AP1 is essential for generation of autophagosomes from the trans-Golgi network

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Summary
Despite recent advances in understanding the functions of autophagy in developmental and pathological conditions, the underlying mechanism of where and how autophagosomal structures acquire membrane remains enigmatic. Here, we provide evidence that post-Golgi membrane traffic plays a crucial role in autophagosome formation. Increased secretion of constitutive cargo from the trans-Golgi network (TGN) to the plasma membrane induced the formation of microtubule-associated protein light chain 3 (LC3)-positive structures. At the early phase of autophagy, LC3 associated with and then budded off from a distinct TGN domain without constitutive TGN-to-plasma cargo and TGN-to-endosome proteins. The clathrin adaptor protein AP1 and clathrin localized to starvation- and rapamycin-induced autophagosomes. Dysfunction of the AP1-dependent clathrin coating at the TGN but not at the plasma membrane prevented autophagosome formation. Our results thus suggest an essential role of the TGN in autophagosome biogenesis, providing membrane to autophagosomes through an AP1-dependent pathway.

Key words: AP1, LC3, Autophagosome, Membrane trafficking, Trans-Golgi network

Introduction
Autophagy is a highly conserved process in eukaryotic cells and is a mechanism for the turnover of cytoplasmic materials in a lysosome-dependent pathway. It is used either to provide nutrients during starvation or as a quality control that eliminates obsolete macromolecules and organelles during cell growth. In addition to the identification of more than 30 autophagy-related genes (Atgs) whose products are required for autophagic vacuole formation and development, recent studies have revealed that autophagy is involved in multiple physiological and pathological processes, including immunity, aging, neurodegenerative diseases and tumorigenesis.

Despite the progress achieved in understanding the molecular basis of autophagy and its important role in physiological and pathological situations, two crucial issues remain unclear: the origin of the smooth membrane cisternae, and the mechanism by which autophagosomal structures acquire membrane. To date, extensive evidence suggests that the autophagic membranes are derived from pre-existing cytoplasmic membrane compartments including the endoplasmic reticulum (ER), the Golgi complex and mitochondria (Juhasz and Neufeld, 2006; Reggiori and Tooze, 2009). Because it is the largest intracellular membrane source in eukaryotic cells, the ER appears to be the origin of autophagosomal membrane based on the discovery of ER marker enzymes in pre-autophagosomal structures (Arstila and Trump, 1968; Ericsson, 1969). In addition, it has been found that phosphatidylinositol 3-phosphate [PtdIns(3)P]-enriched membranes dynamically connect to the ER and provide a membrane platform for autophagosomes (Axe et al., 2008). Recently, this physical connection between the ER and autophagosomes has been confirmed by 3D electron tomography (Hayashi-Nishino et al., 2009; Yla-Anttila et al., 2009). Based on these observations, it has been proposed that a portion of the ER is cleared of ribosomes and folds onto itself to form the isolation membrane (IM), which is a forming autophagosome cradled between two ER membranes (Axe et al., 2008; Hayashi-Nishino et al., 2009).

However, many studies have also reported that some Atg proteins essential for autophagosome formation act at sites outside the ER. Beclin1, the mammalian homolog of yeast Atg6, which is important in mediating the localization of other autophagy proteins to pre-autophagosomal structures, functions mainly at the trans-Golgi network (TGN) as part of a class III PI3K complex (Kihara et al., 2001). The sole transmembrane Atg protein, Atg9, is located in the TGN and travels between the TGN and endosomes in mammalian cells (Young et al., 2006). The Golgi-resident small GTPase, Rab33B, interacts with Atg16L and modulates autophagosome formation (Itoh et al., 2008). In yeast, subunits of the conserved oligomeric Golgi complex localize to the phagophore assembly site and are required for the formation of double-membrane cytoplasm-to-vacuole targeting vesicles and autophagosomes (Yen et al., 2010). All these observations highlight the importance of the Golgi complex, including the TGN, in the biogenesis of autophagosomes. In addition to its localization in the TGN, Atg9 in yeast also targets to mitochondria and travels between mitochondria and the pre-autophagosomal structure (PAS) (Reggiori et al., 2005). In mammalian cells, Atg5 and microtubule-associated light chain 3 (LC3) transiently localize to punctae on mitochondria, and the tail-anchor of a mitochondrial outer membrane protein also labels autophagosome membranes and is sufficient to deliver another
outer mitochondrial membrane protein to autophagosomes (Hailey et al., 2010). These data indicate a connection between mitochondria and autophagosomal structures, and confirm that mitochondria contribute membrane to autophagosomes. More recently, it has been reported in mammalian cells that the coat protein clathrin interacts with Atg16L and is required for the formation of Atg16L-positive autophagosome precursors from the plasma membrane by endocytosis, suggesting that the plasma membrane is a membrane reservoir for inducible autophagosome formation (Ravikumar et al. 2010).

LC3 is the first protein shown to specifically label autophagosomal membranes in mammalian cells (Kabeya et al., 2000) and it is involved in both the origin and elongation of the autophagosomal membranes. Association of LC3 with autophagosomal membranes requires relocation of the protein from the nucleus (Darke et al., 2010), several steps of post-translational modification of pro-LC3, including cleavage at its C-terminal G120 site to form a soluble LC3-I, and a subsequent attachment of phosphatidylethanolamine (PE) to form membrane-bound LC3-II (Tanida et al., 2004; Sou et al., 2006). In this study, using LC3-II as an autophagosomal membrane marker, we investigated the regulatory function of intracellular membrane trafficking in autophagosome formation by modulating the protein secretory pathway at different steps in mammalian cells. We showed that a transient increase in constitutive cargo flow from the TGN to the plasma membrane initiates the generation of LC3-positive vesicles. Blockage of post-Golgi transport, by disrupting AP1-dependent clathrin coating in the TGN, inhibited autophagy. We also showed association of LC3 with the TGN, and AP1 and clathrin with autophagosomes, during autophagy. Our observations suggest a crucial contribution of TGN membrane and AP1 and clathrin coats to autophagosome formation.

**Results**

**ER export is essential for autophagosome formation**

To investigate a possible regulatory effect of the secretory pathway on autophagy in mammalian cells, we assessed the effect of disrupting ER–Golgi trafficking by overexpression of different mutants that interfere with the core functions of the small GTPases Sar1 and Arf1 in the formation of COPII and COPI vesicles (Pucadyil and Schmid, 2009). In HEK293 cells expressing green fluorescent protein (GFP)-tagged LC3, one hour of starvation triggered a dramatic increase in the generation of autophagosomes in the cytoplasm, indicated as GFP–LC3-positive spot-like structures (Fig. 1A). Transient expression of human influenza hemagglutinin (HA)-tagged Arf1T31N, a constitutively inactive Arf1 (Dascher and Balch, 1994; Klausner et al., 1992), Sar1T39N, a constitutively inactive Sar1 (Barlowe et al., 1994; Kuge et al., 1994; Shima et al., 1998) or Sar1H79G, a constitutively active Sar1 (Aridor et al., 1995), prevented starvation-induced autophagosome formation (Fig. 1A,B). Because the amount of LC3-II represents membrane-bound LC3 and the level of autophagy (Kabeya et al., 2000), we measured by western blot the level of LC3-II during starvation with or without the lysosome inhibitor bafilomycin A1 (BafA1). As a result, transiently expressed Arf1T31N–HA significantly suppressed the starvation- and BafA1-induced elevation of LC3-II (Fig. 1C). These data are consistent with previous observations in yeast (Hamasaki et al., 2003) and suggest that the secretory pathway is essential to autophagosome formation in mammalian cells.

**Increased TGN-to-plasma traffic stimulates the formation of LC3-vesicles**

To test whether increased secretory traffic affects the formation of autophagosomes, we visualized the change in intracellular localization of GFP–LC3 during release of a bolus of the thermo-reversible folding mutant, ts045 vesicular stomatitis virus G protein fused to fluorescent protein Cherry at its cytoplasmic tail (VSVG–Cherry), from the ER into the secretory pathway by shifting the temperature from 40°C to 32°C (Bergmann, 1989; Presley et al., 1997). There were no dramatic changes in GFP–LC3 distribution when VSVG–Cherry was localized in the ER at 40°C or reached the juxtanuclear Golgi complex within 40 minutes after changing the temperature (Fig. 2A). Unexpectedly, in ~45% of cells expressing VSVG–Cherry and GFP–LC3, by 120 minutes

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**Fig. 1. The early secretory pathway is essential for autophagosome formation.**

(A) HEK293 cells transiently expressing GFP–LC3 or GFP–LC3 and HA-tagged Arf1 or Sar1 mutants, were incubated in starvation medium for 1 hour and imaged by confocal microscopy. Scale bars: 10 μm. (B) Statistical analysis of the numbers of LC3 dots per cell in A. Quantification of autophagosomes per cell was done using the Axiovision automatic measurement program on the Zeiss LSM510 Meta as described in the Materials and Methods. The values reported are means ± s.e.m.; ***p<0.0001 vs control. (C) HEK293 cells with or without Arf1T31N–HA expression were cultured in starvation medium with or without BafA1 for 1 hour; the cellular LC3 level was assessed by western blot. The LC3-II to LC3-I ratio was evaluated by densitometric analysis.
after the temperature shift, when a large amount of the VSVG–Cherry left the Golgi complex and arrived at the plasma membrane, a large number of spot-like GFP–LC3-positive structures appeared in the cytoplasm, representing increased autophagic vesicles (Fig. 2A). This observation implies that increased Golgi-to-plasma-membrane (PM) traffic promotes the basal level of autophagosomes.

To confirm that the Golgi-to-PM transport in the secretory pathway enhances autophagosome formation, we designed another experiment by shifting the culture temperature of HEK293 cells expressing GFP–LC3 and VSVG–Cherry from 40˚C to 20˚C for 2 hours to directly accumulate VSVG–Cherry in the TGN, then changing the temperature to 32˚C, allowing the cargo to go to the PM (Griffiths et al., 1985; Matlin and Simons, 1983). Surprisingly, at 20˚C, in ~80% of the cells, GFP–LC3 was found to strongly associate with the TGN as demonstrated by colocalization of GFP–LC3 with VSVG–Cherry in the perinuclear region (Fig. 2B). Changing the temperature from 20˚C to 32˚C for 15 minutes, when part of the VSVG–Cherry left the TGN for the plasma membrane, the GFP–LC3 started to bud off from the TGN separate from the VSVG–Cherry. After changing the temperature for 60 minutes, when more VSVG–Cherry left the TGN, GFP–LC3 dispersed into small-spot structures, which morphologically resembled autophagosomes (Fig. 2B).

To determine the specificity of the association of GFP–LC3 with the TGN, we introduced a mutation changing the Gly120 to Ala (LC3G120A) in LC3, which affects the cleavage at the C-terminal region of pro-LC3 and results in failure to form LC3-I and LC3-II (Kabeya et al., 2000). When expressed in the cells with VSVG–Cherry, GFP–LC3G120A never bound to the TGN with the same loading of VSVG–Cherry onto the TGN by shifting the temperature from 40˚C to 20˚C for 2 hours (Fig. 2C). This result strongly suggests that binding of LC3 to the TGN is mediated by a specific interaction between LC3 and the TGN membrane, and TGN-associated LC3 is the membrane-bound LC3–PE.

To exclude the possibility that these events were purely a result of temperature change, we performed the same experiments in cells expressing LC3–GFP only or LC3–GFP with pmCherry-C1. The results showed that the temperature change itself did not stimulate the membrane association of LC3 and the subsequent production of LC3-positive vesicles in any of the cells (supplementary material Fig. S1).

Blockade of post-Golgi transport prevents starvation- or rapamycin-induced autophagosome formation

To obtain further evidence that TGN-to-PM traffic regulates autophagosome formation, we then assessed the generation of autophagosomes when TGN-to-PM transport was blocked. Clathrin adaptor protein 180 (AP180) and EGFR pathway substrate clone 15 (Eps15) mediate the formation of TGN-derived vesicles. Overexpression of an AP180 C-terminal domain (AP180C, residues 530–915) or an Eps15 deletion mutant lacking the second and third N-terminal EH domains (Eps15A95/295) has a dominant-negative effect on clathrin coating at the TGN (Chi et al., 2008; Zhao et al., 2001). When expressed in cells, AP180C–YFP and YFP–Eps15A95/295 almost fully prevented the transport of VSVG–CFP from the TGN to the PM (Fig. 3A) (Chi et al., 2008). In these cells, induction of autophagosome-like vesicles by releasing VSVG–CFP from TGN-to-PM transport was also diminished, although Cherry–LC3 was still recruited to the TGN by VSVG loading (Fig. 3B,C). The effect of AP180C–YFP and YFP–Eps15A95/295 expression was also determined in autophagosome formation induced by starvation. Starvation that induced typical autophagy in control cells resulted in fewer autophagosomes in cells expressing AP180C–YFP or YFP–Eps15A95/295 (Fig. 3D,F), suggesting a fundamental role of exit from the TGN in the regulation of autophagosome biogenesis.

Fig. 2. Enhanced TGN-to-PM transport stimulates autophagosome formation. (A) HEK293 cells transiently expressing GFP–LC3 and VSVG–Cherry were incubated at 40˚C overnight to retain VSVG–Cherry in the ER. Then the cells were imaged over time upon shifting the culture temperature from 40˚C to 32˚C. (B) HEK293 cells expressing GFP–LC3 and VSVG–Cherry were incubated at 40˚C overnight, followed by culture for 2 hours at 20˚C to retain VSVG–Cherry in the TGN. Then the culture temperature was shifted to 32˚C, and the cells were imaged at indicated time points after the temperature shift. (C) HEK293 cells expressing GFP–LC3G120A and VSVG–Cherry were incubated at 40˚C overnight, followed by 2 hours culture at 20˚C to retain VSVG–Cherry in the TGN and imaged. All fluorescence images were confocal images of optical slice thickness ~1 μm. Scale bars: 10 μm.
AP180C or Eps15Δ95/295 removes clathrin from not only the TGN but also the PM (Benmerah et al., 1998; Ford et al., 2001; Lui-Roberts et al., 2005). To ensure that the observed reduction of autophagosomes in AP180 and Eps15 mutant cells was a direct effect on the TGN and did not result from disruption of endocytosis, we used another mutant Eps15 protein, which lacked a 14 amino acid motif (Eps15\textsuperscript{D14aa}). Expression of Eps15\textsuperscript{D14aa–MYC} selectively reduces the exit of secretory proteins from the TGN by binding to AP1, but not AP2 (Chi et al., 2008). We found that expression of Eps15\textsuperscript{D14aa–MYC} also dramatically suppressed starvation-induced autophagosome formation in HEK293 cells (Fig. 3E,F).

Finally, we measured the LC3–PE level in cells overexpressing AP180 or Eps15 deletion mutants by western blot. As expected, the starvation- and BafA1-induced elevation in LC3–PE level was clearly suppressed (Fig. 3G), indicating that autophagosomes failed to form in these mutant cells. Collectively, these data suggest that not only the constitutive TGN-to-PM traffic but also general transport from the TGN is required for the formation of autophagosomes.

**AP1 localizes to starvation- and rapamycin-induced autophagosomes**

Our results showing that AP180 and Eps15 mutants disrupted the formation of LC3-positive vesicles from the TGN strongly suggested a requirement for a clathrin coat in the process. Because recruitment of clathrin to the TGN is mainly mediated by its adaptor protein AP1, it can assemble into a coat lattice, therefore we checked the distribution of AP1 during autophagy. HEK293 cells were transiently transfected with AP180-YFP or Eps15Δ14aa–Myc. Cells were starved for 1 hour with or without BafA1, and analyzed by western blot. The LC3-II to LC3-I ratio was evaluated by densitometric analysis.
γ-adaptin (a subunit of the AP1 complex) and Cherry–LC3. In these cells, γ-adaptin–GFP presented a typical cytosol and TGN localization. Surprisingly, upon starvation for 1 hour or rapamycin treatment for 20 hours, in ~30% of cells, γ-adaptin–GFP redistributed to the formed autophagosomes showing a perfect colocalization with membrane-bound Cherry–LC3. In these cells, nearly every autophagosome contained γ-adaptin–GFP (Fig. 4A).

Previous studies have shown that Atg9 localizes to the TGN and cycles between the TGN and its peripheral pool during autophagy (Young et al., 2006). To further determine the action of AP1, especially in the early stage of autophagy, we expressed γ-adaptin–GFP and Cherry–Atg9 in the cells and analyzed their distribution during starvation or rapamycin treatment. At a very early phase of starvation (20 minutes) or rapamycin treatment (10 hours), colocalization of γ-adaptin–GFP and Cherry–Atg9 was detected (Fig. 4B). The specificity of AP1–LC3 and AP1–Atg9 colocalization was verified by visualizing GFP–α-adaptin (a subunit of the AP2 complex) with Cherry–LC3 and GFP–α-adaptin with Cherry–Atg9. As a result, neither starvation nor rapamycin treatment caused colocalization of AP2 with LC3 or Atg9 in any of the cells (supplementary material Fig. S2).

We also analyzed the colocalization of LC3 with clathrin during starvation. In GFP–LC3-expressing HEK293 cells, after starvation, the cells were stained with a specific anti-clathrin heavy-chain antibody. We found that in some cells (~30%), clathrin also distributed to the GFP–LC3-positive autophagosomes (Fig. 4C). These data thereby strongly suggested an involvement of AP1 and the clathrin coat in the formation of autophagosomes.

Because the colocalization of AP1 or clathrin with GFP–LC3-positive autophagosomes was only found in ~30% of cells, we then tested whether the remaining LC3-positive structures colocalized with other membrane organelles such as ER or mitochondria. Using a YFP-tagged mitochondrial matrix protein as a marker for mitochondria and a GFP-tagged KDEL receptor for the ER, we found that during starvation, Cherry–LC3 rarely colocalized with these markers (supplementary material Fig. S3).

**LC3 binds to the TGN and forms LC3 vesicles from the TGN during starvation-induced autophagy**

We next asked whether we could detect the outgrowth of similar vesicles from the TGN membrane in response to starvation. In fact, in cells overexpressing LC3, we often found high LC3 signals in the peri-nuclear region during autophagy. We therefore chose to stain starved HEK293 cells with antibodies against LC3 and TGN46 to check the intracellular localization of these endogenous proteins. We found that as early as 15 minutes of starvation, endogenous LC3 accumulated in peri-nuclear TGN46-positive sites (Fig. 5A). Nevertheless, LC3 was segregated from TGN46, suggesting it is associated with a unique non-TGN46-containing TGN sub-compartment. Over 30 minutes of starvation, the LC3 started to disperse from the TGN area and distribute randomly in the cytoplasm (Fig. 5A).

To determine the localization of LC3 on the TGN more precisely, we took a series of images through the Z-axis of cells stained with antibodies against LC3 and TGN46. The stacks were reconstructed to create 3D images. A typical 3D image of a cell starved for 20 minutes showed that on the TGN, most of the LC3 did not fuse with TGN46 (Fig. 5B), confirming that LC3 was associated with a non-TGN46-containing sub-compartment of the TGN membrane.

We further visualized the budding process of GFP–LC3 from the TGN during starvation-induced autophagy in living cells. We found in some cells that GFP–LC3-containing membrane pulled off from the TGN as tubular processes that extended for several micrometers (supplementary material Fig. S4). GFP–LC3
accumulated at the tips of these tubules, where they formed a ball-like mass. After a variable time, the tip regions detached and moved outward as separate elements. Sometimes during the detachment, the tip region was further divided into two vesicles (Fig. 5C).

We performed the same experiments to observe the dynamics of the GFP–LC3G120A mutant during starvation. We found, similar to VSVG loading (Fig. 2C), starvation never recruited the GFP–LC3G120A to the peri-nuclear region (Fig. 5D), further confirming a specific interaction of LC3–PE and the TGN membrane. Taken together, these data suggest that the TGN membrane is a source of membrane for autophagosomal structures.

Separation of LC3-containing vesicles from TGN-to-endosome traffic

Once formed, autophagosomes are required to fuse with the endosomal compartments and further fuse with lysosomes for maturation. Although the role of the endosomal system in autophagosome formation remains to be understood, results from yeast studies indicate that endosomes might not be essential for autophagosome assembly (Reggiori et al., 2004). The observation that LC3-positive vesicles bud off from a distinct domain of the TGN and remain separated from the constitutive cargos (VSVG) prompted us to clarify whether this occurs in TGN-to-endosome transport. We first observed the location of TGN38 (rat homolog of human TGN46) during autophagosome formation triggered by enhanced TGN-to-PM transport. We imaged the distribution of LC3–Cherry and CFP–TGN38 over time after shifting from 20°C to 32°C. Shown are the distributions of LC3–Cherry and CFP–TGN38 at 60 minutes after the shift. (B) HEK293 cells transiently expressing LC3–Cherry and CFP–CI-MRP were left untreated or starved for 1 hour and imaged by confocal microscopy. C, control; S, starvation. Scale bars: 10 μm.
TGN46 during release of VSVG–CFP from the TGN. We found that, similar to the observations in starved cells, YFP–LC3 on the TGN caused by VSVG–CFP loading was initially segregated from TGN46, and this segregation became clearer when VSVG began to leave the TGN (supplementary material Fig. S5).

We further visualized the mannose-6-phosphate receptors (MPRs) in starvation-induced autophagy. In cells co-transfected with CFP–CI-MRP and Cherry–LC3, during the entire process of starvation, MRP rarely targeted to the LC3-positive vesicles (Fig. 6B).

Taken together, these results confirmed that the LC3-positive vesicles budded from the TGN contain neither the constitutive cargo nor the components of TGN-to-endosome traffic.

**Knockdown of AP1, but not AP2, inhibits autophagy**

Given that AP1 and clathrin were targeted to autophagosomes, we performed RNAi to identify the necessity for AP1 and clathrin in autophagosome formation, with siRNA against AP2 as a control. The efficacy of the designed siRNA was confirmed by western blot (Fig. 7A). The effect of the siRNA treatment was assessed first by immunostaining. Compared with the control cells in which γ-adaptin or clathrin heavy-chain presented normal expression and peri-nuclear localization, knockdown cells displayed much fainter staining and a lack of TGN association and the number of starvation-induced LC3-positive autophagosomes was dramatically reduced (Fig. 7B,C). Data from western blots confirmed that knockdown of γ-adaptin or clathrin heavy-chain in HEK293 cells dramatically reduced the lipidated LC3-II levels triggered by starvation, with or without BafA1 treatment (Fig. 7D). Interestingly, the suppression of autophagosome number and LC3-II level was not detected in AP2-knockdown cells (Fig. 7B,C,D). These findings strongly suggest that AP1-mediated clathrin coating in the TGN but not AP2-mediated endocytosis plays a crucial role in starvation-induced autophagosome formation.

**Discussion**

The debate on the origin of the autophagosomal membrane and the formation of the autophagosome remains the most pivotal question for understanding autophagy. In this study, through our analysis of AP1-mediated events, we have shown that functional TGN membranes are required for autophagosome formation. Our data showed how modulation of the secretory traffic modified the formation of autophagosomes in mammalian cells. By inactivation of the small GTPases that function in the early secretory pathway, we determined in animal cells the essential role of ER export in starvation-stimulated autophagy, which is consistent with the conclusion reached in yeast studies (Hamasaki et al., 2003; Ishihara et al., 2001). Nonetheless, owing to the close relationship between differential membrane trafficking routes, interruption of the early secretory pathway alters the late...
transport to a great degree and breaks down the Golgi structure, which is central to intracellular trafficking. Observation of the time course of cargo flow and specific modulation of the TGN-to-PM traffic allowed us to determine that the post-Golgi transport regulated the process of autophagosome formation. Although surprising, it is not unreasonable to propose that a forced increase in the export of constitutively secreted proteins from the TGN stimulates the formation of autophagosomes. Many studies have suggested that lateral segregation in the TGN is the primary sorting event and that there is potential interdependence between the different domains (Gleson et al., 2004; Hirschberg et al., 1998; Keller et al., 2001). In addition to the known necessity of AP1 and clathrin for TGN–endosome transport, the fact that dysfunction of AP1 and clathrin impeded VSVG transport to the PM supports the domain segregation model. Instead of functioning directly in the formation of TGN-to-PM carriers, AP1 and clathrin contribute to the constitutive cargo transport, possibly by facilitating the lateral segregation of the TGN membrane. Our results suggest the existence of specific discrete domains in the TGN membrane to which LC3 is recruited and from which the LC3-containing vesicles bud off. These specific domains are exclusive of the constitutive cargos and components destined for the endosomes; the formation and subsequent budding off of these domains are influenced by the post-Golgi traffic. This can explain not only the recent results from the yeast Saccharomyces cerevisiae showing that the post-Golgi Sec proteins and Golgi exit are required for autophagy (Geng et al., 2010; van der Vaart et al., 2010), but also an early report on animal cells demonstrating that post-Golgi but not ER or cis-Golgi membrane proteins are included in the limiting membranes of autophagosomes (Yamamoto et al., 1990). In addition, our results strongly imply that the formation of these LC3-positive vesicles is used by the cell to regulate intracellular membrane partitioning and redistribution.

Our finding showing the recruitment of LC3 to the TGN suggests that the TGN is not only a donor site for early LC3 assembly but also a membrane source for the formation of double-membrane autophagosomes. Instead of an accumulation of formed autophagic vesicles to the TGN, our observations indicate a direct association of LC3 with the TGN membrane in response to the presence of a mass of VSVG at the TGN or starvation-initiated signaling. During the time course of VSVG loading, very few LC3-containing vesicles were observed in the cytosol before VSVG arrived at the TGN, and LC3 vesicles were formed only when VSVG started to leave the TGN. Similarly, in starved cells, accumulation of LC3 in the peri-nuclear area occurred at a very early stage before LC3 vesicles were present in the cytosol. That the expression of AP180C or Eps15Δ95/295 blocked the formation of LC3 vesicles but not the VSVG-controlled association of LC3 with the TGN membrane further supports this conclusion. With regard to the membrane association of LC3, in vitro studies have pointed out the involvement of the orderly membrane recruitment of a series of Atg proteins, including the VPS34–beclin-1 and Atg5–Atg12–Atg16 complexes. Nevertheless, the precise role of these complexes and the underlying mechanism of LC3 binding, especially in mammalian cells, are still unclear. Our data suggest that a crucial step is the specific association of lipid-modified LC3 with TGN membranes. Possibly, during autophagy, upon association with the specific domain of the TGN, LC3 vesicles carry LC3 to the isolation membranes derived from the ER or other membrane compartments. During these processes, Atg9 plays an essential role in facilitating the docking of LC3–PE with the complexes, and AP1-mediated clathrin coating contributes to the budding off of LC3 vesicles from the TGN, although it is currently unknown whether the Atg5–Atg12–Atg16 complex exists in the TGN.

Our results are partially inconsistent with a recent report showing that knockdown of clathrin and AP2, but not AP1, inhibits the formation of autophagosomes (Ravikumar et al., 2010). However, the effect of knockdown of AP2 or epsin-1 on autophagosome formation appeared to be weak in the presence of BafA1, but not in samples without BafA1 treatment, and this has been interpreted as a possible decrease in LC3 II level in the absence of BafA1 and a low gene knockdown effect (Ravikumar et al., 2010). In our experimental system, even in the absence of BafA1, we showed clearly that knockdown of AP1 but not AP2 blocked starvation-initiated autophagosome formation. This was also confirmed by the effect of overexpression of the specific Eps15 mutant Eps15Δ14aa. Combined with the association of LC3 with the TGN and LC3 vesicles budding off the TGN, our observations support the conclusion that clathrin regulates the formation of autophagosomes by mainly functioning in the TGN but not the PM. We do not exclude a possible contribution of the PM to the autophagosome, at least in part because of the intimate membrane traffic between the TGN and the PM. To further clarify this issue, it will be especially crucial and interesting to find out whether the Atg16-positive and LC3-negative vesicles formed through clathrin–Atg16 interaction (Ravikumar et al., 2010) fuse with the TGN membrane (not the early or medial Golgi).

Materials and Methods

**DNA constructs, reagents and antibodies**

Arf1T31N–HA, Sar1T39N–HA, Sar1H79G–HA, VSVG–CFP, VSVG–Cherry, KDEL–GFP, YFP–Eps15Δ95/295 and CFP–TGN38 were described previously (Peters et al., 1995; Presley et al., 1997; Puertollano et al., 2001; Scales et al., 1997; Ward et al., 2001; Wu et al., 2003; Zaal et al., 1999). γ-adaptin–GFP, CFP–CI-MPR and AP180C–GFP were from Juan S. Bonifacino (National Institutes of Health, Bethesda, MD). Mito–YFP was from Clontech. GFP–α-adaptin was from Lois E. Greene (National Institutes of Health, Bethesda, MD). Eps15Δ14aa–Myc was kindly provided by Mark A. McNiven (Mayo Clinic College of Medicine, Rochester, MN). The cDNA of LC3 was a gift from Yoshinori Ohsumi (Tokyo Institute of Technology, Tokyo, Japan). GFP–LC3 was made by cloning the cDNA of LC3 into a PEGFP-C1 vector (Clontech Laboratories) using the BglII(5’) and SalI (3’) restriction sites. Cherry–LC3 or YFP–LC3 were made by changing the GFP in GFP–LC3 plasmid to mCherry or YFP, using the AgeI and BspEI restriction sites. The point mutation for glycine to alanine at position 120 of LC3 (GFP–LC3G120A) was created by PCR-based site-directed mutagenesis using LC3 sense primer (5’-GCTCCAGGACGCTCCGGAACACTGCTGCGTACCATAGG-3’) and LC3 antisense primer (5’-GTATGTAACAGCCAGTGCTGCGGACGTC-CTCCGGAGGCC-3’). Cherry–GFP was made by cloning the human autophagy 9-like 1 protein ORF (GenBank: BK004018.1) into a pmCherry-C1 vector (Clontech) using the EcoR1 (5’) and XbaI (3’) restriction sites. Rapamycin and BafA1 were purchased from Sigma and used at 100 nM.

The following antibodies were used: rabbit polyclonal antibody against LC3, mouse monoclonal antibody against β-actin (Sigma); mouse monoclonal antibodies against γ-adaptin (a subunit of AP1), α-adaptin (a subunit of AP2) and GFP (BD Biosciences); sheep polyclonal antibody against TGN46 (AbD Serotec); mouse monoclonal antibody against Myc and HA (Santa Cruz); mouse monoclonal antibody clathrin heavy chain (Cell Signaling). Alexa-Fluor-488- and Alexa-Fluor-546-tagged secondary antibodies were from Molecular Probes. Secondary antibodies goat anti-rabbit IRDye 800CW and goat anti-mouse IRDye 680 were from LI-COR Biosciences.

**Cell culture and transfection**

HEK293 cells were grown in DMEM supplemented with 10% FBS, 2 mM glutamine, 100 U/ml penicillin and 100 U/ml streptomycin, at 37°C under 5% CO2. Transient transfections were performed using Lipofectamine 2000 according
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