromatograph had an initial temperature of 95°C for 1 min followed by a 40°C/min ramp to 110°C with a hold time of 2 min. This was followed by a second 5°C/min ramp to 250°C and then a third ramp to 350°C and a final hold time of 3 min. A 30-m Restek Rxi-5 MS column with a 5-ml guard column was employed for analysis. Data were collected by MassLynx 4.1. Data analysis for free cellular amino acids was performed using QuanLynx, which quantified the area under the curve for each amino acid. All data were saved to an Excel spread sheet for further analysis. The GC-MS analysis gives relative results of the area under the curve for the same amino acid and is not quantitative relative to other amino acids within the same sample. Since the GC-MS analyses compare levels of the same metabolite/amino acid between different samples/treatments, the results are presented as ratios of infected to uninfected cells.

**RESULTS**

**Triggering expression of ankB upon attachment of *L. pneumophila* to human macrophages.** Upon entry of the ankB mutant to macrophages or amoebas, the bacterium exhibits a dramatic starvation response and differentiation into the nonreplicative phase, but both phenotypes are circumvented by amino acid supplementation (32). Similar phenotypes are also exhibited by the wild-type strain upon entry into proteasome-inhibited cells, and in both cases the respective phenotypes are circumvented upon supplementation of amino acids (32). We tested the hypothesis that WT *L. pneumophila* likely employs AnkB during initial stages of interaction with the host cell to circumvent the amino acid starvation response and the associated phenotypic modulations. We determined whether attachment of *L. pneumophila* to host cells triggered expression of AnkB. To determine this, human monocyte-derived macrophages (hMDMs) were pretreated with cytochalasin D for 30 min to block phagocytosis and then infected with wild-type *L. pneumophila* at an MOI of 10 for 0, 7.5, and 15 min. This bacterial attachment protocol resulted in attachment of ~50% of the cells in the monolayers. Blocking of bacterial entry into hMDMs by cytochalasin D was confirmed by sterilization of the infected cytochalasin D-treated monolayers by gentamicin treatment, indicating that the bacteria were extracellular. Following RNA purification and cDNA synthesis, expression of the ankB gene was determined by real-time qPCR using expression of the 16S RNA as an internal control, as
we described previously (11, 18, 20). Expression of the constitutively expressed mompS gene was used as a control. Compared to expression of ankB at 0 min postattachment, its expression was increased 9-fold and 26-fold at 7.5 and 15 min, respectively, following bacterial attachment (Fig. 1). During the course of the attachment experiment, no change was observed in expression of mompS (Fig. 1). These data show that transcription of ankB is triggered immediately upon attachment of L. pneumophila to hMDMs.

Translocation of AnkB into macrophages upon bacterial attachment. To determine temporal and spatial translocation of AnkB upon intimate attachment of L. pneumophila to macrophages, we generated an L. pneumophila strain expressing a β-lactamase-AnkB reporter fusion construct to monitor real-time translocation of the native AnkB effector injected by attached extracellular bacteria. Since the β-lactamase-AnkB reporter fusion was clearly injected by attached extracellular bacteria (Fig. 2), our microscopy findings suggested that the injected AnkB was likely localized exclusively beneath bacterial attachment sites.

To overcome the above-mentioned caveat and to determine whether the injected native AnkB by attached extracellular bacteria was located beneath bacterial attachment sites, the cytosol of live hMDMs was preloaded with anti-AnkB antibodies prior to bacterial attachment. This would allow the antibody to bind AnkB upon its injection by attached extracellular bacteria (13, 41, 42). This strategy also provides a clear and more solid interpretation of the data, since the anti-AnkB antibody is loaded to the host cell cytosol prior to inoculation of the bacteria. To load the host cell cytosol with the anti-AnkB antibodies prior to infection, the plasma membrane of live hMDMs was preferentially permeabilized with a low concentration of digitonin (13, 41, 42). After loading the cells with the antibody and allowing the cells to heal the membrane damage for few minutes, the integrity of the plasma membrane was confirmed by impermeability to trypan blue. Without digitonin treatment, the plasma membrane of hMDMs was impermeable to anti-AnkB antibodies, as expected (Fig. 3). The antibody-loaded hMDMs were treated with cytochalasin D to prevent phagocytosis and then infected at an MOI of 10 with WT L. pneumophila or the isogenic mutant dotA or ankB. The cells were then fixed and processed for confocal microscopy. This bacterial attachment protocol resulted in an average attachment of 1 or 2 bacteria/cell in ~50% of the cells in the monolayers. To allow differentiation between extracellular and intracellular bacteria, extracellular L. pneumophila cells were labeled with specific antibodies prior to permeabilization of the infected cells. When the hMDMs were permeabilized with digitonin and loaded with anti-AnkB antibodies, the loaded antibody was detectable as red fluorescence.
patches throughout the cytosol of ∼98% of the cells, indicating successful loading of the cells with anti-AnkB antibodies to prior to infection (Fig. 3). When hMDMs were infected with wild-type \textit{L. pneumophila}, 52% of attached extracellular bacteria colocalized with AnkB exclusively beneath the site of bacterial attachment (Fig. 3). As expected, the dotA translocation-defective mutant and the \textit{ankB} mutant did not colocalize with AnkB (5% and 0%, respectively) (Student’s \( t \) test, \( P < 0.007 \) and 0.003, respectively) (Fig. 3).

Since AnkB is hydrophilic and its anchoring to the LCV membrane is mediated by host farnesylation within amoebas and macrophages (22, 25), we determined whether host farnesylation was required for the exclusive localization of AnkB to the cytosolic side of the plasma membrane beneath the sites of bacterial attachment. The cytosol of hMDMs was preloaded with anti-AnkB antisera prior to infection, as described above. Cytochalasin-D treated cells were infected by the farnesylation-defective \textit{ankB} \(_{169C-A}\) substitution mutant in the CaaX motif (22). The data showed that infection by the \textit{ankB} \(_{169C-A}\) substitution mutant resulted in failure to anchor AnkB to the plasma membrane beneath bacterial attachment sites (Student’s \( t \) test, \( P < 0.005 \)) (Fig. 3). Therefore, host farnesylation anchors the injected AnkB by attached extracellular \textit{L. pneumophila} to the cytosolic side of the plasma membrane directly and exclusively beneath bacterial attachment sites. This is the first demonstration of farnesylation-mediated anchoring of an injected bacterial effector to the inner leaflet of the plasma membrane beneath bacterial attachment sites.

**Recruitment of the host farnesylation machinery to the plasma membrane beneath attached extracellular \textit{L. pneumophila}**. Since during infection host-mediated farnesylation of AnkB anchors it to the LCV membrane and AnkB was exclusively localized beneath bacterial attachment sites, we tested the hypothesis that the host farnesylation enzymes FTase, IcmT, and RCE1 were recruited to the plasma membrane by attached extracellular bacteria to anchor AnkB to the plasma membrane. Cytochalasin D-treated hMDMs were infected with wild-type \textit{L. pneumophila} and the isogenic mutants \textit{ankB} and \textit{dotA} at an MOI of 10 for 15 min. The cells were then immediately fixed and processed for confocal microscopy to determine if FTase, IcmT, and RCE1 were recruited beneath the sites of bacterial attachment. The data showed that FTase, IcmT, and RCE1 were all recruited beneath attachment sites of wild-type bacteria at a frequency of ∼85% (Fig. 4A, B, and C). In contrast, FTase, IcmT, and RCE1 were recruited at a frequency of only ∼10%, beneath attachment sites of the \textit{dotA} mutant (Student’s \( t \) test, \( P < 0.006, 0.007, \) and 0.006, respectively) (Fig. 4). FTase, IcmT, and RCE1 were recruited at a significantly reduced frequency of 43, 47, and 41%, respectively (Student’s \( t \) test, \( P < 0.01 \)), beneath attachment sites of the \textit{ankB} mutant bacteria (Fig. 4). The moderate reduction in recruitment of the host enzymes by the \textit{ankB} mutant is most likely due to the fact that the other ∼12 farnesylated effectors of \textit{Legionella} injected by the \textit{ankB} mutant (28, 29) interact with the host farnesylation enzymes, while the translocation-defective \textit{dotA} mutant is severely defective in recruitment of the host enzymes (22). These data indicate that upon attachment of \textit{L. pneumophila} the Dot/Icm apparatus is essential for recruitment of the host enzymes into the plasma membrane beneath the sites of bacterial attachment. This is the first example of recruitment of the host farnesylation machinery by attached extracellular bacteria to anchor an injected bacterial effector to the inner leaflet of the plasma membrane beneath bacterial attachment sites.
effector to the cytosolic side of the plasma membrane exclusively beneath bacterial attachment sites.

Recruitment of the host SCF1 ubiquitin ligase complex beneath attached extracellular *L. pneumophila*. During ectopic expression, the AnkB effector interacts with the host Skp1 component of the SCF1 E3-ubiquitin ligase complex, but the location of this interaction during infection is not known (21, 23). Since AnkB is exclusively localized to the LCV membrane during infection and to the plasma membrane beneath attached extracellular *L. pneumophila*, we tested the hypothesis that the SCF1 ubiquitin ligase was recruited to the LCV, where it interacts with AnkB, and that this recruitment was rapidly initiated at the plasma membrane beneath bacterial attachment sites. To determine recruitment of SCF1 to the LCV, hMDMs were infected at an MOI of 10 for 1 h with wild-type *L. pneumophila* or the isogenic *dotA* or *ankB* mutant defective isogenic mutant. The data showed that both Skp1 and Cul1 components of the SCF1 were recruited to the LCV of the WT strain at a frequency of 82 to 84% (Fig. 5). Recruitment of both host cell components was dependent on a functional Dot/Icm translocation system, since Skp1 and Cul1 were recruited at a frequency of only ~10% by attached translocation-defective *dotA* mutant bacteria (Student’s *t* test, *P* < 0.008) (Fig. 5). Only 34 and 30% of attached *ankB* mutant bacteria recruited Skp1 and Cul1, respectively (Fig. 5), which was significantly less than wild-type bacteria (Student’s *t* test, *P* < 0.01). These data show that the Dot/Icm translocation system of *L. pneumophila* is essential for recruitment of the SCF1 ubiquitin ligase to the LCV and that this recruitment is initiated at the plasma membrane beneath sites of bacterial attachment. This is the first example of recruitment of the host SCF1 to a pathogen-containing vacuole and the initiation of this process at the cytosolic side of the plasma membrane beneath bacterial attachment sites.

Elevated levels of cellular amino acids triggered by attached extracellular *L. pneumophila*. The ultimate function of the LCV membrane-anchored AnkB effector is to generate high levels of cellular amino acids through host proteasomal degradation of K48-linked polyubiquitinated proteins (32). Therefore, we determined whether the injected AnkB by attached extracellular *L. pneumophila* resulted in elevated levels of cellular amino acids through degradation of the polyubiquitinated proteins assembled beneath bacterial attachment sites (21). To achieve this, cytochalasin D-treated hMDMs were infected by the wild-type strain or the isogenic *ankB* mutant *L. pneumophila*. Trypan blue staining of the cells showed that there was no detectable effect of cytochalasin D on permeability of the plasma membrane. The hMDMs were lysed, and the relative levels of free amino acids were determined by GC-MS. The data showed that attached wild-type *L. pneumophila* triggered a rapid rise in the levels of amino acids, relative to uninfected cells (Student’s *t* test, *P* < 0.001) (Fig. 6). In contrast, attachment of the *ankB* mutant bacteria to hMDMs did

![FIG 4](http://iai.asm.org/) The farnesylation machinery components FTα, RCE1, and IcmT are recruited beneath attachment sites of *L. pneumophila* to hMDMs. The hMDMs pretreated with cytochalasin D were infected by wild-type (WT) *L. pneumophila* and the isogenic *dotA* or *ankB* mutant for 15 min. Representative confocal microscopy images of infected hMDMs showing colocalization of FTα (A), RCE1 (B), and IcmT (C) proteins to attached WT, *ankB* mutant, or *dotA* mutant bacteria. Bacteria were labeled with anti-Lpn antibody (green), and FTα, RCE1, and IcmT were labeled with the respective specific antibodies (red) and then analyzed by confocal microscopy. The arrowheads indicate intense colocalization of FTα, RCE1, or IcmT with the WT strain. The numbers in the merged images of all panels are means and standard deviations of the frequency of recruitment of FTα, RCE1, or IcmT beneath attached extracellular bacteria. The data represent analyses of 100 infected cells and are representative of three independent experiments.
not alter cellular levels of amino acids relative to uninfected cells (Fig. 6). Thus, translocation of AnkB by attached extracellular L. pneumophila results in increased levels of cellular amino acids, which are needed to block a potential starvation response and differentiation of L. pneumophila into the nonreplicative phase (see model in Fig. 7). This is the first example of a strategy by an intracellular pathogen to trigger rapid nutritional remodeling of the host cell upon attachment to the plasma membrane, and as a result, a gratuitous surplus of cellular amino acids is generated to support proliferation of the incoming pathogen.

DISCUSSION

Upon entry of WT L. pneumophila into proteasome-inhibited cells or entry of the ankB mutant into untreated cells, both populations of bacteria exhibit a dramatic starvation response (32), which leads to bacterial differentiation into the nonreplicative phase (34, 35), and these differentiations are totally overcome upon supplementation of amino acids (32). Based on these observations, we tested the hypothesis that the function of AnkB is likely to be rapidly triggered upon attachment of L. pneumophila to the host cell to ensure that levels of amino acids in the host cell are sufficient to prevent a starvation response and a counterproductive differentiation into the nonreplicative motile phase. Our data show that both expression and translocation of AnkB are critically dependent on AnkB, since other farnesylated Dot/Icm-translocated effectors are likely to be involved in recruitment of the farnesylases to the cytosolic side of the plasma membrane exclusively beneath bacterial attachment sites.

We have previously demonstrated that the farnesylases cytosolic FTase and ER-bound RCE1 and IcmT localize to the LCV in a Dot/Icm-dependent manner (22). Our data show that all three farnesylases are rapidly recruited to the plasma membrane beneath bacterial attachment sites (Fig. 7). Recruitment of these enzymes is Dot/Icm dependent but is only partially dependent on AnkB, since other farnesylated Dot/Icm-translocated effectors are likely to be involved in recruitment of the farnesylases machinery at the plasma membrane (28, 29).

During infection, the host SCF1 ubiquitin ligase is recruited to the LCV, indicating that AnkB-SCF1 interaction occurs on the LCV membrane (21, 27). Importantly, recruitment of SCF1 is initiated at the inner leaflet of the plasma membrane beneath bacterial attachment sites, similar to the host farnesyl machinery (Fig. 7). This recruitment is only partially dependent on AnkB, indicating that other L. pneumophila F-box proteins (23, 31, 48, 49) are likely to be involved in recruitment of the SCF1 complex to the plasma membrane beneath sites of bacterial attachment. The ability of L. pneumophila to rapidly recruit both the farnesylases machinery and the E3 SCF1 ubiquitin ligase machinery to the cytosolic side of the plasma membrane beneath sites of bacterial attachment allows AnkB to assemble K48-linked polyubiquitinated proteins on the plasma membrane beneath bacterial attachment sites (Fig. 7) (21, 32). Proteasomal degradation of the AnkB-assembled K48-linked polyubiquitinated proteins triggered by attached bacteria, AnkB is the first L. pneumophila effector known to be exclusively anchored to the cytosolic side of the plasma membrane beneath the sites of bacterial attachment (see model in Fig. 7). Upon phagocytosis and initial formation of the phagosomal membrane from the plasma membrane, AnkB remains exclusively anchored to the outer leaflet of the LCV membrane (22), which is derived from the inner leaflet of the plasma membrane. Specific anchoring of AnkB to the inner leaflet of the plasma membrane upon bacterial attachment is dependent on eukaryotic farnesylation of the cysteine residue within the C-terminal CaaX motif of AnkB (22). Anchoring of AnkB to the plasma membrane by attached extracellular bacteria is essential for the biological function of AnkB (see model in Fig. 7). It would be interesting to determine the subcellular location of the AnkB homologue of the Paris strain of L. pneumophila (23), which lacks the farnesylation motif, upon bacterial attachment as well as after formation of the LCV. It is most likely that the structural differences in AnkB between various strains of L. pneumophila account for functional differences and role of the effector in intracellular replication (26), similar to what has been recently shown for the AnkI effector (19) (LegAS4/RomA) of two different strains of L. pneumophila (26, 46, 47). However, genomic redundancy may also play a factor (1, 9). Nevertheless, AnkB is the first example of an injected effector by attached extracellular L. pneumophila in which host-mediated farnesylations anchors it to the cytosolic side of the plasma membrane exclusively beneath bacterial attachment sites.
The limiting amino acid in eukaryotes (40) but ironically is a metabolically preferable source of carbon and energy for L. pneumophila. The availability of higher levels of cell amino acids upon entry of L. pneumophila circumvents a potential starvation response and differentiation into the motile nonreplicative phase for the entering bacterium. The elevated cellular levels of amino acids trigger differentiation of L. pneumophila into the replicative phase and are major sources of carbon and energy that feed the TCA cycle to power intracellular bacterial proliferation.

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