

Distinct Roles in Autophagy and Importance in Infectivity of the Two ATG4 Cysteine Peptidases of *Leishmania major*^{*}

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Background: ATG4 is a cysteine peptidase crucial for macroautophagy.

Results: Gene deletion mutants show that the two ATG4s of *Leishmania* perform distinct roles, although there is some redundancy.

Conclusion: ATG4s are not individually essential but macroautophagy, a process important in the virulence of the parasite, requires one.

Significance: Highlights the distinct roles of ATG4 isoforms and their importance for autophagy and parasite infectivity.

Macroautophagy in *Leishmania*, which is important for the cellular remodeling required during differentiation, relies upon the hydrolytic activity of two ATG4 cysteine peptidases (ATG4.1 and ATG4.2). We have investigated the individual contributions of each ATG4 to *Leishmania major* by generating individual gene deletion mutants ($\Delta atg4.1$ and $\Delta atg4.2$); double mutants could not be generated, indicating that ATG4 activity is required for parasite viability. Both mutants were viable as promastigotes and infected macrophages *in vitro* and mice, but $\Delta atg4.2$ survived poorly irrespective of infection with promastigotes or amastigotes, whereas this was the case only when promastigotes of $\Delta atg4.1$ were used. Promastigotes of $\Delta atg4.2$ but not $\Delta atg4.1$ were more susceptible than wild type promastigotes to starvation and oxidative stresses, which correlated with increased reactive oxygen species levels and oxidatively damaged proteins in the cells as well as impaired mitochondrial function. The antioxidant *N*-acetylcysteine reversed this phenotype, reducing both basal and induced autophagy and restoring mitochondrial function, indicating a relationship between reactive oxygen species levels and autophagy. Deletion of ATG4.2 had a more dramatic effect upon autophagy than did deletion of ATG4.1. This phenotype is consistent with a reduced efficiency in the autophagic process in $\Delta atg4.2$, possibly due to ATG4.2 having a key role in removal of ATG8 from mature autophagosomes and thus facilitating delivery to the lysosomal network. These findings show that there is a level of functional redundancy between the two ATG4s, and that ATG4.2 appears to be the more important. Moreover, the low infectivity of $\Delta atg4.2$ demonstrates that autophagy is important for the virulence of the parasite.

Macroautophagy (hereafter autophagy) is an intracellular process that sequesters cytosol and organelles within autophagosomes for delivery to and degradation within lysosomes. One

main physiological significance of this process in most cell types is recycling of intracellular materials during nutrient deprivation, but in addition it plays important parts in cellular differentiation, tissue remodeling, growth control, size regulation, mitochondrial homeostasis, cellular immunity, adaptation to stresses, and unconventional protein secretion (1–6). The many steps comprising autophagy rely on a large repertoire of proteins mainly from the ATG² family, of which more than 30 have been described. Central to the process are the ATG5-ATG12 conjugation and ATG8-lipidation pathways, which are coordinated by the activity of the ATG6-Vps34 complex (7–9).

ATG4, a cysteine peptidase of Clan CA, family C54, is a key component of the ATG8-lipidation pathway and is essential for autophagosome biogenesis. ATG4 is responsible for cleavage of ATG8 after a glycine residue close to its C terminus, facilitating conjugation of ATG8 to the lipid phosphatidylethanolamine (PE). ATG8-PE attaches to the phagophore and is involved in elongation of the phagophore and closure of the autophagosome. Subsequently, ATG4 cleaves ATG8 from the PE on the outside of the autophagosome membrane, the reverse of the initial conjugation reaction (10–13), which allows interaction and fusion of the autophagosome with the endosomal-lysosomal compartments (14–16). The conjugation and deconjugation activities of ATG4 are crucial for autophagosome genesis and function.

Mammals and plants have multiple orthologues of the single yeast ATG4. There appears to be some functional redundancy (17) but the four ATG4 orthologues in humans have distinct roles in autophagosome genesis and cell survival, which is reflected in their preferred substrates among the six ATG8 orthologues. ATG4B has activity toward all of the ATG8 homologues and is the key regulator for autophagy in humans, its absence has severe consequences in the cell (18–21). ATG4D is predominantly involved in mitophagy and apoptosis (22, 23),

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² The abbreviations used are: ATG/ATG, autophagy related gene/protein; H₂DCF-DA, 2',7'-dichlorodihydrofluorescein diacetate; PE, phosphatidylethanolamine; ROS, reactive oxygen species; NAC, *N*-acetylcysteine; SE, scanning electron microscopy.

whereas ATG4C has restricted substrate specificity and is only required for survival during prolonged stress (19).

Human ATG4A and ATG4B are redox regulated, they are the target for direct but reversible oxidation of a regulatory cysteine residue by reactive oxygen species (ROS), which inactivates the deconjugation activity (24, 25). The catalytic cysteine of ATG4 that processes ATG8 prior to its conjugation to PE is not susceptible to ROS oxidation. Its continued activity ensures that the lipidated ATG8 is formed and, in the absence of the deconjugation activity of ATG4, accumulates within the cell and is available for increased autophagosome biogenesis under conditions of oxidative stress and starvation. Moreover, ROS can up-regulate the expression of ATG4 in a JNK signaling-dependent manner to ensure that an adequate amount of processed ATG8 is available for autophagosome biogenesis after autophagic induction in MEF-7 cells (26). This up-regulation of ATG4 expression can be abrogated by either JNK knock-down or addition of the ROS scavenger *N*-acetylcysteine (NAC). The relationship between ROS and JNK activation was established previously (27, 28), but this link with ATG4 demonstrates a clear correlation between ROS production and the induction of autophagy. Oxidative stress caused by high ROS levels causes extensive and irreversible structural changes to proteins and organelles within cells (29, 30), thus the induction of autophagy is a mechanism to maintain quality control, cellular homeostasis, and cellular energetic balance (24). Further evidence of this link comes from the findings that autophagy-deficient mutants have high ROS levels. Human ATG4C and ATG4D are not redox regulated in the same way and neither are the ATG4s of yeast and plants, suggesting that these latter organisms must deal with ROS stress differently.

Leishmania, the causative agent of important diseases known as the leishmaniasis, which are prevalent in many tropical and subtropical regions, carries out autophagy, comprising two conjugation pathways similar to yeast and higher eukaryotes, which is important for cellular differentiation, mitochondrial function, and phospholipid homeostasis (31–33). It contains two genes encoding ATG4, the two enzymes are designated ATG4.1 and ATG4.2 and share only 19% identity (33). They have differential activity toward the unusual four classes of ATG8 of the parasite (32). ATG4.1 preferentially processes ATG8, ATG8B, and ATG8C, whereas ATG4.2 preferentially cleaves ATG8A (32). This suggests that the two forms of ATG4 have differing roles in the parasite.

The main aim of this study was to determine whether the two ATG4s are individually essential for *Leishmania major* and to elucidate the effects that removal of each ATG4 gene has on the parasite. Our approach using targeted gene deletion has also enabled us to obtain insights into the roles of each enzyme, including the parts that they may play in dealing with ROS, the relationship between ROS and autophagy in *Leishmania*, and the importance of the ATG4s for parasite survival and infectivity.

EXPERIMENTAL PROCEDURES

Parasite Handling Procedures—Promastigotes of *L. major* (MHOM/IL/80/Friedlin, designated WT for this study) were grown in modified Eagle's medium (designated complete

HOMEM medium) with 10% (v/v) heat-inactivated fetal calf serum at 25 °C, as described previously (34). Parasite numbers were estimated using an improved Neubauer hemocytometer. In this report, except when stated otherwise, early log, mid-log, and early stationary phases of promastigote growth correspond to $\sim 5 \times 10^5$, 5×10^6 , and 9×10^6 parasites ml^{-1} , respectively. The following antibiotics were added to the cultures of the *Δatg4* mutants and the derived cell lines as follows: hygromycin B (Sigma) at 50 $\mu\text{g ml}^{-1}$; phleomycin (Cayla, France) at 10 $\mu\text{g ml}^{-1}$, puromycin (Calbiochem) at 10 $\mu\text{g ml}^{-1}$; blasticidin S (Calbiochem) at 10 $\mu\text{g ml}^{-1}$; and neomycin (G418, Geneticin, Invitrogen) at 25 $\mu\text{g ml}^{-1}$.

Metacyclogenesis and Infectivity of *Leishmania* to Macrophages and Mice—Metacyclic promastigotes in *L. major* stationary phase cultures were assessed by using either the peanut agglutinin method (35) or Western blot analysis using antibodies to the metacyclic promastigote-specific protein, HASPB (see below).

Infectivity of *Leishmania* lines to macrophages was determined using peritoneal macrophages from CD1 mice resuspended in RPMI media (Sigma) at $5 \times 10^5 \text{ ml}^{-1}$ and left to adhere onto coverslips overnight. These cells were infected with stationary phase promastigote cultures or amastigotes harvested from mice footpad lesions at a ratio of ~ 2 promastigotes or 0.5 amastigotes per macrophage and incubated for up to 5 days at 32 °C in 5% CO_2 , 95% air. Non-phagocytosed promastigotes or amastigotes were removed after 24 h by washing four times with RPMI. Parasite abundance within the macrophages after 1 and 5 days was determined after the cells were fixed in methanol and stained with Giemsa for 10 min.

The infectivity to mice was determined using groups of 5 mice that were inoculated subcutaneously within a footpad with 5×10^5 stationary phase *L. major* promastigotes or amastigotes, harvested from mice footpad lesions, suspended in 200 μl of PBS, pH 7.4. The thickness of the lesion in infected footpads was measured using a caliper over a 5–12-week period.

Amastigote Isolation from Infected Mice and Transformation to Promastigotes—Amastigotes were excised, into cold PBS containing 50 $\mu\text{g ml}^{-1}$ of gentamycin (Sigma), from footpad lesions of mice, inoculated 5 weeks earlier with 5×10^5 stationary phase promastigotes, after removal of the skin. The lesion material was ground in a glass tissue grinder and centrifuged at $150 \times g$ for 1 min at 4 °C to remove the large debris. The supernatant was then centrifuged at $1700 \times g$ for 15 min and the subsequent pellet was resuspended and treated as required by the design of the experiment.

The number of *L. major* amastigotes in footpad lesions of infected mice was determined using the limiting dilution assay as described previously (36). Briefly, amastigotes of *L. major* were excised from footpads and resuspended in 10 ml of complete HOMEM medium with 50 $\mu\text{g ml}^{-1}$ of gentamycin. The parasite suspension was serially diluted in duplicate flasks, incubated for 5 days at 26 °C, and then inspected daily for parasite growth. For quantifying the morphological forms in these cultures, parasites within the culture flasks were classified according to the following criteria: amastigotes, ovoid to round but lacking an emergent flagellum, or promastigotes, elongated bodies with a flagellum equal to or longer than the cell body

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length. Other parasites were designated as intermediate forms. A minimum of 200 cells was examined and the resulting differential counts were expressed as percentages.

Western Blot Analyses—Parasites were harvested at $1,000 \times g$ for 10 min, washed twice in PBS, and the pellets were either used immediately or stored at -20°C . Parasite lysates used for this analysis were produced by resuspension of parasite pellets in lysis buffer, comprising 0.25 M sucrose, 0.25% Triton X-100, 10 mM EDTA, and a mixture of peptidase inhibitors (10 μM E-64, 2 mM 1,10-phenanthroline, 4 μM pepstatin A, and 1 mM phenylmethylsulfonyl fluoride). After a 10-min incubation in ice, lysates were centrifuged at $15,700 \times g$ for 30 min at 4°C , and the resulting supernatant (designated soluble fraction) was retained for analysis. Protein concentrations were determined according to the Bradford procedure (Bio-Rad) using bovine serum albumin as the protein standard.

Western blot analysis was performed on *L. major* lysates prepared as described above. The primary anti-GFP (Abcam Ltd.) and the secondary anti-rabbit antibodies were used at a 1:2,000 (v/v) and 1:5,000 (v/v) dilutions, respectively. The primary anti-His (Invitrogen) and the secondary anti-mouse antibodies were used at 1:1,000 (v/v) and 1:50,000 (v/v), respectively. To assess metacyclogenesis, the metacyclic promastigote-specific proteins SHERP and HASPB were monitored using anti-SHERP and anti-HASPB antibodies (produced in rabbit) at a dilution of 1:5,000 (v/v). The antibodies were kindly provided by Prof. Deborah Smith (University of York). Loading controls used the anti-TbEF1 α antibody (Upstate) at a dilution of 1:5,000 (v/v). The West-Pico chemiluminescence detection system (Pierce) was used to visualize antigens.

Generation of *L. major* ATG4 Null Mutants and Re-expressing Cell Lines—To generate the *ATG4.1* null mutant, the 1014-bp 5' flank fragment of *ATG4.1* (LmjF32.3890) was amplified by PCR from *L. major* genomic DNA with primers NT164 (5'-AAG CTT CCT CGT CAC CTT CCT CGA TGA CAG GCG GGT-3') and NT165 (5'-GTC GAC TCC GGG GGT GCA GAC GGG GCT GTA G-3'), digested with HindIII and Sall, and inserted into the HindIII/Sall-digested pGL345-HYG (37) to give pGL345ATG4.1-HYG5'. The 3' flank fragment was amplified by PCR with primers NT166 (5'-CCC GGG TCG TAG CGC CAG GAC GCT GAT CGT CAT AGC-3') and NT167 (5'-AGA TCT ATC ACG CTG CAC AGT ATC ACC CGG T-3'). The resulting 1008-bp fragment was digested with SmaI and BglII and cloned into the same restriction sites in the pGL345ATG4.1-HYG5' plasmid to give pGL345ATG4.1HYG5'3'. The cassette used for transfection of *L. major* promastigotes was released by HindIII/BglII digestion. The pGL345ATG4.1BSD5'3' plasmid used for replacement of the second allele was generated from plasmid pGL345ATG4.1HYG5'3' by replacing the SpeI/BamHI ORF of the hygromycin resistance gene with a SpeI/BamHI ORF of the blasticidin S-deaminase gene. A population of parasites resistance to hygromycin was generated after transfection with pGL345ATG4.1HYG5'3' (see below). This population was used for a second round of transfection with the pGL345ATG4.1BSD5'3' construct. Two blasticidin-resistant clones from independent events, designated $\Delta atg4.1a$ and

$\Delta atg4.1b$, were selected for analysis. The generation of the $\Delta atg4.2$ line has been described previously (33).

To generate the lines re-expressing *ATG4.1*, the *ATG4.1* gene modified with a polyhistidine tag at the C-terminal end (to verify protein expression) was inserted in the pRIB-Pur bearing the puromycin resistance gene (38). PCR primers NT158 (5'-CTC GAG ATG GGC ACG AAC GCC AAA GTG GCA GAG-3') and NT159 (5'-ATC GAT CTA ATG ATG ATG ATG GCT CGG TGG AGA AGA GAT TGA ATT-3') produced the polyhistidine-tagged version of *ATG4.1* containing BglII and BamHI sites, respectively. The 1077-bp fragment was digested by BglII/BamHI and cloned into pRIB-Pur vector previously digested with the same restriction enzymes to give the pRIB-PurATG4.1 plasmid. The integration cassette from this plasmid was excised by digestion with PacI and PmeI before electroporation of *L. major* promastigotes. This fragment possesses the 5' and 3' flanking regions for chromosomal integration at the ribosomal locus. Proteins at this locus are constitutively expressed in both promastigote and amastigote life cycle forms (39). This cell line was designated $\Delta atg4.1::ATG4.1$. The pRIB-PurATG4.2 plasmid was made similarly using the polyhistidine *ATG4.2* but was amplified using primers NT160 (5'-CTC GAG ATG GCT CCG CTC CGT GCA AGA TT-3') and NT161 (5'-AGA TCT TCA ATG ATG ATG ATG ATG ATG ATG ATG ATC CAG ATA CTC CCA-3'). The resultant cell line was designated $\Delta atg4.2::ATG4.2$. The creation of cell lines expressing GFP-ATG8 and *ATG4.2* have been described previously (33).

L. major promastigotes were electroporated (NucleofectorTM) with 15 μg of DNA of the extra chromosomal constructs, namely pN-GFP-ATG8 and pN-ATG4.2, and 30 μg of the pRIB-PurATG4.1 or pRIB-PurATG4.2 cassettes. Transfectants were selected with the appropriate antibiotic at the concentrations detailed above. All PCR assays were carried out in a GeneAmp 9600 PCR system (PerkinElmer Life Sciences) for 30 cycles of denaturation (94°C , 15 s), annealing (65°C , 15 s), and extension (72°C , 2 min). For the attempts to generate the *ATG4* double null, the pGL345ATG4.1NEO5'3' and pGL345ATG4.1BLE5'3' plasmids were generated from plasmid pGL345ATG4.1HYG5'3' by replacing the SpeI/BamHI ORF of the hygromycin resistance gene with SpeI/BamHI ORFs of the neomycin resistance and bleomycin resistance genes, respectively.

Southern Blot Analyses—Genomic DNA was extracted according to standard procedures (40). DNA (10 μg) was digested with XhoI, fractionated by agarose gel electrophoresis, nicked, denatured, neutralized, and blotted onto HybondTM-N⁺ membrane (Amersham Biosciences) by capillary transfer. The probe was prepared from a 1014-bp HindIII/Sall 5' flank fragment excised from the T-vector (Promega) using a random priming kit (Prime-It; Stratagene) and purified on Microspin S-200 HR columns (Amersham Biosciences). Filters were hybridized overnight at 65°C with a [α -³²P]dATP-labeled 5' flank fragment probe in Church-Gilbert hybridization solution. Filters were washed under high stringency (15 min at 65°C with $2 \times \text{SSC}$, 0.1% SDS and twice with $0.2 \times \text{SSC}$, 0.1% SDS) and exposed to x-ray film (Konica Medical Film).

Monitoring and Quantification of Autophagy—For live imaging, promastigotes of WT[pN-ATG8], $\Delta atg4.1$ [pN-ATG8],

$\Delta atg4.1::ATG4.1$ [pN-ATG8], $\Delta atg4.2$ [pN-ATG8], and $\Delta atg4.2::ATG4.2$ [pN-ATG8] in HOMEEM growth medium were mounted on coverslips and observed by fluorescence microscopy equipped with a FITC filter set and images were captured. The occurrence of GFP-ATG8-containing puncta (putative autophagosomes) within these cells was assessed in at least 100 cells from 3 independent experiments.

Use of H_2DCF -DA, Alamar Blue, MitoTracker Red, and MitoTracker Green—For assessing the levels of ROS in the promastigotes, cells at 1×10^7 promastigotes ml^{-1} were harvested by centrifugation, washed once in serum-free HOMEEM, and 2×10^6 cells in 200 μl were incubated with 0.1 mM H_2DCF -DA (Molecular Probes) for 2 h at 26 °C and fluorescence was measured using a microtiter plate reader (excitation 380–420 nm, barrier filter 520 nm). The Alamar Blue assay was used to evaluate metabolic activity and cell viability, 4×10^6 promastigotes of WT and $\Delta atg4$ lines at log phase of growth were subjected to starvation conditions (PBS) for up to 8 h at 26 °C or normal growth conditions. Alamar Blue (resazurine salt, Sigma) was added to a final concentration of 0.0125% for the final hour of incubation and its reduction was measured using the fluorescence microtiter plate reader (excitation 550 nm, barrier filter 590 nm). For assessing mitochondrial morphology and mitochondrial membrane potential of the respective lines, 10^7 promastigotes were incubated with either 0.2 μM MitoTracker GreenTM (Invitrogen) or 0.1 μM MitoTracker Red CMXRos (Invitrogen), respectively, for 30 min at 26 °C. Promastigotes were then washed in PBS and either assessed for fluorescence (of MitoTracker Red at excitation 579 nm, barrier filter 599 nm; of MitoTracker Green at excitation 490 nm, barrier filter 516 nm) using a microtiter plate reader or mounted on glass slides for live imaging analysis by fluorescence microscopy using a Nikon TE2000S inverted microscope equipped with FITC and mCherry filter sets. Images were processed using IPLabs 3.7 image processing software (BD Biosciences Bioimaging). For assessing the relationship between autophagy, ROS and mitochondrial morphology and function, NAC at 2.5 or 5 mM was included in some experiments.

Detection of Oxidized Proteins—Carbonylated proteins were detected with an Oxyblot protein oxidation detection kit (Chemicon International Ltd.) according to the manufacturer's protocol. Briefly, promastigote lysates containing 50 μg of protein were incubated with 2,4-dinitrophenylhydrazine for 20 min at room temperature, resolved by SDS-PAGE, and detected by Western blot analysis with anti-2,4-dinitrophenol rabbit antibodies (diluted at 1:200).

Scanning Electron Microscopy (SE) Analyses—Parasites isolated from mice were fixed with 2.5% glutaraldehyde in 0.1 M phosphate buffer, pH 7.4, for 40 min and dehydrated. Samples were dried prior to mounting on an aluminum stub using either silver paint or double stick carbon tape. Samples were then introduced into the chamber of the sputter coater and coated with a very thin film of gold/palladium before SE examination. Body lengths of WT and transgenic lines were measured using the ESI Vision 3.2 Image analysis software (Olympus Soft Imaging Solutions). A minimum of 200 cells was examined for each sample.

Data Processing—Experimental data from macrophage infections, mice infectivity, Alamar Blue, MitoTracker Red, MitoTracker Green, and H_2DCF -DA assays were pooled for comparison using unpaired *t* tests. A *p* value of <0.05 was used as the level of significance. Counts of puncta from ATG-tagged proteins and the measurements of body lengths of promastigotes were performed on at least 100 cells and analyzed statistically as described above.

RESULTS

Targeted Deletion of the *L. major* ATG4 Genes—We have described previously the generation of the *ATG4.2* null mutant ($\Delta atg4.2$) and an add-back line ($\Delta atg4.2::ATG4.2$) and how analysis of the mutants demonstrated that the cysteine peptidase *ATG4.2* is important for differentiation between developmental forms of *Leishmania* (33). To facilitate a comparison between the contributions of the two *ATG4*s in the biology of the parasite, we generated a *ATG4.1* gene deficient cell line (Fig. 1A). The two independent clones obtained (designated $\Delta atg4.1a$ and $\Delta atg4.1b$) were confirmed by Southern blot analysis to lack the *ATG4.1* gene and have the correct integration of the replacement cassette (Fig. 1B). Hybridization of XhoI-digested genomic DNA with a 5'-flank probe revealed a 7.6-kb product corresponding to the *ATG4.1* locus of the WT parasites, which is absent in the $\Delta atg4.1$ lines. In these, a DNA fragment of ~5.0 kb is visible, consistent with correct integration of the *HYG* and *BSD* drug resistance markers in the *ATG4.1* locus. As a control, *ATG4.1* modified with a C-terminal His₆ tag was re-expressed in the $\Delta atg4.1$ lines (to give $\Delta atg4.1::ATG4.1$). Successful expression of *ATG4.1* was confirmed by detection of a 43-kDa protein, which was absent from lysates of WT and $\Delta atg4.1$ promastigotes, by Western blot analysis with anti-His antibody (Fig. 1C). Subsequent phenotypic analyses were performed on both $\Delta atg4.1$ clones, but as they behaved similarly data are presented for $\Delta atg4.1a$ only (hereafter referred to as $\Delta atg4.1$). Although the $\Delta atg4.1$ and $\Delta atg4.2$ null mutants were readily obtained by two successive rounds of targeting their endogenous loci, multiple attempts to generate a line that lacked both *ATG4.1* and *ATG4.2* genes were unsuccessful. Single allele deletions of the second *ATG4* gene in both $\Delta atg4.1$ and $\Delta atg4.2$ lines were successful (to give $\Delta atg4.2/atg4.1^{-/+}$ and $\Delta atg4.1/atg4.2^{-/+}$, respectively), but the second round of allele replacement using these lines resulted in trisomic parasite lines in which both selectable markers had integrated into the appropriate endogenous locus but there appeared an additional copy of the endogenous gene. As both of these loci had been successfully targeted in generating the $\Delta atg4.1$ and $\Delta atg4.2$ mutants, the results of these genetic manipulation attempts suggest strongly that the parasite requires at least one copy of either *ATG4.1* or *ATG4.2*.

***L. major* Mutants Lacking *ATG4*s Form GFP-ATG8-containing Puncta**— $\Delta atg4.1$ and $\Delta atg4.2$ both grew indistinguishably from WT as promastigotes *in vitro*. GFP-ATG8 has been validated as a molecular marker for autophagy in *Leishmania* and has been used to show that autophagy increases during differentiation between life cycle stages with the abundance of autophagosomes in promastigotes being greatest during metacyclogenesis (33, 41). Thus we used this marker protein with the

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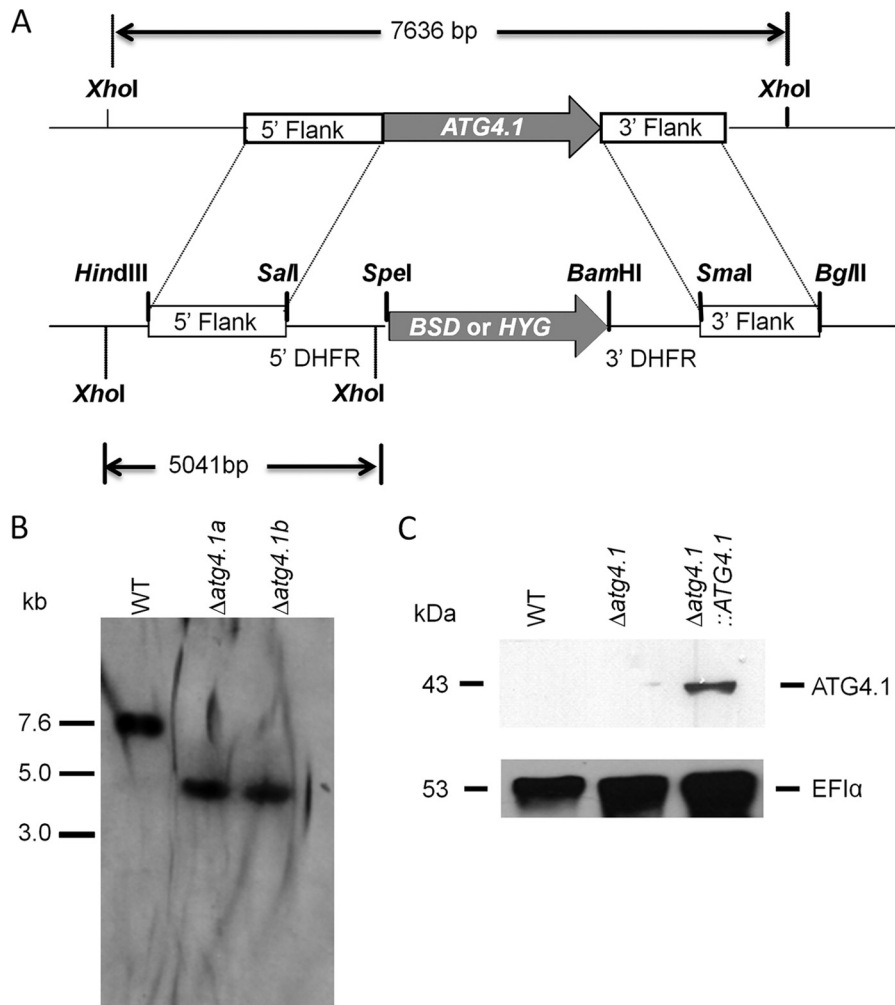


FIGURE 1. Generation of the *ATG4.1* null mutant. *A*, schematic representation of the *ATG4.1* locus and the plasmid constructs used for gene replacement. The *ATG4.1* and antibiotic resistance genes are shown as arrows, flanking DNA sequences are shown as boxes. Restriction enzymes used for the different constructs and the expected sizes of the *XhoI* fragments are shown. 5'-*DHFR* and 3'-*DHFR*, dihydrofolate reductase flanking regions; *BSD*, blasticidin resistance gene; *HYG*, hygromycin resistance gene. *B*, Southern blot analysis of genomic DNA digested with *XhoI*, separated on a 1% agarose gel, blotted onto a nylon membrane, and hybridized with a ^{32}P -labeled DNA probe corresponding to the 5'-flanking region of *ATG4.2*. *C*, Western blot analysis using the α -His antibodies of extracts from early stationary phase promastigotes. *EFl* α served as a loading control.

mutant lines to investigate the involvement of the two ATG4s in autophagy. The lines were transfected with pN-GFP-ATG8 and the occurrence of GFP-labeled puncta was monitored (Fig. 2A). The $\Delta atg4.1$ lines (designated $\Delta atg4.1$ [pN-ATG8] and $\Delta atg4.1::ATG4.1$ [pN-ATG8]) at the mid-logarithmic phase of growth produced one to two GFP-labeled autophagosomes per cell, which was similar to WT cells. In contrast $\Delta atg4.2$ [pN-ATG8] had multiple puncta. With $\Delta atg4.1$, the proportion of cells bearing the GFP-labeled structures peaked at day 7 (when metacyclogenesis was maximal), similarly to the situation with WT promastigotes although the number of cells with puncta was lower (Fig. 2B). The proportion of cells bearing GFP-labeled structures returned to WT levels in $\Delta atg4.1::ATG4.1$ [pN-ATG8]. In both cases, the proportion of cells with GFP-labeled puncta subsequently declined to low levels as stationary phase was reached and metacyclogenesis ended. In contrast, $\Delta atg4.2$ [pN-ATG8] had a higher proportion of cells with autophagosomes at early logarithmic phase of growth, a proportion that increased as the logarithmic phase progressed. Most notably, the cells retained puncta at stationary phase of

growth. $\Delta atg4.2$ promastigotes also were characterized by the presence of multiple puncta (Fig. 2A), which were restored to WT numbers in $\Delta atg4.2::ATG4.2$ [pN-ATG8]. Western blot analysis monitored the proportion of cytosolic and lipid-associated forms of ATG8 (GFP-ATG8* and GFP-ATG8-PE, respectively). All the cell lines at stationary phase of growth contained GFP-ATG8-PE, but the $\Delta atg4.2$ [pN-ATG8] line had a significantly higher proportion of GFP-ATG8-PE relative to GFP-ATG8* (Fig. 2C), which is consistent with the accumulation of autophagosomes as observed by fluorescence microscopy.

We have shown previously that autophagy in *L. major* is induced by starvation (33), so we quantified the number of autophagosomes in the mutants after starvation (Fig. 2D). The proportion of cells with GFP-ATG8 puncta increased dramatically in WT, $\Delta atg4.1$, $\Delta atg4.1::ATG4.1$, and $\Delta atg4.2::ATG4.2$ promastigotes. In contrast, the proportion of $\Delta atg4.2$ promastigotes with GFP-ATG8 puncta was initially relatively high and this only increased a little under starvation conditions.

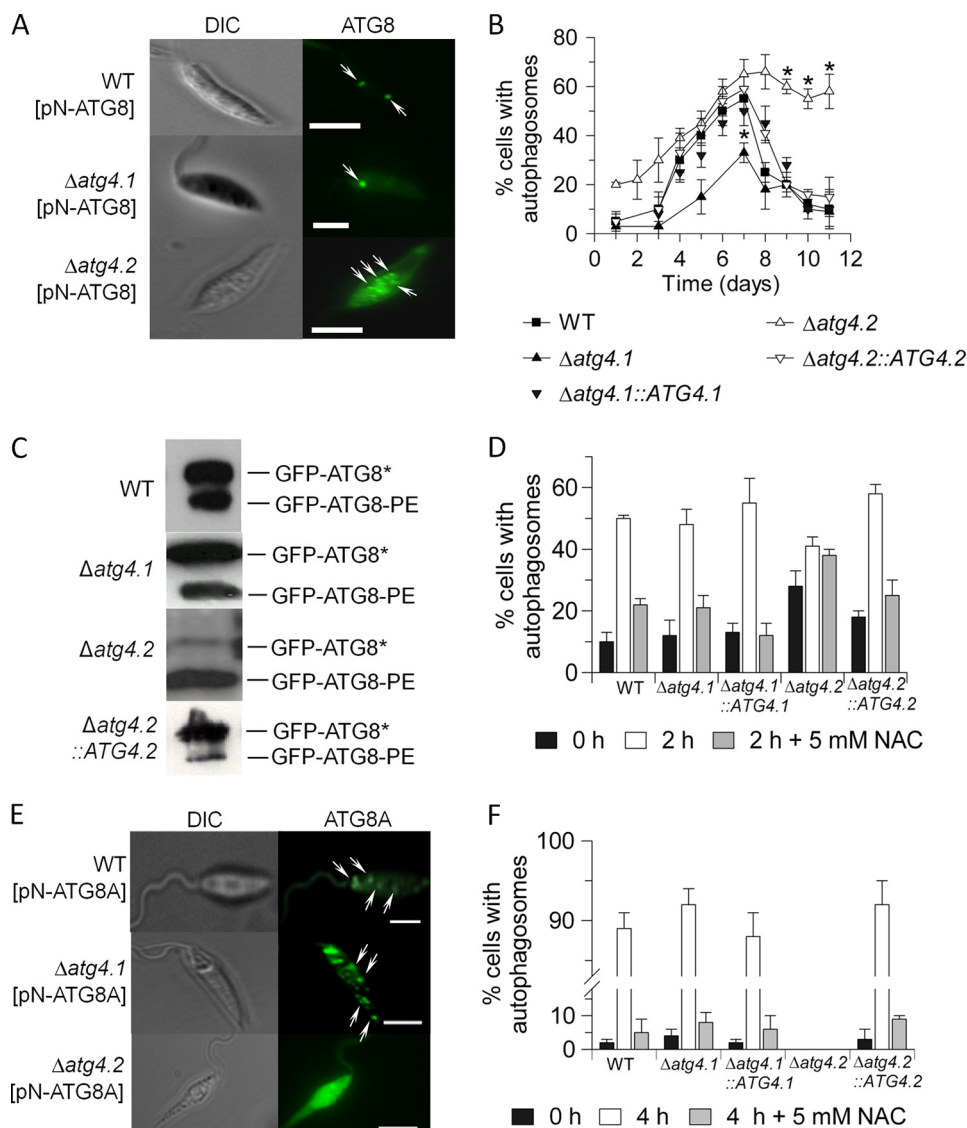


FIGURE 2. Autophagosome formation in promastigotes of the $\Delta atg4$ mutants. *A*, occurrence of GFP-ATG8 puncta in *L. major* promastigotes in normal medium. The arrows indicate autophagosomes. Scale bar, 10 μ m. *B*, the proportion of GFP-ATG8-expressing promastigotes with autophagosomes was assessed during growth *in vitro*. Data are mean \pm S.D. from three independent experiments. Asterisks denote values statistically different from WT cells, $p < 0.05$. *C*, Western blot analysis of GFP-ATG8-PE. Promastigote cell extracts expressing GFP-ATG8 were separated by SDS-PAGE in the presence of 6 M urea and analyzed by Western blotting with an anti-GFP antibody. The faster migrating, lipidated bands are labeled GFP-ATG8-PE, whereas the un-lipidated bands migrating more slowly are labeled GFP-ATG8*. *D*, the occurrence of GFP-ATG8-labeled autophagosomes resulting from starvation of early log stage promastigotes in PBS for 2 h with or without added NAC. Data are mean \pm S.D. from three independent experiments. *E*, occurrence of GFP-ATG8A puncta in starved *L. major* promastigotes. The arrows indicate autophagosomes. Scale bar, 10 μ m. *F*, the occurrence of GFP-ATG8A-labeled autophagosomes resulting from starvation of early log stage promastigotes in PBS with or without added NAC. Note the discontinuity in the ordinate axis.

The Requirement for the Different ATG4s in Autophagy Involving ATG8A—Previously, we showed that the preferred substrate for ATG4.2 was ATG8A (32). To investigate further the relative roles of ATG4.1 and ATG4.2 in the biogenesis of ATG8A-containing puncta, $\Delta atg4.1$ [pN-ATG8A] and $\Delta atg4.2$ [pN-ATG8A] lines were generated, incubated in starvation medium, and monitored for the formation of ATG8A-containing puncta (Fig. 2E). $\Delta atg4.1$ [pN-ATG8A] produced multiple GFP-ATG8A-labeled structures per cell, similar to WT promastigotes. In contrast $\Delta atg4.2$ [pN-ATG8A] produced no such puncta, as described previously (32). The proportion of promastigotes with GFP-ATG8A puncta under starvation conditions was $\sim 90\%$ in WT, $\Delta atg4.1$, $\Delta atg4.1::ATG4.1$, and $\Delta atg4.2::ATG4.2$ (Fig. 2F), a great increase from 2 to 5% in

non-starved cells. No puncta were detected in $\Delta atg4.2$ promastigotes, strongly supporting the hypothesis that ATG8A-dependent puncta formation is totally dependent upon ATG4.2.

Metacyclogenesis and Infectivity of the $\Delta atg4$ Mutants—We reported previously that $\Delta atg4.2$ promastigotes are defective in metacyclogenesis (33), thus we investigated whether $\Delta atg4.1$ has a similar defect. $\Delta atg4.1$ promastigotes differ from $\Delta atg4.2$ promastigotes in that they apparently differentiated fully to metacyclic promastigotes, as judged by the lectin peanut agglutinin assay (Fig. 3A) and the presence of the metacyclic-specific markers, SHERP and HASPB (Fig. 3B); the process seemed delayed and there was a significant difference in the number of metacyclic promastigotes at day 7 between the WT parasites

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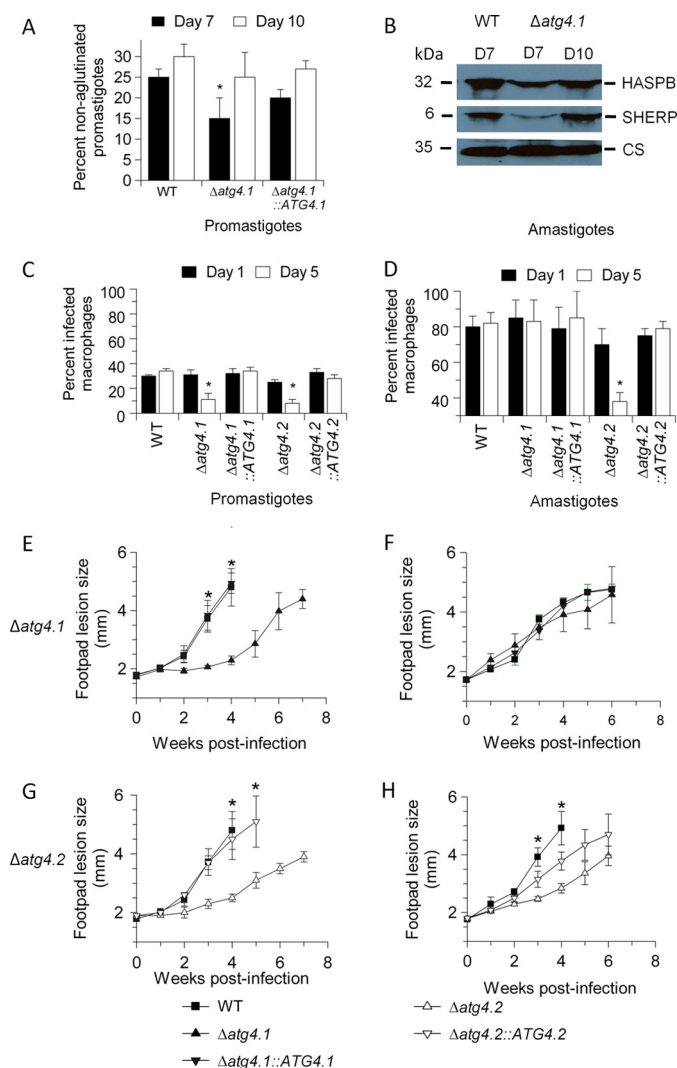


FIGURE 3. Metacylogenesis and infectivity of *L. major* ATG4 null mutants. A, the percentage of metacyclic promastigotes in stationary phase populations of promastigotes was determined by the peanut lectin agglutination method. Values shown are the mean \pm S.D. from three independent experiments. *, data for $\Delta atg4.1$ differed significantly from those for WT promastigotes at day 7 ($p < 0.01$). B, extracts of 10^7 stationary phase promastigotes at day 7 (D7) or day 10 (D10) were assessed for metacyclic-specific proteins by Western blot analysis with anti-SHERP and anti-HASPB antibody. Cysteine synthase (CS) served as a loading control. C and D, mouse peritoneal macrophages were infected *in vitro* with stationary phase promastigotes or amastigotes isolated from lesions in BALB/c mice. Cells were assessed at 1 and 5 days post-infection for the percentage of infected macrophages. Data are mean \pm S.D. and are representative of three independent experiments. *, infection level of ATG4 null mutant lines differed significantly from those of WT parasites ($p < 0.05$). E–H, BALB/c mice were infected with 2×10^7 promastigotes or amastigotes and footpad lesion size was monitored weekly. Data are mean \pm S.D. of five mice. *, infection level of ATG4 null mutant lines differed significantly from those of WT or the re-expressing line ($p < 0.05$).

and $\Delta atg4.1$ ($p < 0.05$), but by day 10 there was no difference (Fig. 3A).

We compared the two mutants for infectivity to macrophages *in vitro* and mice. Stationary phase promastigotes of both $\Delta atg4.1$ and $\Delta atg4.2$ infected macrophages *in vitro* in similar numbers to WT parasites but each of them survived poorly over 5 days with many macrophages clearing the infection (Fig. 3C). In contrast, amastigotes of $\Delta atg4.1$, purified from mice, were as infective as WT amastigotes and proliferated equally well (Fig. 3D). Amastigotes of $\Delta atg4.2$ also infected macro-

phages well but survived poorly over 5 days (Fig. 3D, right panel).

Analysis of the infectivity of $\Delta atg4.1$ and $\Delta atg4.2$ lines to BALB/c mice using promastigotes and amastigotes yielded results that correlated well with those from the *in vitro* analysis using macrophages (Fig. 3, E–H). Lesions resulting from inoculation of stationary phase $\Delta atg4.1$ and $\Delta atg4.2$ promastigotes were slower growing than those of WT parasites, this was reversed by re-expression of the genes (Figs. 3, E and G). In contrast, inoculation of $\Delta atg4.1$ amastigotes generated lesions similar to those of WT amastigotes (Fig. 3F), whereas inoculation of $\Delta atg4.2$ amastigotes resulted in slower growing lesions (Fig. 3H). Lesion development with amastigotes of $\Delta atg4.2::ATG4.2$ was at an intermediate level between WT and $\Delta atg4.2$ (Fig. 3H). These findings suggest that both ATG4s play some role in facilitating infection of mice by promastigotes, and that ATG4.2 is also important in the survival and multiplication in mice of inoculated amastigotes, whereas ATG4.1 is not.

Morphology of $\Delta atg4.2$ in Mice—The finding that inoculation of $\Delta atg4.2$ amastigotes into BALB/c resulted in slower growing lesions than WT parasites prompted us to investigate whether the lesions were equivalent in terms of parasite numbers per volume. The results showed that the number of amastigotes contained within lesions of comparable width was significantly lower for $\Delta atg4.2$ than with WT and $\Delta atg4.1$ ($p < 0.05$; Fig. 4A). Scanning electron microscopy (SE) analysis of $\Delta atg4.2$ purified from lesions showed that they had a longer body length on average than WT amastigotes ($p < 0.05$; Fig. 4, B and C), a difference largely reversed by re-expression of ATG4.2 in $\Delta atg4.2::ATG4.2$. $\Delta atg4.1$ amastigotes were not significantly different in size from WT amastigotes ($p > 0.05$; data not shown).

Transformation of the parasites extracted from lesions in mice back to promastigotes was assessed by inoculating them into 5 ml of medium at the cell density of $2 \times 10^5 \text{ ml}^{-1}$. Under these conditions, $\Delta atg4.2$ parasites differentiated into promastigotes more quickly than the WT and $\Delta atg4.1$ parasites (Fig. 4D).

Use of the $\Delta atg4$ Mutants to Study the Relationship between ROS Levels and Autophagy in *Leishmania*—The relationship between ROS levels and induction of autophagy known for mammalian cells prompted us to study whether there is a correlation between ROS levels and the abundance of autophagosomes in *Leishmania*. The approaches we adopted were measuring ROS levels in the mutants, correlating these with autophagosome generation, and measuring the sensitivity of the autophagy-compromised mutants to oxidative stress. We also modulated the ROS levels in promastigotes through the addition of the antioxidant NAC (known to scavenge H_2O_2 , hydroxyl radical, and superoxide within cells and to help maintain a reduced intracellular environment (42)), and monitored how this affected autophagy.

NAC added to the starvation buffer inhibited the biogenesis of GFP-ATG8-containing puncta in all cell lines except $\Delta atg4.2$ promastigotes (Fig. 2D). The proportion of $\Delta atg4.2$ promastigotes with GFP-ATG8-containing autophagosomes was not statistically different whether or not NAC was added (Fig. 2D). These data show an apparent correlation between increased

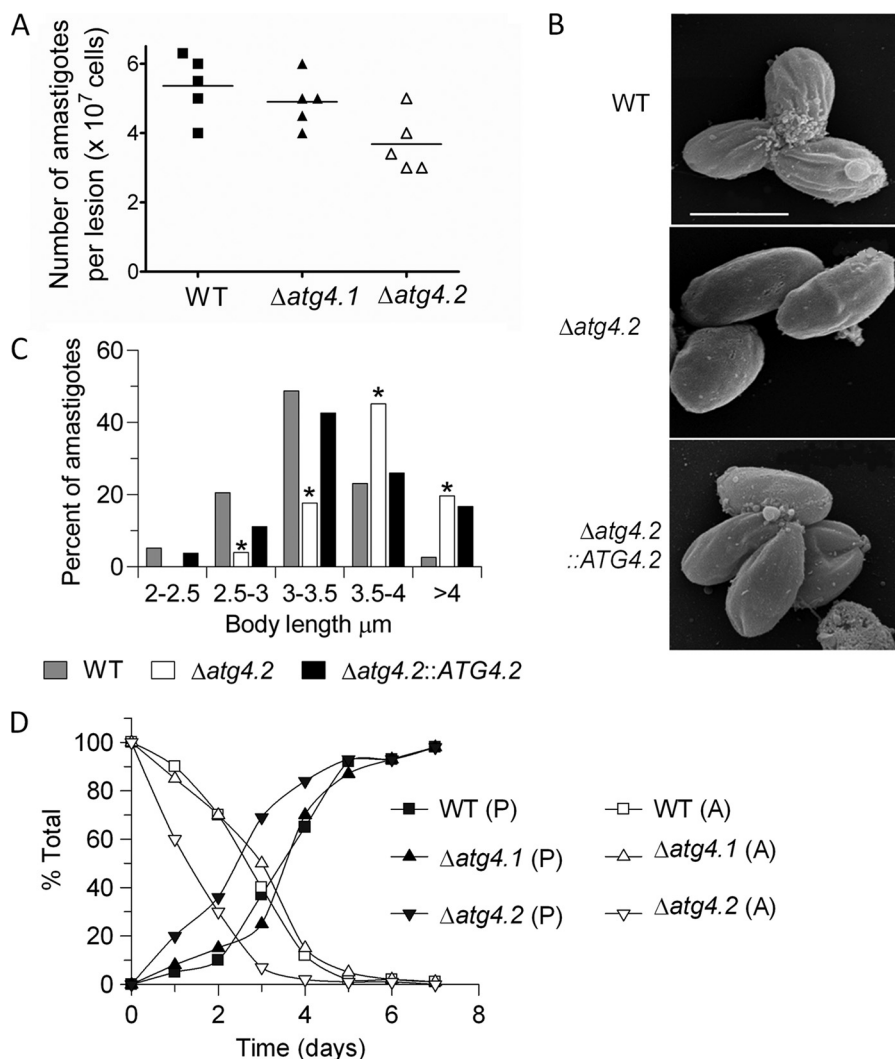


FIGURE 4. **Morphology and transformation of amastigotes of $\Delta atg4.2$.** A, parasite load in footpad lesions of infected BALB/c mice. The lesions from five mice were dissected when they reached 4 mm in thickness, the tissue was disrupted in PBS, and the number of parasites was determined microscopically. The lines indicate the means, which for WT and $\Delta atg4.1$ are significantly different from $\Delta atg4.2$, $p < 0.05$. B, scanning electron micrographs. Bar represents 4 μm . C, histogram of body lengths of amastigotes isolated from lesions in mice. The abscissa shows body length ranges in μm ; the ordinate indicates percent of cells counted. *, data for $\Delta atg4.2$ differed significantly from those for WT parasites in all class intervals ($p < 0.05$). D, transformation of amastigotes isolated from BALB/c mice. The data show the disappearance of amastigotes (A) and the appearance of promastigotes (P) over 7 days.

ROS levels and number of GFP-ATG8 puncta in starved promastigotes (except for $\Delta atg4.2$) and are consistent with autophagy being induced by ROS as reported for other cells (24, 25, 43). However, the accumulation of GFP-ATG8 containing puncta in $\Delta atg4.2$ even when ROS levels had been diminished using NAC confirm that this mutant has dysfunctional autophagic machinery.

The addition of NAC also abrogated the formation of the GFP-ATG8A structures (Fig. 2F). These data suggest that ATG8A-dependent puncta formation is similar to autophagy involving ATG8 in being induced by ROS.

Increased ROS Levels and Oxidatively Damaged Proteins in $\Delta atg4$ —To elucidate the parts played by each *L. major* ATG4 in any autophagic response to counter ROS, we measured the levels of ROS in normal and starved promastigotes of the various lines using 2,7-dichlorofluorescein ($H_2DCF-DA$) (24, 25, 44, 45). This revealed that ROS levels in $\Delta atg4.2$ were 2-fold higher ($p < 0.01$) than in WT promastigotes at early log phase (day 2),

whereas those in $\Delta atg4.1$ were similar to WT (Fig. 5A, compare with Fig. 2B). ROS levels were increased 40-fold in WT promastigotes during a 2-h starvation, a similar scale of increase also occurred in the two null mutant lines (Fig. 5A). Addition of NAC at 2.5 mM to the starvation medium suppressed ROS levels at 2 h very greatly in promastigotes of all lines except $\Delta atg4.2$ (Fig. 5A). More NAC (5 mM) was required to produce an equivalent reduction of ROS in $\Delta atg4.2$ (Fig. 5A).

ROS levels during *in vitro* growth of promastigotes were also assessed to determine whether there was any apparent relationship between these and the occurrence of autophagy. The levels in WT promastigotes increased as early log phase progressed to mid log phase and early stationary phase but then declined at stationary phase (Fig. 5B, time points equivalent to days 2, 4, 6, and 9 in Fig. 2). Thus the period at which ROS levels were highest coincided with the time when autophagy was also elevated. The same profile occurred with $\Delta atg4.1$ and $\Delta atg4.1::ATG4.1$. However, the situation was different in the

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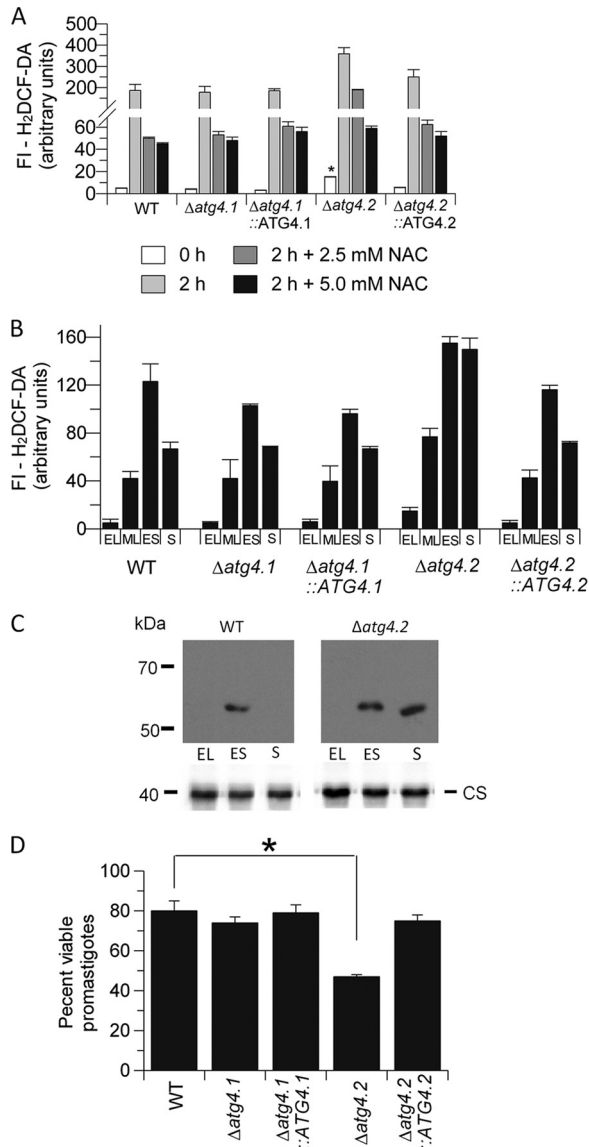


FIGURE 5. Increased ROS levels and oxidatively damaged proteins in $\Delta atg4.2$ promastigotes. *A*, ROS levels, measured using 50 mM H₂DCF-DA, in early log promastigotes incubated for 2 h at 27 °C in PBS with or without NAC. Note the discontinuity in the ordinate axis. Data are mean \pm S.D. from independent experiments. *, ROS levels in $\Delta atg4.2$ were significantly higher than in WT promastigotes, $p < 0.001$. *B*, ROS levels in promastigotes at early log (EL), mid log (ML), early stationary (ES), and stationary (S) phases of growth (days 2, 4, 6, and 9, respectively). *C*, occurrence of carbonylated protein in extracts of promastigotes at the growth phases detailed in *B*. Cysteine synthase (CS) served as a loading control. *D*, log phase promastigotes were starved for 2 h in PBS and then assessed for the proportion of viable cells compared with WT promastigotes in complete HOMEM medium using a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide colorimetric assay. *, denotes statistically significant difference, $p < 0.05$.

$\Delta atg4.2$ line, DCF-DA fluorescence intensity was 2-fold higher than its WT counterpart at early log phase, increased as log phase progressed, and was maintained at this high level during stationary phase. This phenotype was restored to WT levels in the $\Delta atg4.2::ATG4.2$ re-expressor.

One consequence of elevated ROS in a cell is the unwanted oxidation of proteins, which can be removed via autophagy. Protein carbonyl content has been shown to be a biomarker for protein oxidation (46), thus we investigated the presence of carbonylated proteins in parasite extracts by derivatization

with 2,4-dinitrophenylhydrazine. The stable dinitrophenylhydrazone product was assessed by Western blot analysis. Just a single protein of ~ 60 kDa was detected in extracts of WT promastigotes, as previously reported (47, 48), and only in those of promastigotes at early stationary phase of growth when ROS levels are highest (Fig. 5C). However, the same analysis with $\Delta atg4.2$ yielded the equivalent protein also in extracts of stationary phase promastigotes (Fig. 5C). Thus for this mutant there is an apparent correlation between elevated ROS levels, protein carbonylation, and restricted autophagic flux.

$\Delta atg4.2$ Promastigotes Are Less Able to Withstand Starvation and Oxidative Stresses—Autophagy generally has a role in protecting against starvation and other stresses, thus we tested the hypothesis that deletion of *ATG4* would hinder autophagic flux and make promastigotes of *L. major* more susceptible to stresses. $\Delta atg4.2$ promastigotes were indeed less able to withstand starvation in PBS, with viability, as assessed using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay, after 2 h starvation being 42% less than that of WT, $\Delta atg4.1$, $\Delta atg4.1::ATG4.1$, and $\Delta atg4.2::ATG4.2$ promastigotes under the same conditions (Fig. 5D). Viability of the WT itself was reduced to $\sim 80\%$ compared with that when in growth medium.

The sensitivity of the various lines to exogenous hydrogen peroxide was also assessed. The results show that $\Delta atg4.2$ promastigotes are more susceptible than the other lines, with an IC₅₀ of 0.21 ± 0.02 mM compared with IC₅₀ values of around 0.5 mM for the other lines. These findings suggest that deletion of *ATG4.1* does not interfere with the ability of the promastigote to withstand these stresses, whereas removal of *ATG4.2* does have a detrimental effect.

$\Delta atg4.2$ Promastigotes Have a Dysfunctional Mitochondrion—There is increasing evidence that general autophagy is required for the maintenance of mitochondrial homeostasis, and we have recently shown this to be so for *Leishmania* by generating mutants lacking ATG5 (31). Thus we investigated whether removal of the individual ATG4s from *L. major* had an impact upon the mitochondrion of the parasite. MitoTracker Green staining, which is used to assess total mitochondrial mass, showed that $\Delta atg4.1$ promastigotes were similar to WT promastigotes and also that more than three-quarters of $\Delta atg4.2$ promastigotes at stationary phase of growth contained a reticulate mitochondrion characteristic of WT promastigotes (Fig. 6A). The remainder ($\sim 20\%$) of $\Delta atg4.2$ promastigotes had collapsed or fragmented mitochondrial morphology. However, there was a similar overall intensity of the fluorescence in all the cell lines, consistent with there being little difference in mitochondrial mass and this remained in the presence of 5 mM NAC (data not shown). A much lower fluorescence intensity was observed with MitoTracker Red (which assesses mitochondrial membrane potential) in $\Delta atg4.2$ compared with WT promastigotes (Fig. 6B, closed bars; $p < 0.05$), suggesting that mitochondrial function was impaired in this null mutant. No such difference was noted for the other mutants (Fig. 6B, closed bars). Addition of NAC to normal medium greatly increased the mitochondrial membrane potential of $\Delta atg4.2$ promastigotes by 4 h (Fig. 6B, open bars) and also suppressed ROS levels (Fig. 6C, open bars) although the mitochondrial network was

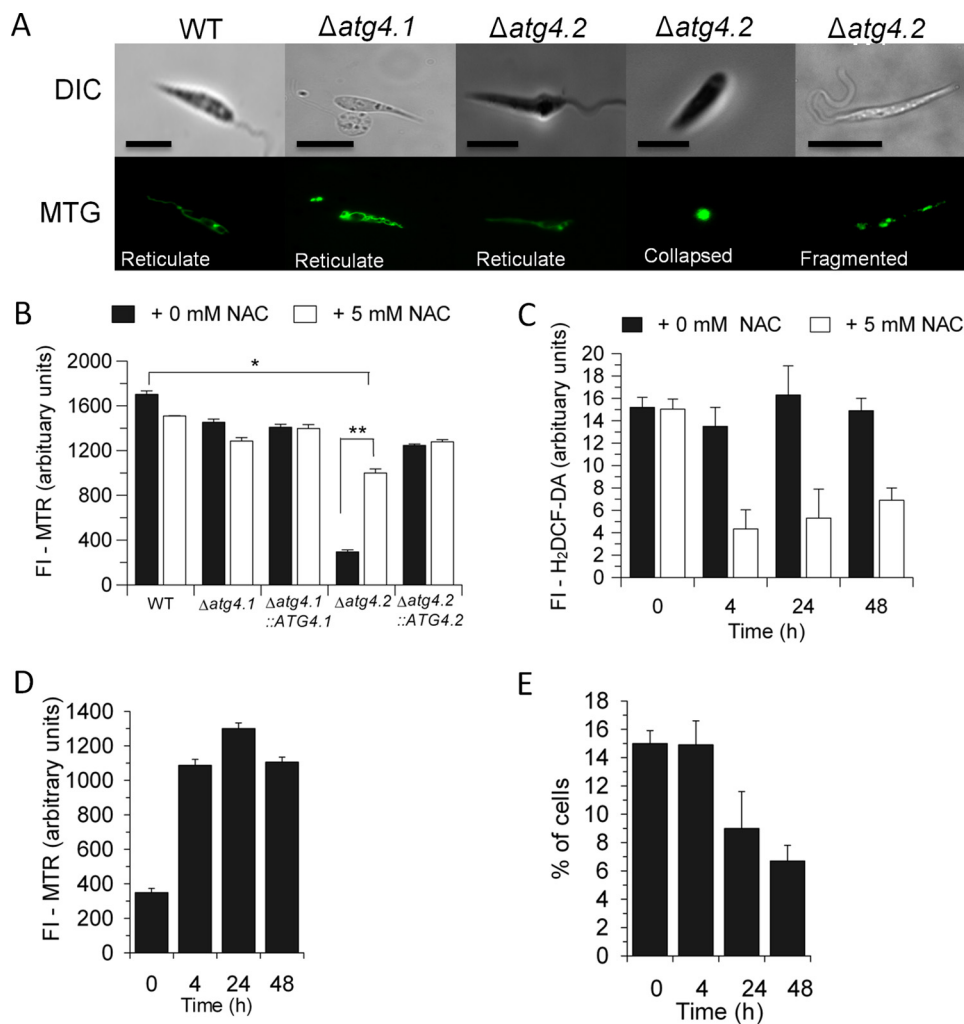


FIGURE 6. Evidence for a dysfunctional mitochondrion in $\Delta atg4.2$ but not $\Delta atg4.1$. A, the types of mitochondrial morphology observed by fluorescence microscopy in promastigotes stained with MitoTracker Green. Scale bar, 10 μm . B, the fluorescent intensity from MitoTracker Red, at 0.1 μM , of 2×10^6 promastigotes in normal medium after 4 h incubation at 26 $^{\circ}\text{C}$ with and without NAC. Data are mean \pm S.D. from three independent experiments. * and ** denote statistically significant differences between, respectively, WT and $\Delta atg4.2$ and $\Delta atg4.2$ with and without NAC. C–E, effect of addition of NAC to $\Delta atg4.2$ promastigotes at $2 \times 10^6 \text{ ml}^{-1}$ in normal medium over 48 h. C, ROS levels measured using 50 μM H₂DCF-DA. D, fluorescent intensity from MTR. E, the proportion of $\Delta atg4.2$ promastigotes with an abnormal mitochondrial morphology.

still disrupted in $15 \pm 1\%$ of promastigotes at this time point (Fig. 6E). Prolonged incubation in normal medium containing NAC for up to 48 h maintained ROS at relatively low levels (Fig. 6C, open bars) and the mitochondrial membrane potential close to but significantly lower than that of WT promastigotes (Fig. 6, compare D with B) and there was also a significant decrease in the proportion of cells with a defective mitochondrion (to $6 \pm 1\%$, Fig. 6E). These results suggest that the lack of ATG4.2 and consequent impaired autophagic flux is responsible for the increased levels of ROS and that these lead to impaired mitochondrial homeostasis and function.

DISCUSSION

This study aimed to elucidate the roles of the two ATG4 enzymes in *L. major* by generating gene deletion mutants. Both individual null mutants were generated successfully using promastigotes and their phenotypes were subsequently analyzed. The main defects observed for $\Delta atg4.1$ were the fewer promastigotes that contained GFP-ATG8-containing autophagosomes (Fig. 2B) and poorer ability of promastigotes to success-

fully infect macrophages *in vitro* (Fig. 3, C and D) and in mice (Fig. 3, E–H), even though they appeared to undergo metacyclogenesis *in vitro* similarly to, although slightly slower than, WT parasites (Fig. 3, A and B). The capacity of this mutant to produce autophagosomes, albeit less abundantly than the WT parasites, implies some redundancy in function between the two ATG4s. We have previously shown that although ATG4.2 has higher activity toward the ATG8A group of paralogues, it can hydrolyze ATG8 *in vitro* after prolonged incubation (32) and this is supported by the observation in this study of GFP-ATG8-containing puncta in mutants lacking ATG4.1. However, puncta containing ATG8B and ATG8C, also preferred substrates for ATG4.1 (32), were not formed in $\Delta atg4.1$ (data not shown). Thus the viability of $\Delta atg4.1$ shows that these proteins and the puncta that they form do not have an essential role in the parasite. The reduced ability of promastigotes of $\Delta atg4.1$ to infect macrophages and mice may reflect a reduction in flux through the autophagic pathway and so, perhaps, a reduced capacity for differentiation between life cycle forms. Thus these

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data are consistent with the contributions of ATG4.1 to autophagy being significant but not essential for *Leishmania* development.

The deletion of *ATG4.2* had more profound effects upon the parasite. The lack of *ATG4.2* markedly reduces the ability of the parasite, whether as promastigotes and amastigotes, to infect and survive in both macrophages *in vitro* (Fig. 3, *C* and *D*) and also mice (Fig. 3, *E–H*). As $\Delta atg4.2$ had a significantly higher amount of lipidated ATG8 (Fig. 2*C*) and a higher number of GFP-ATG8-containing puncta at stationary phase of growth (Fig. 2*B*), we conclude that the lower virulence of this mutant is due primarily to a block in the delivery of autophagosomes to the lysosomal network, probably due to poor cleavage of ATG8 from the mature autophagosomes, and thus a reduction in flux through the ATG8-dependent autophagic pathway. To further characterize the identity of the higher number of GFP-ATG8-containing puncta in $\Delta atg4.2$ at stationary phase of growth, we transfected the line with mC-ATG5, which also labels autophagosomes (31). These analyses showed that most of the puncta were labeled with GFP-ATG8 only, very rarely was labeling observed with mC-ATG5 (data not shown). As ATG5 is released from nascent autophagosomes before they mature, this is entirely consistent with our idea that the additional puncta in $\Delta atg4.2$ are mature autophagosomes. Unfortunately there are no other markers for autophagosomes currently available for use with *Leishmania*. However, electron microscopic analyses also revealed that structures typical of autophagosomes (as exemplified in Ref. 41) were more abundant in $\Delta atg4.2$ than in $\Delta atg4.1$ or WT parasites (data not shown).

The significance of the deconjugation of ATG8-PE by ATG4 has been described previously for yeast (49). In this organism, with just a single *ATG4* gene, the expression of a mutated Atg8 (truncated at the glycine required for conjugation to PE, which avoids the need for the initial processing by ATG4) in an *atg4*-deletion mutant produced a low autophagic efficiency but high levels of ATG8-PE and increased numbers of autophagosomes in the cytosol. These results with yeast and also now *Leishmania* $\Delta atg4.2$ confirm that ATG4s have delipidation activity that is required at the late stage of the autophagic cycle. This conclusion is further supported by an inhibitory effect on autophagosome occurrence of overexpressing ATG4.2 in *Leishmania* promastigotes (32) and HsAtg4b in human cells (49). We have also shown previously that recombinant ATG4.2 can cleave ATG8-PE and release ATG8 (32), which adds further support for the important role of ATG4.2 at this stage of the autophagic process.

The increase in number of autophagosomes resulting from deletion of *ATG4.2* contrasts with our finding (detailed in Ref. 41) that inhibiting lysosomal peptidases caused an increase in undigested autophagosomal material within the lysosomal network, an effect not observed in this current study, but not an increased number of mature autophagosomes. These differing effects resulting from the two interventions are consistent with the role we postulate for *ATG4.2* in autophagy in *Leishmania*.

A block in autophagy in $\Delta atg4.2$ would also account for other observed phenotypic characteristics of $\Delta atg4.2$, including the reduced ability to maintain viability during oxidant and starvation stresses and a lower ability to induce autophagy when

nutrients were limiting. It is also likely that such reduced flux would hinder cellular differentiation of the parasite and so could explain why the forms of the parasite isolated from mice were abnormally large and differentiated more rapidly to promastigotes, they were not fully differentiated to amastigotes even in a mammal (Fig. 4). Similar findings have been reported for mouse fibroblast cells, which when defective in autophagy were unable to reduce their size in response to starvation (50), and *Arabidopsis*, where mutants lacking the two ATG4s (named *atg4a4b-1*) were unable to form autophagosomes under nutrient-limiting conditions (51).

ATG4.2 of *Leishmania* also has a high activity toward ATG8A, the paralogue of ATG8 that forms puncta primarily under starvation conditions (32). *ATG4.1* does not cleave ATG8A, which explains why ATG8A-labeled puncta were not formed in $\Delta atg4.2$ (Fig. 2, *F* and *G*). This inability could play some part in determining the phenotype of this null line, but as yet the full significance of the ATG8A-containing putative autophagosomes is unknown.

Autophagy plays a part in protecting against ROS-mediated damage of proteins and organelles in eukaryotic cells (52), and its absence results in elevated ROS levels and consequently accumulation of oxidized proteins, a lower mitochondrial membrane potential and dysfunctional mitochondria (52, 53). Moreover, autophagy can be induced by elevated ROS (24–26). Our investigation of the relationship between ROS and autophagy in *Leishmania* has shown that ROS levels do vary as *Leishmania* promastigotes grow *in vitro* with, interestingly, higher levels occurring during the phase when metacyclogenesis occurs and autophagy is elevated (Fig. 5*B*). $\Delta atg4.2$ had higher levels of ROS than WT throughout this growth, with notably higher levels at stationary phase when the numbers of autophagosomes are also markedly elevated. This is consistent again with autophagic flux being reduced in this mutant and a role for autophagy in dealing with ROS stress, hypotheses that can be tested in future studies.

There was also a prominent depolarization of the mitochondrion and some fragmentation in $\Delta atg4.2$ (Fig. 6). We have recently shown that mutants lacking ATG5, which do not produce autophagosomes, have an even more markedly dysfunctional mitochondrion with an increased mitochondrial mass (31), thus the data for $\Delta atg4.2$ are consistent with this. There is increasing evidence for a close inter-relationship between autophagy and mitochondrial homeostasis (54–56) and these data with *Leishmania* $\Delta atg4.2$ support this.

Our experiments using the antioxidant NAC in parasites starved for 2 h showed that this agent resulted in lower ROS levels in all lines (Fig. 5*A*) and also to decreased production of autophagosomes containing ATG8 and ATG8A (Fig. 2) and restoration of mitochondrial integrity (Fig. 6). This is consistent with ROS being an important regulator of autophagy in *Leishmania* and also supports ROS as a mediator of mitochondrion damage, although direct evidence needs to be obtained in future studies to confirm this.

In human cells, ROS modulates the autophagic pathway during amino acid starvation (25). This appears to occur through ROS inhibiting the deconjugation activity of ATG4, causing ATG8-PE to accumulate at the pre-autophagosomal mem-

branes. This fosters autophagosome biogenesis, and so autophagy to remove more rapidly oxidized proteins and damaged organelles (24, 45). This mechanism cannot operate in *Leishmania*, however, as neither *Leishmania* ATG4 is affected directly by ROS, as demonstrated by the lack of inhibition by 5 mM H₂O₂ of the hydrolytic activities of recombinant ATG4.1 and ATG4.2 toward their preferred substrates, His-ATG8-HA and His-ATG8A-HA, respectively (data not shown). Yeast also similarly differs from mammalian cells in this way and this organism has a regulatory mechanism in which ATG4 is blocked from ATG8-PE in nascent autophagosomes by the ATG12-ATG5 complex, ATG18 and ATG21 (8). It remains to be determined whether such a blockage functions in *Leishmania* but we have shown that the parasite does have the ATG5-ATG12 conjugation pathway (31) and a gene encoding ATG18 is present in the *L. major* genome (www.geneDB.org, LmjF29.1575). However, there is not currently strong evidence for a gene encoding ATG21 and so if such inhibition occurs the precise mechanism is likely to differ from that in yeast.

Our results overall show that the ATG4s of *L. major* are different, there is some redundancy in function between them especially in autophagosome biogenesis, but that ATG4.2 would appear to be more important than ATG4.1 for the later stages of the autophagic pathway when ATG8 is cleaved from the autophagosomes. Neither is individually essential, and the finding that cells lacking both *ATG4.1* and *ATG4.2* could not be obtained, even though both loci were demonstrated to be targetable by the gene deletion constructs, suggests that ATG4 performs essential roles in *Leishmania*. The activity of the enzymes toward not only ATG8 but also ATG8A, both of which function in autophagy in some conditions, makes autophagy in *Leishmania* an interesting system to investigate in terms of the roles of the individual ATG4 proteins.

The results presented also support the idea that autophagy is important for differentiation of the parasite, and especially in enabling promastigotes to infect mammals. Importantly, however, ATG4.2 appears to have a constitutive role in the growth of the parasite in mammals. This is an important finding that opens up many new avenues for further investigation on the roles of autophagy in the *Leishmania*-mammal interaction.

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