The Chaperone BAG6 Regulates Cellular Homeostasis between Autophagy and Apoptosis by Holding LC3B

**HIGHLIGHTS**

- AMFR and USP13 regulate CASP3-mediated BAG6 cleavage.
- CASP3 cleaves BAG6 and the cleaved N-terminal BAG6 binds to LC3-I.
- BAG6 binds to LC3B-I through its LIR1 motif to inhibit autophagosome formation.
- BAG6 regulates cellular homeostasis between autophagy and apoptosis.

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**Unstressed**

- AMFR and USP13 balanced
- Full-length BAG6
- Autophagy
- Cell survival

**ER stress**

- AMFR overexpression or USP13 knockdown
- Ub-TXN
- CASP3 activation
- N-terminal BAG6
- LC3-I/Pro-LC3
- Apoptosis
- Cell death
The Chaperone BAG6 Regulates Cellular Homeostasis between Autophagy and Apoptosis by Holding LC3B

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SUMMARY
AMFR/gp78 and USP13 are a pair of ubiquitin ligase and deubiquitinase that ensure the accuracy of endoplasmic reticulum-associated degradation (ERAD). Depletion of USP13 leads to caspase activation and cleavage of the ER chaperone BAG6, which is reversed by knockdown of AMFR. However, the mechanism and physiological relevance of this regulation are still unclear. Here, by using the NEDDylator system, we screened out TXN as a substrate of AMFR and USP13 and showed its involvement in regulating CASP3 activation and BAG6 cleavage. Furthermore, we showed that the cleaved N-terminal BAG6 is located in the cytosol and interacts with both LC3B-I and unprocessed form of LC3B (Pro-LC3B) through the LIR1 motif to suppress autophagy. An NMR approach verified the direct interaction between BAG6 LIR1 and LC3B-I or Pro-LC3B. Collectively, our findings uncover a mechanism that converts BAG6 from an ERAD regulator to an autophagy tuner and apoptosis inducer during ER stress.

INTRODUCTION
In eukaryotic cells, endoplasmic reticulum-associated degradation (ERAD) plays a crucial role in maintaining cellular homeostasis by monitoring and degrading misfolded or unfolded proteins in the ER (Brodsky, 2012; Preston and Brodsky, 2017; Ruggiano et al., 2014; Smith et al., 2011). Dysfunction in the ERAD pathway can lead to irreversible ER stress, which eventually activates cellular apoptosis pathway (Hetz, 2012; Hetz and Papa, 2018; Tabas and Ron, 2011). Various E3 ubiquitin ligases selectively recognize these different aberrant substrates. At the same time, some deubiquitinases (DUBs) are also recruited to keep the specificity of modification. Among them, AMFR/gp78 and USP13 are a pair of ubiquitin ligase and deubiquitinase functioning in ERAD (Chen et al., 2006; Fang et al., 2001; Liu et al., 2014). Previously we have reported that AMFR ubiquitinates not only ERAD substrates but also ERAD machinery protein UBL4A, one of the components of BAG6 complex, leading to the disassembly of BAG6 complex. On the other hand, USP13 antagonizes AMFR by deubiquitinating UBL4A to maintain the integrity of BAG6 complex and therefore promotes ERAD (Liu et al., 2014). Previously, we and other groups have demonstrated that BAG6 is a multifunctional chaperone protein that can hold hydrophobic domain of tail-anchored (TA) membrane proteins (Leznicki et al., 2010; Mariappan et al., 2010) or misfolded proteins (Ernst et al., 2011; Hessa et al., 2011; Mariappan et al., 2010; Minami et al., 2010; Wang et al., 2011) and direct them to distinct destinations. Therefore, the integrity of BAG6 complex is necessary for holding client proteins from aggregation. Additionally, we have also found that overexpression of AMFR or knockdown of USP13 results in the cleavage of BAG6 by CASP3 (Liu et al., 2014). A CASP3 cleavage site, DEQD, has been reported in the C terminus of BAG6 (Wu et al., 2004). However, the detailed mechanisms of how CASP3 is activated to cleave BAG6 in response to USP13 depletion are still unknown.

Autophagy is a highly conserved degradation process that recycles cellular components and defends against intracellular pathogens (Bento et al., 2016; Mizushima et al., 2011). It is initiated from the biogenesis of a crescent-shaped isolation membrane (IM), named phagophore, which expands to form a double membrane vesicle called autophagosome. The autophagosome contains cellular components including damaged organelles, protein aggregates, and invasive microbes, which are degraded by the hydrolases after autophagosome fusion with lysosome (Galluzzi et al., 2014, 2017; He and Klionsky, 2009). The initiation and maturation of autophagosome are strictly regulated by autophagy-related proteins. Upon autophagy induction, the activated ULK1/ATG13/RB1CC1 kinase complex transfers to the autophagosome formation site enriched with phosphatidylinositol (PtdIns) (Karanasios et al., 2013; Nishimura et al., 2017).
BAG6 participates in a variety of cellular processes, including protein quality control, gene regulation, and apoptosis (Lee and Ye, 2013). BAG6 was initially found as an apoptosis regulator by interacting with Reaper, a central apoptosis effector in Drosophila (Thress et al., 1998; Thress et al., 1999). Subsequent studies discovered BAG6 functions as an anti-apoptotic regulator and interacts with multiple apoptotic modulators (Desmots et al., 2008; Kumar et al., 2004; Minami et al., 2007). Depletion of Bag6 in mouse causes cell death and proliferation defects, resulting in embryonic lethality (Desmots et al., 2005). Besides, BAG6 is necessary for DNA damage-induced apoptosis by controlling the TP53/p53 signaling pathway in human cells (Sasaki et al., 2007). Interestingly, it has been recently reported that BAG6 also promotes autophagy by transporting EP300/p300 into the nucleus to hyperacetylate ATG7 to suppress autophagy (Sebti et al., 2014a, 2014b). Therefore, it seems that BAG6 may function in both apoptotic and autophagic pathways. We then questioned whether BAG6 is a switcher of autophagy and apoptosis during ER stress.

Here, by using a NEDDylator system, we identified the substrates of AMFR and USP13, which might mediate the activation of CASP3 and consequence of BAG6 cleavage. Based on mass spectrometry analysis, we have identified multiple substrates and demonstrated TXN (thioredoxin) as a substrate of AMFR and USP13 involved in CASP3 activation and BAG6 cleavage. Ubiquitinated TXN accompanies CASP3 activation to cleave BAG6. Cleaved BAG6 remains in the cytosol and interacts with excessive LC3B-I or the unprocessed form of LC3B (Pro-LC3B) through the LIR1 motif to suppress autophagy. Our NMR data further verified that BAG6 binds to LC3B-I or Pro-LC3B through some specific amino acids and BAG6 binding to LC3B-I can stabilize LC3B-I structure. Therefore, our study provides evidence that ERAD machinery proteins can regulate the balance between autophagy and apoptosis.

RESULTS
Depletion of USP13 Induces Apoptosis and Suppresses Autophagy

Impaired ERAD pathway causes prolonged ER stress, which activates alternative signal pathways and promotes apoptosis (Hetz, 2012; Hetz and Papa, 2018; Tabas and Ron, 2011). USP13 knockdown results in loss of ER homeostasis and prolonged ER stress, so we examined whether USP13 depletion activates apoptosis in cells. As shown in Figure 1A, the cell viability was decreased in USP13 knockdown cells, and the knockdown cells were tended to apoptosis compared with the control cells. MG132 is a proteasome inhibitor and has been shown to induce apoptosis (Emanuele et al., 2002; Giuliano et al., 1999; Lauricella et al., 2003; Maclaren et al., 2001; Rock et al., 1994). It is easier to detect the apoptotic change in cells with MG132 treatment. In our experiment, with MG132 treatment, the cell viability of both the control and USP13 knockdown cells was significantly decreased (Figure 1A). Because apoptosis and autophagy are two pathways intertwined in cells to control cell fate, we examined whether USP13 could regulate autophagy. To monitor the autophagic flux, we employed a HeLa cell line stably expressing mRFP-GFP-LC3B (Yan et al., 2018). The autophagosomes bearing LC3B appeared as yellow puncta (with both GFP and RFP signals), but once fused with lysosomes, the autolysosomes were detected as red puncta (with only RFP signal in lysosome) because green fluorescence signal was quenched by low pH. Interestingly, we found that, under both normal and starvation (EBSS treatment) conditions, USP13 knockdown substantially reduced the number of yellow puncta (autophagosomes) (Figures 1B and 1C). In contrast, there was no significant difference in the number of red puncta (autolysosomes) between the control and USP13 knockdown cells (Figure 1C). These data suggest that USP13 knockdown may suppress the early step of autophagy. Chloroquine (CQ) is a drug that inhibits the fusion of autophagosome with lysosome, which leads to the accumulation of autophagosomes in cells. It is commonly used to measure changes in autophagic flux. The accumulation of yellow puncta was reduced in USP13 knockdown than in the control cells under CQ treatment (Figures 1B and 1C).
Figure 1. Depletion of USP13 Induces BAG6 Cleavage and Suppresses Autophagy

(A) USP13 knockdown decreases cell viability with or without MG132 treatment. Cells were transfected with control shRNA, USP13 shRNA-1, or USP13 shRNA-2. Cells were treated with DMSO or MG132 (10 μM) for 12 h. Cell viability was measured by MTS assay. Data are represented as mean ± SD from three independent experiments. *p < 0.05 (one-way ANOVA).
We showed previously that knockdown of USP13 promotes the cleavage of BAG6. This phenotype is further enhanced by treating cells with the proteasome inhibitor MG132 (Liu et al., 2014). To explore the physiological function of this regulation, we first identified the protease responsible for BAG6 cleavage under these conditions. Since the cleavage site is predicted to be close to the C terminus, which harbors a known caspase cleavage site, we tested whether BAG6 cleavage induced by USP13 depletion could be due to the activation of caspase. Interestingly, we found that CASP3 was robustly cleaved and activated in USP13 knockdown cells compared with that in control cells (Figure 1D). We then tested BAG6 cleavage in the control and USP13 knockdown cells. Cells were solubilized by an NP40-containing lysis buffer. We analyzed both the NP40-soluble and -insoluble fractions by immunoblotting, because we found previously that cleaved BAG6 preferentially accumulates in NP40-insoluble fractions. As expected, knockdown of USP13 generated a fast migrating BAG6 species, which was more accumulated in the presence of MG132 but was blocked with the treatment of a pan-caspase inhibitor Z-VD-FMK (Figure 1E). It was reported previously that MG132 treatment results in CASP3 and CASP7 activation (Emanuele et al., 2002; Giuliano et al., 1999; Gray et al., 2010; Lauricella et al., 2003). The treatment of MG132 in our assay is to magnify the effect of caspase activation. It is worth pointing out that, with MG132 treatment, in some cases BAG6 is also cleaved in the control cells. Similar observation was obtained in CRISPR cells that have USP13 deficiency-induced BAG6 cleavage (Figure 2A). A single mutation introduced at the cleavage site (D995A) completely abolished cleavage by the caspases (Figure S1E). These results suggest that USP13 depletion activates caspases and induces BAG6 cleavage.

**A NEDDylator System to Screen AMFR Substrates**

We previously reported that USP13 antagonizes E3 AMFR-mediated ubiquitination in the ERAD pathway. Interestingly, overexpression of AMFR also promotes BAG6 cleavage (Liu et al., 2014), whereas knockdown of AMFR in USP13 knockdown cells suppresses USP13 deficiency-induced BAG6 cleavage (Figure 2A). Thus, it seemed that the two enzymes may antagonize each other to modify substrate ubiquitination during apoptosis. To screen for such substrates, we employed a well-developed NEDDylator system, which captures the substrates of a specific E3 by conjugating the target proteins with ubiquitin-like NEDD8 molecule (Figure 2B) (Zhuang et al., 2013).
**Figure 2. Screening of AMFR Substrates by the NEDDylator System**

(A) The membrane fraction BAG6 is more cleaved in response to USP13 knockdown and is rescued by co-transfecting with AMFR shRNA. Cells were transfected with control shRNA, USP13 shRNA-1, AMFR shRNA-1, or (USP13 shRNA-1 + AMFR shRNA-1) and were treated with MG132 (10 μM, 12 h). Samples were prepared using the same procedure as in Figure 1E and were immunoblotted with the indicated antibodies.

(B) Schematic representation of the procedure for identifying AMFR substrates using the NEDDylator system. NEDDylator is transfected into AMFR KO cells, cell lysates are immunoprecipitated with Ni-NTA beads, eluted with streptavidin beads, and purified proteins are identified by mass spectrometry.

(C) Heatmap showing the expression levels of different proteins in response to NEDDylator treatment. The proteins are grouped into three categories: TM domains, RING, and CUE-G2BR.

(D) Western blot analysis of whole cell extracts treated with different NEDDylator constructs. The blot shows the expression of AMFR and NEDD8 proteins.

(E) List of proteins identified using mass spectrometry. The proteins are categorized into four groups: ER-localized proteins and ERAD-related proteins, mitochondria-localized proteins, apoptosis-related proteins, and autophagy-related proteins.

- **ER-localized proteins and ERAD-related proteins**
  - AMFR
  - FAF2
  - HSPA1A
  - Pdia6
  - Hsp90b1
  - VCP
  - Bag6
  - Hspa5
  - Hspaa4l
  - Hspa4
  - Canx
  - Rpn1

- **Mitochondria-localized proteins**
  - Trp1
  - Hspd1
  - Hsdc17b10
  - Immi
  - Atp5a1
  - Etfb
  - Mdh2
  - Hspa9
  - Timm44

- **Apoptosis-related proteins**
  - Cse1l
  - Prdx2
  - Slc25a6
  - DSP
  - Txn
  - Pdia6
  - Slc25a5
  - Actn4
  - Dnm1l
  - Pdcd6ip
  - Flna
  - Pdia6
  - Api5
  - Hnrrnpk
  - Naca
  - Ddx42

- **Autophagy-related proteins**
  - Cisd2
  - Cltc
  - Anxa7
  - Atg6v1e1
  - Rab5c

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The membrane fraction BAG6 is more cleaved in response to USP13 knockdown and is rescued by co-transfecting with AMFR shRNA. Cells were transfected with control shRNA, USP13 shRNA-1, AMFR shRNA-1, or (USP13 shRNA-1 + AMFR shRNA-1) and were treated with MG132 (10 μM, 12 h). Samples were prepared using the same procedure as in Figure 1E and were immunoblotted with the indicated antibodies.
AMFR contains multiple transmembrane (TM) domains at the N terminus, followed by a RING, a CUE, a G2BR, and a VIM motif at the C terminus. The RING and G2BR domains are responsible for interacting with the ubiquitin-conjugating enzyme (E2) UBE2G2, the CUE domain recognizes ubiquitin chain on substrates, and the VIM motif interacts with VCP/p97. We first verified that AMFR itself has no NEDDylation activity (Figure S2A). To transfer NEDD8 to AMFR substrates, we designed three NEDDylators by fusing the NEDD8 E2 conjugating enzyme UBE2M with the different length of AMFR (Figure 2C). The resulting chimeras were named as NEDDylator<sup>ARATION</sup>, NEDDylator<sup>CU2+G2BR</sup>, and NEDDylator<sup>CUE</sup> (Figure 2C). We then transfected these three NEDDylators into AMFR knockout cells, together with His-biotin-tagged HB-NEDD8. As expected, only NEDDylator<sup>CUE+G2BR</sup> and NEDDylator<sup>CUE</sup>, but not NEDDylator<sup>ARATION</sup>, could transfer NEDD8 to FAF2 (Figure S2B). Therefore, NEDDylator<sup>CUE</sup> and NEDDylator<sup>CUE+G2BR</sup> were used for the later experiments.

Affinity purification and mass spectrometry identified 371 proteins that were modified by NEDD8 in the presence of NEDDylator<sup>CUE+G2BR</sup> and 110 proteins in the presence of NEDDylator<sup>CUE</sup> (Table S1, related to Figure 2). We then selected substrates that were possible targets that activate caspases and divided them into four groups: ERAD-related proteins, mitochondria-localized proteins, apoptosis-related proteins, and autophagy-related proteins. In each group, we found substrates previously identified as AMFR-interacting proteins including AMFR itself, FAF2 and BAG6 in the ERAD pathway; HSPD1 and IMMT on mitochondrion; and CSE1L as an apoptosis-related protein, which validates our approach of four groups based on the signal pathways they involved.

**AMFR Cooperates with USP13 to Regulate the Ubiquitination of TXN**

Among the identified proteins, we focused on TXN, a redox protein that can denitrosylate target proteins to regulate protein activity (Benhar et al., 2008; Mitchell et al., 2007; Rössig et al., 1999). TXN reduces S-nitrosylated CASP3 to CASP3 to activate apoptosis (Benhar et al., 2008). TXN maintains the activity of cytosolic CASP3, whereas TXN2 regulates the activity of mitochondrial CASP3 (Benhar et al., 2008).

To validate TXN as a substrate of AMFR, we first expressed both FLAG-TXN and AMFR constructs in cells and tested the interaction of the two proteins. The result showed that FLAG-TXN could pull down AMFR in cells (Figure S3A). In vitro GST pulldown assay using the transmembrane-deleted recombinant AMFR (GST-AMFRΔTM) showed that it stably interacted with the recombinant His-MYC-TXN (Figure S3C). The interaction was further verified at the endogenous level using an anti-AMFR antibody to co-immunoprecipitate TXN (Figure 3A). Furthermore, our data also showed that USP13 interacted with TXN at both the overexpressing and endogenous levels (Figures S3B and 3B). Next, we examined whether TXN is a substrate of AMFR and USP13. To test this idea, FLAG-TXN was transfected into cells together with the AMFR construct and an immunoprecipitation assay using anti-FLAG beads under denaturing condition was performed. The immunoblotting result showed that TXN-conjugated ubiquitin was noticeably increased in AMFR-overexpressing cells (Figure 3C). Consistent with this result, TXN-conjugated ubiquitin was also accumulated in USP13 knockdown cells (Figure 3D). These results verified that TXN is a substrate of AMFR and USP13.
We then tested the interaction of TXN and caspases. Upon induction of apoptosis by TNF-α treatment, FLAG-TXN was found to interact with CASP3, CASP7, and slight amount of CASP2 (Figure S3D). These data implied the possible activation of caspases by TXN. Interestingly, overexpression of TXN induced the cleavage of CASP3 and BAG6, which could be suppressed by knockdown of AMFR, indicating ubiquitination of TXN by AMFR might be essential for TXN denitrosylation activity (Figure 3E). On the other hand, CASP3 cleavage induced by AMFR overexpression or USP13 knockdown was completely blocked when the cells were co-expressing TXN knockdown shRNA (Figures 3F and 3G). To further examine whether TXN
reduces S-nitrosylated CASP3, we employed a biotin switch assay (Figure S3E). The biotin switch assay has been used extensively to measure protein S-nitrosylation (Forrester et al., 2009). We found a decrease of S-nitrosylated CASP3 (SNO-CASP3) in TXN-overexpressing cells (Figure S3F), suggesting TXN reduction of S-nitrosylated CASP3. Collectively, these results demonstrated that AMFR cooperates with USP13 to regulate the ubiquitination of TXN, which then influences S-nitrosylation of CASP3 and its activation, as well as BAG6 cleavage.

Cleaved BAG6 Suppresses Autophagy

The above results prompted us to speculate that the cleaved BAG6 is a potential regulator of autophagy and apoptosis. To see whether cleaved BAG6 could have a regulatory role in autophagy, we first transfected BAG6 and its mutants into HeLa cells stably expressing mRFP-GFP-LC3B.Cells overexpressing BAG6 greatly promoted yellow and red LC3B puncta formation under starvation condition (Figures 4A and 4C). Fluorescence imaging showed that BAG6 was exclusively localized in the nucleus (Figure 4B). In contrast, BAG6(1-991aa)-overexpressing cells had fewer yellow LC3B puncta compared with the control cells (Figures 4A and 4C). BAG6(1-991aa) was localized dominantly in the cytosol (Figure 4B). The caspase cleavage site mutant BAG6D995A, which was localized in the nucleus (Figure 4B), also induced yellow and red LC3B puncta formation similar to the full-length BAG6 (Figures 4A and 4C). The phenotype was further supported by the immunoblotting result, which showed that BAG6 and BAG6D995A promoted, whereas BAG6(1-991aa) inhibited, LC3B-I to LC3B-II transformation (Figure 4D).

To further address the specific importance of cleaved BAG6 in autophagic regulation, we tested BAG6 and its mutants in BAG6 knockout cells. Transiently expressing mRFP-GFP-LC3B was used to indicate the change of autophagy. Similarly to that observed in the wild-type background, BAG6 or BAG6D995A overexpression promoted, whereas BAG6(1-991aa) overexpression inhibited, yellow and red LC3B puncta formation compared with the control knockout cells (Figures S4A and S4B). These data suggest that cleaved BAG6 remains in the cytosol and prevents autophagy.

We then postulated that the cleaved BAG6 may interact with the cytosolic autophagy-related proteins to regulate autophagy. We therefore employed a co-immunoprecipitation assay to screen a collection of autophagy-related proteins to see which one interacts with BAG6. We found that LC3B was co-precipitated with FLAG-BAG6(1-991aa) (Figure 5A). Thus, BAG6 may be involved in an early stage of autophagosome formation. WIPI2 is an autophagy marker for early stage, and it is recruited to nascent autophagosomes by interacting with PtdIns3P at the onset of autophagosome formation (Dooley et al., 2015). Compared with the control, there was slightly more WIPI recruitment in BAG6- and BAG6D995A-overexpressing cells, but less in BAG6(1-991aa)-overexpressing cells (Figures S4C and S4D). These results suggest that the cleaved BAG6 suppresses autophagy at an early stage.

BAG6 Interacts with LC3B via the LIR1 Motif

Given that BAG6 is a holdase chaperone that interacts with the hydrophobic residues of the substrate for either translocation or degradation, we suspected that the cleaved BAG6 suppresses autophagy by holding LC3 from being lipidated. We first verified that the two proteins interacted with each other at the endogenous level by co-immunoprecipitation (Figure 5B). To see which domain of BAG6 is responsible for the interaction with LC3B, we expressed a series of BAG6 truncation mutants in cells and performed co-immunoprecipitation experiments. The study identified residues 88–266aa as the minimal domain responsible for the interaction with LC3B (Figures 5C and 5D). Interestingly, residues 88–266aa contains a region that was identified previously to participate in substrate hydrophobicity recognition (Tanaka et al., 2016).

Proteins interacting with LC3 usually bear a conserved linear sequence named LIR (LC3-interacting region). LIR sequences consist of a core motif W/F/Y-X-X-L/I/V (x = acidic or hydrophobic residues) flanked by N- and C-terminal sequences (Birgisdottir et al., 2013; Johansen and Lamark, 2020). LIR-containing proteins play crucial roles in autophagy including cargo recognition, autophagosome formation and maturation, as well as many autophagy-regulated signaling pathways (Birgisdottir et al., 2013; Johansen and Lamark, 2020). By sequence analysis, we identified four putative LIR motifs in BAG6 among the region of 89–387aa (LIR1 132–135: YVMV; LIR2 270–273: YVEV; LIR3 292–295: YEVL; LIR4 331–334: FVAL) (Figure 5C). To identify the functional LIR in BAG6, we mutated each of the four LIRs and analyzed the interaction of these mutants with LC3B by co-immunoprecipitation. The result showed that FLAG-BAG6(1-266aa) LIR1Y132A/V135A...
Figure 4. Cleaved BAG6 Remains in the Cytosol and Suppresses Autophagy

(A) HeLa cells stably expressing mRFP-GFP-LC3B were transiently transfected with FLAG-tagged BAG6 or its mutants. Cells were grown under starvation condition and analyzed for LC3B puncta. Nuclei are in blue. Scale bar, 5 μm.

(B) Subcellular localization of BAG6 and its mutants under starvation condition. HeLa cells were transiently transfected with various FLAG-tagged Bag6 or its mutants as in (A). Cells were grown under starvation condition and stained with anti-FLAG antibody in green. Nuclei are in blue. Scale bar, 5 μm.

(C) Quantitative analysis of the number of yellow (RFP+GFP+) and red (RFP+GFP−) puncta per cell as represented in (A). Data are represented as mean ± SD from three independent experiments. *p < 0.05; ***p < 0.001; ns, not significant (one-way ANOVA).

(D) Western blot analysis of samples from (A). mRFP-GFP-LC3B and the endogenous LC3B were examined. The LC3B-II/LC3B-I ratio was labeled to indicate the transition of LC3B under different transfection conditions. See also Figure S4.
Figure 5. BAG6 Interacts with LC3B via a LIR Motif

(A) Immunoprecipitation analysis of the interactions between FLAG-BAG6(1-991aa) and GFP-LC3B. Cells were transfected with GFP-LC3B along with the control or FLAG-BAG6(1-991aa) and FLAG pulldown was performed with anti-FLAG beads. Samples were analyzed by immunoblotting with the indicated antibodies.

(B) Endogenous BAG6 interacts with LC3B in HEK293FT cells. Immunoprecipitation using either IgG or anti-BAG6 was performed. The co-immunoprecipitated LC3B was immunoblotted with an anti-LC3B antibody.

(C) The domain structure of BAG6 and the LIR motifs at the N terminus of BAG6. A series of BAG6 truncations were generated based on the domain information.

(D) BAG6(1-266aa) is responsible for the interaction with GFP-LC3B. The different BAG6 truncations in (C), together with GFP-LC3B, were transfected into HEK293FT cells. FLAG pulldown was performed with anti-FLAG beads. Samples were analyzed by immunoblotting with the indicated antibodies.

(E) BAG6 LIR1 mutants decreases the binding affinity to LC3B. HEK293FT cells were transfected with GFP-LC3B, together with FLAG-tagged wild-type BAG6(1-266aa), LIR1Y132A/V135A, LIR1Y132A/V133A/M134A/V135A (abbreviated as LIR1AAAA), or YVMV deletion (designated as Δ132-135). FLAG pulldown was performed with anti-FLAG beads. Samples were analyzed by immunoblotting with the indicated antibodies. The number below the blots indicates relative intensity of the proteins.

(F and G) The addition of increasing concentration of LC3B-I (F) or Pro-LC3B (G) (from 0 to 50 μM) to BAG6 LIR1 peptide (1 μM) leads to a concentration-dependent increase in anisotropy value. The Kd value was calculated using nonlinear curve fitting. BAG6 LIR1 mutant and the random peptide were used as negative controls. Data are mean ± SD from three independent experiments.

See also Figure S5.
slightly reduced the binding affinity to LC3B. The four amino acid substitution mutant FLAG-BAG6(1-266aa) LIR1Y132A/V135A and the deletion mutant FLAG-BAG6(1-266aa)Δ132-135 greatly decreased the amount of bound LC3B (Figure 5E). In contrast, mutation in LIR2/LIR3/LIR4 motifs did not affect the interaction of BAG6 and LC3B (Figure 5S). Intriguingly, our co-immunoprecipitation studies also suggest that BAG6 might prefer to bind to LC3B-I rather than to LC3B-II (Figures 5A–5E). As shown in Figure 5S, although CQ treatment resulted in more LC3B-II, there was more LC3B-I than LC3B-II co-precipitated by BAG6. To gain more insights into the binding site between BAG6 LIR1 motif and LC3B, we synthesized an FITC-labeled 20-aa peptide encompassing the BAG6 LIR1 motif (residues 124–143) and a mutant peptide bearing two amino acid substitutions in LIR1 (LIR1Y132A/V135A). We also synthesized a random peptide as a negative control. We then incubated purified LC3B-I at different concentrations (up to 50 μM) with these peptides and measured fluorescence anisotropy. Indeed, addition of LC3B-I increased the anisotropy of wild-type BAG6 LIR-1 peptide but not that of BAG6 mutant peptide or the random control peptide (Figure 5F). The dissociation constant (Kd) for the binding of BAG6 LIR1 to LC3B-I was 22.23 ± 2.45 μM. These results indicate that BAG6 can directly bind to LC3B in a LIR1-dependent manner. We also introduced Pro-LC3B, the unprocessed form of LC3B, in this assay, because the protein sequence of Pro-LC3B is very similar to that of LC3B-I. Similar result was obtained with Pro-LC3B, with Kd = 17.67 ± 2.37 μM (Figure 5G). These results suggest that the cleaved BAG6 may bind to Pro-LC3B/LC3B-I and sequester them from being further lipidated.

To explore how BAG6 LIR1 affects autophagy, we transfected the BAG6 LIR1 mutants into HeLa cells stably expressing mRFP-GFP-LC3B. BAG6Y132A/V135A and BAG6D995A/Y132A/V135A stimulated autophagy in a similar extent as their wild-type LIR1 counterparts (Figures 4A, 4C and 4D). In contrast, overexpression of BAG6(1-991aa)Y132A/V135A did not suppress autophagy as that of BAG6(1-991aa) (Figures 4A, 4C and 4D), suggesting the involvement of LIR1 in autophagic regulation. Consistent with these results, the mutants in BAG6 knockout cells exhibited similar phenotypes to their counterparts in the wild-type background (Figures S4A and S4B), verifying the specific role of BAG6 in autophagy. The number of WIPI2-positive dots in BAG6(1-991aa)Y132A/V135A-overexpressing cells was also increased compared with that in BAG6(1-991aa)-overexpressing cells (Figures S4C and S4D). Collectively, these results suggest that the cleaved N-terminal BAG6 regulates autophagy through its LIR1 motif.

**BAG6 Knockdown Promotes Autophagosome Formation**

To further verify that BAG6 regulates autophagy, we performed an image-based analysis to see how depletion of BAG6 would affect autophagy. Using the mRFP-GFP-LC3B stable cell line, knockdown of BAG6 led to a noticeable increase of both yellow and red LC3B puncta number under both normal and starvation conditions (Figures 6A and 6B). Immunoblotting result further verified the transition of LC3B-I to LC3B-II under BAG6 knockdown condition (Figure 6C). These results verified that BAG6 inhibits the autophagic flux.

Next, we examined whether autophagy attenuation caused by AMFR overexpression or USP13 knockdown could be suppressed by BAG6 knockdown. Both AMFR overexpression and USP13 knockdown resulted in a decrease in the number of yellow LC3B puncta under normal and starvation conditions (Figures S6A and S6B). Interestingly, BAG6 knockdown completely released the autophagy inhibition caused by AMFR overexpression or USP13 knockdown, and the phenotype was similar to that of BAG6 single knockdown, as indicated by the increased number of yellow and red LC3 puncta in these conditions (Figures S6A and S6B). These data suggest that BAG6 functions downstream of AMFR and USP13 in autophagic pathway.

**NMR Identifies the Direct Interaction between BAG6 LIR1 Motif and LC3B**

To elucidate how BAG6 interacts with LC3B and the functional consequence of this interaction, we used NMR to further characterize the interaction between BAG6 LIR1 peptide and LC3B. BAG6 LIR1 and mutant peptides were synthesized and HPLC verified (Figures S7A and S7B). The 1H-15N HSQC spectra were then collected. In the absence of BAG6 LIR1, the 1H-15N HSQC spectra of LC3B-I and Pro-LC3B were different (Figure 7A). Although the concentrations of the two samples were the same, the spectra of Pro-LC3B showed weaker peak intensities. At the same time, there were differences in the peak positions. The results indicate that the two proteins adopted different structures despite significant sequence homology. The weaker spectrum profile for Pro-LC3B also suggests that Pro-LC3B may exist in several different conformational states, with only one major state showing peaks.
We then titrated different amounts of BAG6 LIR1 or BAG6 LIR1 mutant peptide into the NMR samples at the ratio of 0, 0.5, 1, 1.5, and 2 and measured the $^1$H-$^{15}$N HSQC spectra. The spectra of Pro-LC3B changed significantly upon addition of BAG6 LIR1, but the majority of the LC3B-I peaks remained unaffected by BAG6 LIR1. Intriguingly, the HSQC spectrum of Pro-LC3B in the presence of BAG6 LIR1 at 0.5 ratio was almost identical to that of LC3B-I in the presence of BAG6 LIR1 at 0.5 ratio (Figure 7B). These results suggest that Pro-LC3B undergoes a dramatic conformational switch upon interaction with the BAG6 LIR1 and the structure of Pro-LC3B bound by BAG6 LIR1 is similar to LC3B-I. In addition, upon addition of BAG6 LIR1, some peaks in the spectra of LC3B-I (Figure 7C) and Pro-LC3B (Figure 7D) were either shifted or eliminated. For example, the insets in Figures 7C and 7D showed an expanded area where two peaks exhibited significantly broadening and finally disappeared when excess of BAG6 LIR1 was present (arrow), whereas the other two peaks only showed negligible changes upon titration of BAG6 LIR1. The disappearing residue peaks were further identified, including 4E, 12T, 19D, 33V, 51K, 52F, 53L, 81L, 85G, 96S, and 108F (Figure S7C). Most of these residues were in or near the hydrophobic pockets on the surface of LC3B, implying the location of LC3B interacting with BAG6.

Figure 6. Depletion of BAG6 Induces Autophagy
(A) Depletion of BAG6 induces aggregation of LC3B-positive puncta in cells under both normal and starvation conditions. HeLa cells stably expressing mRFP-GFP-LC3B were transfected with control shRNA, BAG6 shRNA-1, or shRNA-2 and treated with normal medium or EBSS for 4 h. BAG6 is in cyan (Alexa 633) and nuclei are in blue. The arrows indicate the cells with BAG6 knockdown. Scale bar, 5 μm.
(B) Quantitative analysis of the number of yellow (RFP+GFP+) and red (RFP+GFP-) puncta per cell as represented in (A). Data are represented as mean ± SD from three independent experiments. *p < 0.05; **p < 0.01, ***p < 0.001 (one-way ANOVA).
(C) Western blot analysis of the samples as represented in (A). The knockdown efficiency of BAG6 was shown. mRFP-GFP-LC3B and the endogenous LC3B were also examined. The LC3B-II/LC3B-I ratio was labeled to indicate the transition of LC3B in BAG6 knockdown cells.
See also Figure S6.
Disappearance of residue peaks during the titration suggests an intermediate timescale of exchange for those residues between free and bound states. To exclude possible artifacts in the stepwise titration, solution mixtures with 0.2 mM Pro-LC3B and 0.4 mM BAG6 LIR1 were directly prepared without the stepwise titration and their HSQC spectra were also acquired. The spectra were identical to those acquired for the titration experiments (Figure S7D), indicating there were no artifacts introduced in the stepwise titration.

The interaction of BAG6 LIR1 mutant with LC3B-I or Pro-LC3B was also studied using a similar NMR titration strategy. Unlike BAG6 LIR1, addition of BAG6 mutant to LC3B-I (Figure S7E) or Pro-LC3B (Figure S7F)
solution caused no significant spectra change with only minor chemical shift changes for a few residues. The lack of significant disturbance in spectra indicated that this BAG6 LIR1 mutant did not interact with LC3B-I or Pro-LC3B strongly and thus confirmed that the mutations introduced into BAG6 LIR1 did inhibit its interaction with LC3B-I or Pro-LC3B. Collectively, our NMR results suggest that BAG6 LIR1 can hold Pro-LC3B or LC3B-I in a very stable state and could possibly shield LC3B to be further recruited to the autophagosome.

**DISCUSSION**

Here we showed that AMFR and USP13, previously identified E3 and DUB in the ERAD pathway, also regulate caspase activity and BAG6 cleavage to balance apoptosis and autophagy. By a powerful NEDDylation capturing system, we identified TXN as a common substrate of USP13 and AMFR and showed that TXN ubiquitination results in CASP3 activation and BAG6 cleavage. We have further discovered that BAG6 can bind to LC3B through its LIR1 motif to directly inhibit autophagosome formation. Therefore, we propose the model that BAG6 is a regulator that balances apoptosis and autophagy (Figure 8).

BAG6 has a “holdase” activity, which shields exposed hydrophobic segments of misfolded polypeptide to prevent aggregation. Given this, we questioned whether BAG6 holds autophagy-related proteins to regulate autophagy. Since we have detected BAG6 preferentially binding to LC3B-I in cells, one possibility is that BAG6 holds LC3B-I to prevent it from forming autophagosome. By sequence analysis, we found that BAG6 has a LIR1 (132–135aa) motif, which may specifically interact with LC3B. Our in vitro anisotropy measurement verified that BAG6 binds to LC3B-I as well as the newly synthesized Pro-LC3B with high affinity (Figures 5F and 5G). Owing to the technical difficulty to synthesize LC3B-II in vitro, we were not able to test whether BAG6 has lower affinity for LC3B-II. In line with this, NMR studies also showed that BAG6 LIR1 could bind to and stabilize LC3B-I structure. We found that the HSQC spectra of LC3-I with BAG6 LIR at the ratio of 0.5 remained unchanged for 10 days, whereas the HSQC spectra of LC3-I showed changes only after 2 or 3 days. Interestingly, the NMR titration experiments further showed that Pro-LC3B had two different conformational states. With the addition of BAG6 LIR1, the conformational state of Pro-LC3B changed to a state similar to LC3B-I, suggesting that BAG6 binds to Pro-LC3B to stabilize the structure similar to LC3B-I (Figures 7A and 7B). Given that endogenous full-length BAG6 shuttles between nucleus and cytosol, we propose that full-length BAG6 interacts with a limited amount of Pro-LC3B/LC3B-I, thus autophagy would not be affected. Although BAG6 gets cleaved, it remains in the cytosol and therefore holds excessive amount of Pro-LC3B/LC3B-I and prevents them to autophagosomes. In support of this hypothesis, the LIR1 mutant BAG6(1-991aa)Y132A/V135A abolished its ability to inhibit autophagy (Figures 4A, 4C, and 4D). In summary, our data showed that BAG6 subcellular localization and its LIR1 are essential for its

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**Figure 8. A Simple Model of BAG6 Regulation of Cellular Homeostasis between Autophagy and Apoptosis**

Under normal condition, BAG6 shuttles between nucleus and cytosol to keep cellular homeostasis. Under ER stress condition such as AMFR overexpression or USP13 knockdown, BAG6 is cleaved by CASP3, which is activated by ubiquitinated TXN. The cleaved BAG6 is localized in the cytosol and holds LC3B to inhibit autophagy. Apoptosis is induced.
regulation in autophagy. It is also worth pointing out that it does not rule out the possibility that BAG6 can also regulate autophagy through interacting with other autophagy-related proteins.

Apoptosis and autophagy are two interconnected pathways that determine cell fate between death and survival in response to cell stress (Song et al., 2017). Common upstream signaling of these two pathways has been found (Song et al., 2017; Tsapras and Nezis, 2017). Upon ER stress, autophagy is activated preceding apoptosis to protect cells by degrading protein aggregations. Induced autophagy can block apoptosis by inhibiting the activation of caspases to reduce cellular injury, whereas autophagy can also induce apoptosis when the duration or degree of ER stress reaches the limit of cellular adaptive mechanisms. During the apoptosis stage, activated caspases can cleave autophagy-related proteins, including ATG5, BECN1 (beclin 1), and ATG4D, to inactivate autophagy (Doherty and Baehrecke, 2018; Song et al., 2017; Tsapras and Nezis, 2017). Our data showed that USP13 knockdown inhibited autophagy, which is consistent with the previous report that USP13 promotes autophagy through deubiquitinating BECN1 to stabilize the protein (Liu et al., 2011). Interestingly, BECN1 has been reported to regulate the balance between apoptosis and autophagy (Wirawan et al., 2010; Zhu et al., 2010). BECN1 is a key component of the PtdIns3K complex, which is essential for initiation of autophagosome formation. At the onset of apoptosis, BECN1 is cleaved by caspases, and the cleaved fragments of BECN1 abolish its autophagy-inducing capacity. Instead, the C terminus of cleaved BECN1 is predominantly recruited to mitochondria to induce the release of pro-apoptotic factors (Wirawan et al., 2010). This is very similar to BAG6 in our case, for the N terminus of BAG6 holds Pro-LC3B/LC3B-I and inhibits autophagy, whereas the C terminus of BAG6 was previously proven to induce apoptosis (Wu et al., 2004).

The difficulty of identification of specific E3 substrates is the weak transient interaction between E3 and the substrates. In addition, rapid degradation of ubiquitinated substrates by proteasome also increased the difficulty. The NEDDylator system is a powerful tool to identify specific E3 substrates by modifying substrates with NEDD8 (Zhuang et al., 2013). TXN, identified by the NEDDylation system, turned out to be a common substrate of AMFR and USP13. TXN participates in the apoptotic pathway by catalyzing denitrosylation of caspases (Benhar et al., 2008; Mitchell et al., 2007; Rossig et al., 1999). S-nitrosylation is a post-translational modification, which mediates transduction of myriad cellular signals by adding nitric oxide (NO) to cysteine residues of target proteins (Sengupta and Holmgren, 2013). Previous studies demonstrated that caspases residing in mitochondria are consistently S-nitrosylated to inhibit their activity (Mannick et al., 2001). After an apoptotic stimulus, caspases are denitrosylated by TXN and rapidly activate the apoptotic pathway. Here, we found that knockdown of USP13 increased the ubiquitination of TXN, as well as the activation of CASP3, suggesting that ubiquitination of TXN is related to the activation of CASP3. Based on these results, we propose that ubiquitination of TXN is involved in regulating TXN activity. Under normal condition, BAG6 shuttles between nucleus and cytosol to keep cellular homeostasis. Knockdown of USP13 or overexpression of AMFR leads to the accumulation of ubiquitinated TXN and increased the activity of TXN. As a result, CASP3 was rapidly activated to cleave BAG6, which then holds LC3B and promotes cell apoptosis (Figure 8).

In summary, we propose a mechanism of ERAD machinery proteins, AMFR, USP13, and BAG6, in maintaining cellular homeostasis by regulating apoptosis and autophagy. The key protein BAG6 functions as a switcher to decide cell fate to either survive or die. These findings broaden our knowledge that BAG6 not only functions to help degrade client proteins but also plays a key role in balancing cellular homeostasis. Designing specific regulators of BAG6 could be a potential therapeutic strategy for its associated diseases.

Limitations of the Study
Our study shows that BAG6 preferentially binds to LC3-I rather than LC3-II in vivo, and future work is needed to validate that BAG6 inhibits LC3 lipidation in vitro. The new function of BAG6 LIR1 domain in autophagy needs to be further developed.

Resource Availability
Lead Contact
Further information should be directed to and will be fulfilled by the Lead Contact, Yanfen Liu (liuyf@shanghaitech.edu.cn).
Materials Availability
Materials are available upon request from Dr. Yanfen Liu.

Data and Code Availability
This study did not generate/analyze datasets.

METHODS
All methods can be found in the accompanying Transparent Methods supplemental file.

SUPPLEMENTAL INFORMATION
Supplemental Information can be found online at https://doi.org/10.1016/j.isci.2020.101708.

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AUTHOR CONTRIBUTIONS
Y.L., J. Lu, Y.C., and X.D. designed the experiments and wrote the manuscript; M.Z., Y.L., and Y.C. designed the experiments of the NEDDylator system; X.D. and J. Lu designed the NMR experiments and analysis of NMR data; Y.C., X.D., Y.K., J. Liu, T.Z., C.Y., Z.W., W.S., J. Lu, and Y.L. performed the experiments.

DECLARATION OF INTERESTS
The authors declare no competing interests.

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Supplemental Information

The Chaperone BAG6 Regulates Cellular Homeostasis between Autophagy and Apoptosis by Holding LC3B

Yuanyuan Chu, Xingqi Dong, Yingjin Kang, Jingnan Liu, Tao Zhang, Cuiwei Yang, Zhangshun Wang, Wangchen Shen, Huanhuan Huo, Min Zhuang, Junxia Lu, and Yanfen Liu
Figure S2

A

-  +  GST-AMFRΔTM

| Input |  +  GST-AMFRΔTM |
|-------|-----------------|
| + E1/E2/Flag-Ub | NEDD8 E1/E2/ HB-NEDD8 |

IB:Flag

IB:NEDD8

Flag-ub

AMFR

B

-  +  GST-AMFRΔTM

| Input |  +  GST-AMFRΔTM |
|-------|-----------------|
| + E1/E2/Flag-Ub | NEDD8 E1/E2/ HB-NEDD8 |

IB:F2

IB:NEDD8

Flag-ub

AMFR

FAF2~NEDD8

FAF2
Figure S4

A (BAG6+ cell line)  

|          | Control | BAG6 | BAG6(1-991aa) | BAG6**D995A** | BAG6**Y132A/V135A** | BAG6(1-991aa)**Y132A/V135A** | BAG6**D995A/Y132A/V135A** | BAG6 |
|----------|---------|------|---------------|--------------|-------------------|-----------------------------|--------------------------|------|

Flag RFP  

GFP  

Starvation  

Merge  


B  

|          | Control | BAG6 | BAG6(1-991aa) | BAG6**D995A** | BAG6**Y132A/V135A** | BAG6(1-991aa)**Y132A/V135A** | BAG6**D995A/Y132A/V135A** | BAG6 |
|----------|---------|------|---------------|--------------|-------------------|-----------------------------|--------------------------|------|

Flag (BAG6)  

Starvation  

Merge  


C (mRFP-GFP-LC3B stable cell line)  

|          | Control | BAG6 | BAG6(1-991aa) | BAG6**D995A** | BAG6**Y132A/V135A** | BAG6(1-991aa)**Y132A/V135A** | BAG6**D995A/Y132A/V135A** | BAG6 |
|----------|---------|------|---------------|--------------|-------------------|-----------------------------|--------------------------|------|

Flag (BAG6)  

Starvation  

Merge  


D  

|          | Control | BAG6 | BAG6(1-991aa) | BAG6**D995A** | BAG6**Y132A/V135A** | BAG6(1-991aa)**Y132A/V135A** | BAG6**D995A/Y132A/V135A** | BAG6 |
|----------|---------|------|---------------|--------------|-------------------|-----------------------------|--------------------------|------|

GFP Merge  

BAG6  

mRFP-GFP-LC3B stable cell line  

Flag  

Control  

The number of puncta per cell  

The number of WIPI2* puncta per cell  

---

*** ****** ns

*** ****** ns

ns  

*** ****** ns

ns  

ns

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ns  

ns

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ns

ns

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ns

ns

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ns

ns

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ns

ns

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ns

ns

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ns

ns

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ns

ns

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ns

ns
### Figure S5

#### A

| Input (16%) | IP: Flag |   |
|-------------|----------|---|
| - + - - -   | - + - - - | Flag-BAG6(1-387aa) |
| - + + - -   | - + + - - | Flag-BAG6(1-387aa) LIR2**** |
| - + - + -   | - + - + - | Flag-BAG6(1-387aa) LIR3**** |
| - + - - +   | - + - - + | Flag-BAG6(1-387aa) LIR4**** |
| + + + + +   | + + + + + | GFP-LC3B |

#### B

| Input (8%) | IP: IgG | IP: BAG6 |
|-----------|--------|---------|
| Control   | Starvation | CQ     |
| Control   | Starvation | CQ     |

| kD | BAG6 | LC3B-I | LC3B-II |
|----|------|--------|--------|
| 150|      |        |        |
| 15 |      |        |        |

GFP-LC3B-I
GFP-LC3B-II
Flag
Figure S6

A mRFP-GFP-LC3B stable cell line

| Condition   | GFP | RFP | Merge |
|-------------|-----|-----|-------|
| Control     | ![Control GFP](image1) | ![Control RFP](image2) | ![Control Merge](image3) |
| shBAG6-1    | ![shBAG6 GFP](image4) | ![shBAG6 RFP](image5) | ![shBAG6 Merge](image6) |
| AMFR        | ![AMFR GFP](image7) | ![AMFR RFP](image8) | ![AMFR Merge](image9) |
| AMFR + shBAG6-1 | ![AMFR+shBAG6 GFP](image10) | ![AMFR+shBAG6 RFP](image11) | ![AMFR+shBAG6 Merge](image12) |
| shUSP13-1   | ![shUSP13-1 GFP](image13) | ![shUSP13-1 RFP](image14) | ![shUSP13-1 Merge](image15) |
| shUSP13-1+  | ![shUSP13-1+ GFP](image16) | ![shUSP13-1+ RFP](image17) | ![shUSP13-1+ Merge](image18) |

B The number of puncta per cell

| Condition   | Red puncta | Yellow puncta |
|-------------|------------|---------------|
| Starvation  | *** ns     | *** ns        |
| shBAG6 -    | ***        | ns            |
| AMFR -      | *          | ns            |
| shUSP13-1   | +          | *** ns        |

* indicates statistical significance, ns indicates non-significant difference.
Figure S7

A

Fragmentor Voltage  | Collision Energy | Ionization Mode |
---------------------|------------------|-----------------|
140                  | 0                | ESI             |

*EBI Scan (0.437 min) Fragg=140.0V Bag66_wildtype.d

Counts (%) vs. Mass-to-Charge (m/z)

B

Fragmentor Voltage  | Collision Energy | Ionization Mode |
---------------------|------------------|-----------------|
140                  | 0                | ESI             |

*EBI Scan (0.447 min) Fragg=140.0V Bag66_mutant.d

Counts (%) vs. Mass-to-Charge (m/z)

C

Blue: BAG6/Pro-LC3B ratio=0
Red: ratio=0.5

D

Green: titration
Red: premixture

E

Black: BAG6/Pro-LC3B ratio=0
Red: ratio=0.5
Blue: ratio=1
Green: ratio=1.5
Maroon: ratio=2

F

Black: BAG6/Pro-LC3B ratio=0
Red: ratio=0.5
Blue: ratio=1
Green: ratio=1.5
Maroon: ratio=2
Supplemental Figure Legends

**Figure S1. Depletion of USP13 induces BAG6 cleavage, which is blocked by caspase depletion, Related to Figure 1.**

(A) The membrane fraction BAG6 was cleaved in USP13 knockout cells which was induced by MG132 and blocked by Z-VAD-FMK. Samples were prepared using the same procedure as in Figure 1E.

(B and C) As in Figure 1F, except different caspase shRNAs were co-transfected with USP13 shRNAs.

(D) Specific cleavage of BAG6 by recombinant purified CASP2/3/7 in vitro. Recombinant BAG6 was incubated with different concentrations of caspase enzymes at 37°C for 1 h.

(E) BAG6<sup>D995A</sup> was not cleaved by recombinant purified CASP2/3/7 in vitro. As in (C), expect using the caspase cleavage site mutant BAG6<sup>D995A</sup> as the substrate.

**Figure S2. NEDDylation efficiency of the three NEDDylators, Related to Figure 2.**

(A) Wild type AMFR exclusively mediates ubiquitination but not NEDDylation on substrates. E1-GST, His-UBE2G2 (E2), Flag-ubiquitin, and GST-AMFRΔTM were incubated for 1 h at 37°C (Left). NEDD8 E1, NEDD8 E2 UBE2M, HB-NEDD8, and GST-AMFRΔTM were incubated for 1 h at 37°C (Right). Reaction samples were stopped by adding sample buffer and analyzed by immunoblotting with Flag antibody.

(B) NEDDylator<sup>CUE+G2BR</sup> and NEDDylator<sup>CUE</sup>, but not NEDDylator<sup>ΔRING</sup>, can efficiently transfer NEDD8 to a known AMFR substrate FAF2. AMFR knockout cells were transfected with NEDDylator<sup>CUE+G2BR</sup>, NEDDylator<sup>CUE</sup>, or NEDDylator<sup>ΔRING</sup>, along with HB-NEDD8. Cells were treated with biotin (5 μM) for 48 h. Samples were immunoprecipitated by streptavidin beads and analyzed by FAF2 antibody.

**Figure S3. TXN interacts with USP13, AMFR and caspases, Related to Figure 3.**

(A and B) Immunoprecipitation analysis of the interactions between Flag-TXN and AMFR (A), or Flag-TXN and GFP-USP13 (B). Cells were transfected with AMFR (A) or GFP-USP13 (B), together with control vector or Flag-TXN. Immunoprecipitation using either IgG or anti-Flag was performed. The co-immunoprecipitated AMFR or USP13 was immunoblotted with an anti-AMFR (A) or anti-GFP antibody (B).

(C) GST-AMFRΔTM interacts with His-MYC-TXN. His-MYC-TXN purified from E. coli was incubated with GST or GST-AMFRΔTM for pulldown analysis. Samples were immunoblotted with an anti-MYC antibody.
(D) Immunoprecipitation analysis of the interactions between Flag-TXN and CASP2/3/7 under TNF-α treatment. Cells were transfected with control vector or Flag-TXN, and treated with TNF-α (10 ng/ml) for 24 h. Flag pulldown was performed with anti-Flag beads. Samples were analyzed by immunoblotting with the indicated antibodies.

(E) Schematic of the biotin switch labeling assay. Three major steps are involved: 1) free thiol-blocking; 2) selective reduction of S-nitrosylation (SNO) sites and labeling of resultant free thiols; 3) detection of biotin-labeled thiols.

(F) S-nitrosylated CASP3 (SNO-CASP3) is decreased in TXN-overexpressing cells. HEK293FT cells were transfected with control vector or Flag-TXN. 24 h after transfection, cells were prepared following the operation instruction of Biotin Switch Assay Kit (S-nitrosylation). After immunoprecipitation by streptavidin beads, the samples were analyzed by immunoblotting with the indicated antibodies. SNO-CASP3 level was decreased in TXN-overexpressing cells. The cleaved CASP3 was increased in TXN-overexpressing cells in the input fraction.

Figure S4. Cleaved BAG6 suppresses autophagy through its LIR1 motif, Related to Figure 4.
(A) Endogenous WIPI2 signal in HeLa cells transfected with Flag-tagged BAG6 or its mutants. Cells were treated with EBSS for 4 h. WIPI2 is in green, Flag signal is in red, and nuclei are in blue. Scale bar, 5 μm.

(B) Quantitative analysis of the number of WIPI2 dots per cell as represented in (A). Data are represented as mean ± SD from three independent experiments. ***p < 0.001; ns, not significant (one-way ANOVA).

(C) BAG6 knockout cells were transiently transfected with mRFP-GFP-LC3B, together with Flag-tagged BAG6 or its mutants. Cells were treated with EBSS for 4 h, and analyzed for LC3B puncta. Flag is in cyan and nuclei are in blue. Scale bar, 5 μm.

(D) Quantitative analysis of the number of yellow (RFP+GFP+) and red (RFP+GFP−) puncta per cell as represented in (C). Data are represented as mean ± SD from three independent experiments. *p < 0.05; ***p < 0.001; ns, not significant (one-way ANOVA).

Figure S5. BAG6 interaction with LC3B is not dependent on LIR2/3/4 motifs, Related to Figure 5.
(A) BAG6 LIR2, LIR3, and LIR4 mutants do not affect the binding affinity to LC3B. Immunoprecipitation analysis of the interactions between Flag-BAG6(1-387aa) mutants and GFP-LC3B. HEK293FT cells were transfected with GFP-LC3B, together with BAG6(1-387aa)
or its LIR mutants: LIR2<sup>Y270A/V271A/E272A/V273A</sup> (abbreviated as LIR2<sup>AAAA</sup>), LIR3<sup>Y292A/E293A/V294A/L295A</sup> (abbreviated as LIR3<sup>AAAA</sup>), LIR4<sup>F331A/V332A/L334A</sup> (abbreviated as LIR4<sup>AAAA</sup>). Flag pulldown was performed with anti-Flag beads. Samples were analyzed by immunoblotting with the indicated antibodies.

(B) Immunoprecipitation analysis of the interaction between BAG6 and LC3B-I/LC3B-II in HEK293FT cells. Cells were grown under normal condition, treated with EBSS for 4 h, or treated with 20 μM CQ in complete medium for 4 h. Immunoprecipitation using either IgG or anti-BAG6 was performed. The co-immunoprecipitated LC3B was immunoblotted with an anti-LC3B antibody.

**Figure S6. Knockdown of BAG6 suppresses AMFR overexpression- or USP13 knockdown-induced autophagy attenuation, Related to Figure 6.**

(A) Autophagy attenuation caused by AMFR overexpression or USP13 knockdown is suppressed by BAG6 knockdown. HeLa cell stably expressing mRFP-GFP-LC3B were transfected with control vector, BAG6 shRNA-1, AMFR, (BAG6 shRNA-1 + AMFR), USP13 shRNA-1, or (BAG6 shRNA-1 + USP13 shRNA-1), and treated with normal medium or EBSS for 4 h. Nuclei are in blue. Scale bar, 5 μm.

(B) Quantitative analysis of the number of yellow (RFP⁺GFP⁺) and red (RFP⁺GFP⁻) puncta per cell as represented in (A). Data are represented as mean ± SD from three independent experiments. *p < 0.05; **p < 0.01; ns, not significant (one-way ANOVA).

**Figure S7. NMR analysis of BAG6 LIR1 mutant interacting with LC3B-I and Pro-LC3B, Related to Figure 7.**

(A and B) Identification of purified BAG6 peptide (A) or BAG6 LIR1 mutant by mass spectrometry.

(C) Identification of the disappearing residue peaks on LC3B during titration.

(D) Overlay ¹H-¹⁵N HSQC spectrum of BAG6 LIR1: Pro-LC3B ratio=2 using titration method (Green) and a directly prepared mixture (Red). The two spectra overlay perfectly.

(E) Overlay ¹H-¹⁵N HSQC spectra of BAG6 LIR1 mutant: LC3B-I titration at the ratio of 0 (Black), 0.5 (Red), 1 (Green), 1.5 (Blue), 2 (Maroon). Inset shows the spectrum change of four residue peaks based on BAG6 LIR1 mutant titration.

(F) Overlay ¹H-¹⁵N HSQC spectra of BAG6 LIR1 mutant: Pro-LC3B titration at the ratio of 0 (Black), 0.5 (Red), 1 (Green), 1.5 (Blue), 2 (Maroon). Inset shows the spectrum change of four
residue peaks based on BAG6 LIR1 mutant titration.
## Transparent Methods

### Key Resources Table

| Oligonucleotides and peptides used in this paper | Source |
|------------------------------------------------|--------|
| FITC-BAG6 LIR1 motif (VHDRNANSYVMGVTFNLPSD) | Zhengzhou Phtdpeptides Co.,Ltd |
| FITC-BAG6 LIR1 mutant motif (VHDRNANSMAGTFNLPSD) | Zhengzhou Phtdpeptides Co.,Ltd |
| FITC-random peptides (YGRKKRQRORYKEGVNYG) | Zhengzhou Phtdpeptides Co.,Ltd |
| Oligonucleotides (5'-3') | |
| AMFR sgRNA-1: GCTAACGGCCTCGCTGCAG | This paper |
| AMFR sgRNA-2: GCTACACCCACGGAATGCA | This paper |
| USP13 sgRNA-1: AAAGACATTTTCGAAAAAC | This paper |
| USP13 sgRNA-2: TACATTGCGCTTTTGGAA | This paper |
| shBAG6 targeting sequence:  
  1. ACCGGAATGCCCAACAGCTATGTCATGGTT  
  2. GCCATCCCCATACAGATCAAT | (Wang et al., 2011) |
| shUSP13 targeting sequence:  
  1. CCTGAATACCTTGGTAGTGCAGATAAAGAA  
  2. GGCATGTGTTAAGGCTTTTGTT | (Liu et al., 2014) |
| shCASP2 targeting sequence:  
  1. GGACATCATCACCTTGGAAAT  
  2. GCACTTCGAGAGGAGAAAGA | This paper |
| shCASP3 targeting sequence:  
  1. GGGAAATGGCATCTGGAA  
  2. CCTGAGATGGGTATGTATA | This paper |
| shCASP7 targeting sequence:  
  1. GCTTCGCCTGCATCCTCTTAA  
  2. GATGGTGTCACACCACAAAG | This paper |
| BAG6 Y132A/V135A mutagenic forward primer: GGAATGCCCAACAGCTGTCATGGCTGGAACC | This paper |
| BAG6 LIR1<sup>AAAA</sup> mutagenic forward primer: GGAATGCCCAACAGCTGTCATGGCTGGAACC | This paper |
| BAG6 Δ132-135 mutagenic forward primer: GGAATGCCCAACAGGGAACCTTCAATC | This paper |
| BAG6 LIR2<sup>AAAA</sup> mutagenic forward primer: ATCCTTCCCTGCGAGGCTGCGGCGCGCTC CAGGAGCTA | This paper |
Cell lines
HEK293FT (PTA-5077) and HeLa cells (CRM-CCL-2) from ATCC were cultured in DMEM (Thermo Fisher Scientific, 10566024) with 10% fetal bovine serum (Gemini, 900-108) and 100 U/ml penicillin G and 100 μg/ml streptomycin (Thermo Fisher Scientific, 15140148) at 37°C under 5% CO₂. HeLa cells stably expressing mRFP-GFP-LC3B were cultured in complete medium containing 300 ng/μl G418 (Invivogen, ant-gn-1). CRISPR/Cas9 technology was used to generate the USP13 and AMFR knockout cells. Target sequences are listed in Key Resource Table. Target sequences were inserted into pGL3-U6-2sgRNA plasmid. Cells were co-transfected with pST1374-N-NLS-Flag-linker-Cas9 and sgRNA plasmids. Isolated knockout clones were verified by western blot and sequencing.

Plasmids
pCDN3.0-AMFR, GST-AMFRΔTM, GFP-USP13 and Flag-BAG6 were described previously (Liu et al., 2014). For BAG6, USP13, CASP2, CASP3, and CASP7 shRNA knockdown, target sequences were cloned into pSUPER.neo. The empty vector was used as the negative control. Target sequences are listed in Key Resource Table. pST1374-N-NLS-Flag-linker-Cas9 plasmid (Addgene, 44758) and pGL3-U6-2sgRNA plasmid (Addgene, 115519) were provide by Professor Xingxu Huang (ShanghaiTech University, China). Construct for AMFR shRNA was provided by Professor Shenyun Fang (University of Maryland, Baltimore, MD). mRFP-GFP-LC3B plasmid was provided by Professor Tamotsu Yoshimori (Osaka University, Japan; Addgene, 21074). EGFP-LC3B was constructed by inserting EGFP-LC3B fragment into pcDNA3.0. The construct for expression of Flag-TXN was generated by cloning the coding DNA fragment into the pRK5-Flag vector. The construct for expression of His-MYC-TXN was generated by cloning the coding DNA fragment into the pET28a vector. The constructs for expression of the various Flag-BAG6 truncations were generated by individually introducing a stop codon at specific positions in Flag-BAG6 plasmid. The BAG6 LIR mutants were generated...
by PCR-based mutagenesis and the mutagenic primers are listed in Key Resource Table. All mutations were confirmed by DNA sequencing. The constructs for expression of His-Pro-LC3B (residues 1-125) and His-LC3B-I (residues 1-120) were generated by cloning the coding DNA fragments into the pET32a vector. The constructs for expression of His-BAG6 LIR1 motif (residues 124-143: VHDRNANSYVMGVTFNLPSD) and the mutant LIR1 Y132A/V135A (VHDRNANSAMGTFNLPSD) were generated by inserting commercially synthesized oligomer into the pET32a vector. Mammalian expression constructs for HB-NEDD8 and NEDD8 E2 UBE2M, and prokaryotic expression constructs for HB-NEDD8, NEDD8 E1, NEDD8 E2 UBE2M, His-CASP2, His-CASP3 and His-CASP7 were kindly provided by Dr. Zhuang Min (Zhuang et al., 2013). NEDDylator system plasmids were generated by fusing the different AMFR fragments with a NEDD8 E2 UBE2M.

Antibodies and reagents

Rabbit anti-USP13, rabbit anti-BAG6, rabbit anti-AMFR were described previously (Liu et al., 2014). Anti-GFP antibody was developed in rabbit using recombinant GFP protein as the immunogen. The antibody was affinity-purified on the immunizing protein immobilized on agarose. Other primary antibodies used were as follows: rabbit anti-Flag (Sigma-Aldrich, F7425); rabbit anti-CASP2 (Abcam, ab179520), rabbit anti-CASP3 (ABclonal, A2156); rabbit anti-Cleaved-CASP3 (Cell Signaling Technology, 9664); rabbit anti-CASP7 (Cell Signaling Technology, 9492); rabbit anti-TXN (Proteintech, 14999-1-AP); rabbit anti-LC3B (Sigma-Aldrich, L7543, for detecting the endogenous LC3B); mouse anti-LC3 (M186-3; for detecting the exogenous LC3B); mouse anti-ACTB (HRP-Direct) (MBL, PM053-7); mouse anti-TOMM20 (Santa Cruz Biotechnology, sc-17764); rabbit anti-Histone H2AX (Abgent, AP20703b-400); rabbit anti-HA (Cell Signaling Technology, 3724); rabbit anti-NEDD8 (Cell Signaling Technology, 2754); mouse anti-MYC (Cell Signaling Technology, 2276); mouse anti-Ubiquitin (Santa Cruz Biotechnology, sc-8017); The secondary antibodies goat anti-mouse IgG (H+L), HRP (111-035-146) and goat anti-rabbit IgG (H+L), HRP (111-035-144) were purchased from Jackson ImmunoResearch Inc. The secondary antibodies goat anti-mouse IgG (H+L), Alexa Fluor 568 (A-11031) and goat anti-rabbit IgG (H+L), Alexa Fluor 633 (A-21071) were purchased from Thermo Fisher Scientific.
Z-VAD-FMK was purchased from Santa Cruz (sc-311561). MG132 was purchased from Medchemexpress (HY-13259). Earle's Balanced Salt Solution (EBSS) was purchased from Sigma-Aldrich (E2888).

**Protein expression and purification**

HB-NEDD8, CASP2, CASP3 and CASP7 were purified from BL21 cells according to previously described method (Scheer et al., 2005; Zhuang et al., 2013). Flag-BAG6 were purified using Flag affinity chromatography procedure (Sigma-Aldrich). GST-AMFRΔTM were purified from *E.coli* by GST affinity chromatography (GE Healthcare) procedure according to the reference (Russell and Wilkinson, 2005). His-Pro-LC3B (residues 1-125) and His-LC3B-I (residues 1-120) were purified using Ni-NTA beads (GE Healthcare) and then the His tag was cleaved by incubating with TEV protease at 4°C for 48 h. Free His tag and His-TEV protease were removed by Ni-NTA beads. Gel filtration size exclusive chromatography with Superdex 75 column (GE Healthcare) was employed for further purification. To purify 15N-labeled proteins and 15N, 13C double labeled proteins, related constructs were transformed into BL21 cells. Cells were grown in LB media containing 100 μg/ml ampicillin and 34 μg/ml chloramphenicol at 37°C for 10 h. Then cells were transferred to 50 ml M9 media and cultured overnight. Next the bacterial fluid were diluted at 1:10 with M9 media and cultivated at 37°C to an OD$_{600nm}$ around 1.0. Cells were then harvested by centrifugation at 8000 × g for 10 min at 4°C and the precipitate was transferred to 15N-labeled or 15N, 13C double labeled M9 media and cultivated at 37°C to an OD$_{600nm}$ around 1.0. Expression of isotope labeled proteins were induced with 0.5 mM IPTG at 16°C overnight. Proteins were purified through Ni-NTA beads using the same procedure as describe above. For purification of His-BAG6 LIR1 motif (residues 124-143) and its mutant His-BAG6 LIR1Y132AV135A, the polypeptides were further purified with HPLC after being cleaved by TEV protease.

**GST pulldown assay, immunoprecipitation, and immunoblotting**

GST-tagged proteins were incubated with GST beads for 1 h in PBS buffer, and the unbound proteins were removed by centrifugation at 1000 × g for 3 min. Next, GST-tagged protein bound beads were incubated with target proteins in NP40 lysis buffer (150 mM sodium chloride, 0.5% NP40 and 50 mM Tris–HCl pH 7.4) for 1 h. Then GST beads were washed three times with NP40 wash buffer (150 mM sodium chloride, 0.1% NP40 and 50 mM Tris–HCl pH 7.4). The samples were eluted by 1 × sample buffer and detected by western blot. For immunoprecipitation experiments, related plasmids were transfected into HEK293FT cells and were extracted by NP40 lysis buffer containing protease inhibitor cocktail. The soluble supernatant fractions were harvested by centrifugation at 17000 × g for 10 min and then used for immunoprecipitation with indicated antibodies. After incubating for 1 h, beads were washed three times with NP40 wash buffer, and then detected by western blot. For immunoprecipitation under denaturing condition, harvested cells were lysed in a buffer with 1% SDS and 5 mM DTT.
The samples were heated at 65°C for 10 min and diluted into 0.1% SDS and 0.5 mM DTT with NP40 lysis buffer. The soluble supernatant fractions were harvested and subjected to immunoprecipitation experiments as described above. Immunoblotting was then performed using polyvinylidene fluoride membrane (Bio-Rad, 1620177) and the indicated antibodies. ECL western blotting detection reagents (PerkinElmer, NEL105001EA) was used to detect the protein signal and the chemiluminescence bands were imaged under Amersham Imager 600 (GE Healthcare Life Sciences, USA). The bands were adjusted within the linear range, and quantified by ImageJ software (NIH).

**Immunofluorescence microscopy**

HeLa cells stably expressing mRFP-GFP-LC3B were seeded on small glass slides and transiently transfected with the indicated constructs. The small glass slides were collected 24 h after transfection and fixed with PBS containing 4% paraformaldehyde and 4% sucrose. After being washed for three times with PBS, the nuclei were labeled with DAPI (Sigma-Aldrich). Finally the samples were mounted with ClearMount™ Mounting solution (Invitrogen, USA). For the immunofluorescence stain studies, fixed cells were washed for four times with PBS, and then incubated with the indicated primary antibodies (1:500) for 1 h, and fluorescent dye-conjugated secondary antibodies (1:600) for 30 min. Images were acquired on Zeiss LSM800 or LSM880 microscope (Zeiss, Germany) with a 63 × 1.4 NA oil objective. Same acquisition parameters were used for a specific set of experiments.

**Cell viability assay**

Cell viability was measured by a cell proliferation assay kit (CellTiter 96® AQueous One Solution) (Promega, G3582), following the manufacturer’s instructions. HeLa cells transfected with control shRNA or USP13 shRNA were seeded in a 96-well plate. Cell death was measured by utilizing CellTiter 96® AQueous One Solution Reagent (contains a novel tetrazolium compound [3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium, inner salt; MTS] and an electron coupling reagent (phenazine ethosulfate; PES)) added to cells for 3 h at 37°C. Absorbance was recorded at 490 nm using a 96-well plate reader (MD SpectraMax i3, USA). Three independent experiments were performed.

**NEDDylator system**

The NEDDylator system were constructed according to reference (Zhuang et al., 2013). Briefly, NEDDylator were generated by fusion expressing substrate binding domain of AMFR with NEDD8 E2 UBE2M. The NEDDylator and HB-NEDD8 were co-transfected into AMFR knockout cells. After being cultured in DMEM containing 5 μM biotin for 48 h, cells were harvested and subject to immunoprecipitation with Ni-NTA column. The bound proteins were then eluted by
low pH buffer and subjected to another round of immunoprecipitation by Pierce™ Streptavidin Magnetic Beads (Thermo Fisher Scientific, 88816). Proteins were then analyzed by mass spectrometry.

**Biotin switch assay**

HEK293FT cells were transfected with the indicated constructs. 24 h after transfection, cells were prepared following the operation instruction of Biotin Switch Assay Kit (S-nitrosylation) (Abcam, ab236207) (also see Figure S3E). First, cells were washed three times with S-nitrosylation wash buffer. Then cells were lysed and blocked with Buffer A containing blocking reagent for 30 min at 4°C (Figure S3E, step 1). The clarified lysates were harvested by centrifugation at 17000 × g for 10 min and the proteins in supernatants were precipitated by ice-cold acetone for 1 h at -20°C. Next the protein pellets were incubated with Buffer B containing reducing and biotin labeling reagents for 1 h (Figure S3E, step 2). Then the proteins were precipitated again by ice-cold acetone for 1 h at -20°C, followed by resuspension with cold S-nitrosylation wash buffer. To detect the S-nitrosylated proteins, samples were immunoprecipitated by streptavidin beads and then immunoblotted with anti-CASP3 antibody to examine the level of S-nitrosylated CASP3 (SNO-CASP3) (Figure S3E, step 3).

**Anisotropy measurement**

FITC-labeled BAG6 LIR1 peptide, BAG6 LIR1 mutant peptide and random peptide were commercially synthesized. These peptides were pre-dissolved in HEPES buffer (150 mM sodium chloride and 10 mM HEPES, pH 7.4) containing 10% dimethyl sulfoxide (DMSO) as a concentrated stock solution. For anisotropy measurement, BAG6 peptides or random peptide were diluted to 1 μM with HEPES buffer and mixed with LC3B-I or Pro-LC3B range from 0 μM to 50 μM. Anisotropy of the mixture was detected using a HORIBA FluoroMax-4 (Japan) with the fluorescence polarization module. Excitation wavelength was 494 nm and emission was detected at 518 nm. Data analysis was conducted using OriginPro (Origin lab cooperation, USA) as well as Microsoft Excel Data Analysis package. The Kd value for the binding was calculated using nonlinear curve fitting in OriginPro.
NMR titration

NMR experiments were performed at 25°C on Agilent 800 MHz spectrometer and 600 MHz spectrometers equipped with a Cryoprobe. 0.2 mM 15N-labeled Pro-LC3B and 0.2 mM 13C,15N-labeled LC3B-I protein were prepared in 50 mM Tris-HCl pH 7.5, 100 mM sodium chloride buffer with 10% D2O. 4 mM wild type BAG6 LIR1 peptide or its mutant was pre-dissolved in 50 μl dimethyl sulfoxide (DMSO) as a concentrated stock solution. For titration experiments, the concentration ratio of BAG6 LIR1 peptide to LC3B-I or Pro-LC3B was controlled in the range from 0 to 2 by adding extra volume of BAG6 LIR1 peptide solution into the mixture. 2D 1H-15N HSQC experiments were carried out with 32 scans, acquiring 1562 (800 MHz spectrometer) or 1024 (600 MHz spectrometer) points in the direct dimension (1H dimension), and 128 points in the indirect dimension (15N dimension). DDS is used to determine 0 ppm of H dimension. All NMR spectra were processed using NMRPipe (Delaglio et al., 1995) and analyzed with Sparky (Goddard and Kneller).

Quantification and statistical analysis

Data are representative of at least three independent experiments. For quantitative analyses, values were expressed as mean ± SD, and error bars represented the standard deviations from counting of 80 cells in each group from three independent experiments. Statistical analyses were performed by using OriginPro and Microsoft Excel Data Analysis package. The significance among multiple groups were obtained using one-way ANOVA followed by Tukey’s multiple comparisons test. ns, not significant; *, p < 0.05; **, p < 0.01; ***, p < 0.001.
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