Degradation of macromolecules by lysosomes is a fundamental event for biological homeostasis. In mammals, three forms of autophagy, macroautophagy, microautophagy and chaperone-mediated autophagy (CMA), have been identified. Here, we showed a novel type of autophagy, in which RNA is taken up directly into lysosomes for degradation. This pathway, which we term “rNautophagy,” is ATP-dependent, and unlike CMA, is independent of HSPA8/Hsc70. LAMP2C, a lysosomal membrane protein, serves as a receptor for this pathway. The cytosolic tail of LAMP2C specifically binds to almost all total RNA derived from mouse brain. The cytosolic sequence of LAMP2C and its affinity for RNA are evolutionarily conserved from nematodes to humans. Our findings shed light on the mechanisms underlying RNA homeostasis in higher eukaryotes.

Discovery of a novel type of autophagy targeting RNA

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Abbreviations: CMA, chaperone-mediated autophagy; RBPs, RNA-binding proteins; LAMP, lysosome-associated membrane protein; LAMP, LAMP/CD68-like membrane protein; HSPA8, heat shock 70 kD protein 8; Hsc70, 70 kD heat shock cognate protein; RP, ribosomal protein; HNRNP, heterogeneous nuclear ribonucleoprotein; GAPDH, glyceraldehyde-3-phosphatedehydrogenase; RNase, ribonuclease; ATPase, adenosine triphosphatase; WT, wild type; KO, knockout; Hs, Homo sapiens; Ce, Caenorhabditis elegans; Dm, Drosophila melanogaster

Regulated degradation of cellular components by lysosomes is essential to maintain biological homeostasis. In mammals, three forms of autophagy, macroautophagy, microautophagy and chaperone-mediated autophagy (CMA), have been identified. Here, we showed a novel type of autophagy, in which RNA is taken up directly into lysosomes for degradation. This pathway, which we term “rNautophagy,” is ATP-dependent, and unlike CMA, is independent of HSPA8/Hsc70. LAMP2C, a lysosomal membrane protein, serves as a receptor for this pathway. The cytosolic tail of LAMP2C specifically binds to almost all total RNA derived from mouse brain. The cytosolic sequence of LAMP2C and its affinity for RNA are evolutionarily conserved from nematodes to humans. Our findings shed light on the mechanisms underlying RNA homeostasis in higher eukaryotes.
C-terminal sequences of both fly and nematode LAMP orthologs exhibited high affinity for RNA, similar to that of LAMP2C (Fig. 2E). Considering the relationship between LAMP2A and CMA, we hypothesized a novel autophagic pathway that directly imports RNA into lysosomes via LAMP2C. The standard method for monitoring direct uptake of molecules into lysosomes is a cell-free system using isolated lysosomes.12-14 Freshly isolated lysosomes were prepared from mouse brain (Fig. S4). We confirmed that the isolated lysosomes were intact and that CMA was working properly in a cell-free system (Fig. S5B). For monitoring uptake of RNA, the lysosomes were incubated with total RNA in the presence or absence of ATP and/or HSPA8. Subsequently, the lysosomes were precipitated by centrifugation, and the levels of RNA remaining in the supernatant were analyzed. In parallel, to assess the presence of RNA taken up inside the lysosomes, we degraded mouse brain lysate because, among various tissue types, Lamp2c mRNA showed the highest expression levels in the brain (Fig. 1D). Lamp2c was expressed in neurons while it was hardly detected in glial cells (Fig. S1). Mass spectrometry analysis of all protein bands detected in the pull-down assay revealed that LAMP2C peptide predominantly interacted with a wide range of RBPs (Fig. 1E; Table S1).

We next examined whether the interactions of RBPs with the cytosolic sequence of LAMP2C were mediated by RNA. The protein interactions of LAMP2C peptide were almost completely abolished by pretreatment of the lysate with protease-free RNase A in a pull-down assay (Fig. 2A). Substitutions of phenylalanines at the RNA-binding sites of HNRNPA1, which has a typical RNA-binding motif, completely abolished its interaction with LAMP2C peptide (Fig. 2B). These data suggest that the interactions of RBPs with the cytosolic tail of LAMP2C are indirect associations mediated by RNA.

We then investigated the interaction between RBAs and the cytosolic sequence of LAMP2C. We confirmed that the RNA specifically interacted with the LAMP2C peptide among LAMP2 constructs (Fig. 2C). Using purified total RNA derived from mouse brain, the cytosolic sequence of LAMP2C was shown to bind directly to a wide range of RNAs (Fig. 2D). The RNA was hardly detected in the flow-through after being pulled down with LAMP2C peptide (Fig. 2D), indicating that LAMP2C peptide bound to almost all RNAs. An endogenous interaction between LAMP2C and RNA was also observed (Fig. S2). In a pull-down assay using purified RNA, RNA partly interacted with the cytosolic sequence of LAMP2B, but not with that of LAMP2A (Fig. S3). Considering that LAMP2B peptide did not interact with RNA in mouse brain lysate, this interaction may be nonphysiological. Intriguingly, the C-terminal sequences of both fly and nematode LAMP orthologs exhibited high affinity for RNA, similar to that of LAMP2C (Fig. 2E).

Considering the relationship between LAMP2A and CMA, we hypothesized a novel autophagic pathway that directly imports RNA into lysosomes via LAMP2C. The standard method for monitoring direct uptake of molecules into lysosomes is a cell-free system using isolated lysosomes.12-14 Freshly isolated lysosomes were prepared from mouse brain (Fig. S4). We confirmed that the isolated lysosomes were intact and that CMA was working properly in a cell-free system (Fig. S5B). For monitoring uptake of RNA, the lysosomes were incubated with total RNA in the presence or absence of ATP and/or HSPA8. Subsequently, the lysosomes were precipitated by centrifugation, and the levels of RNA remaining in the supernatant were analyzed. In parallel, to assess the presence of RNA taken up inside the lysosomes, we degraded...
RNA outside of lysosomes by RNase A treatment after the incubation, and then analyzed the levels of RNase A-resistant RNA. As a result, the amount of RNA outside of lysosomes was dramatically reduced only in the presence of ATP (Fig. 3A). Levels of RNA were reduced over time in the presence of ATP, but remained unchanged in the absence of ATP (Fig. 3C). RNA resistant to exogenous RNase A was detected only in lysosomes incubated in the presence of ATP (Fig. 3B). These results indicate that RNA is taken up by lysosomes in a process that is dependent on ATP. Immunoelectron microscopy of the lysosomes incubated with ATP and RNA, or with ATP only, also showed the presence of RNA taken up in lysosomes (Fig. 3E; Fig. S6). Unlike CMA, HSPA8 had no effect on the uptake of RNA into isolated lysosomes (Fig. 3A and B).

We further investigated the degradation of RNA in lysosomes. Isolated lysosomes and RNA were incubated with or without ATP, and the total levels of RNA in the samples were analyzed. The total
levels of RNA were reduced in a manner that was dependent on ATP (Fig. 3D; Fig. S7), indicating ATP-dependent degradation of RNA by isolated lysosomes. Taken together, our results revealed an autophagic pathway that directly imports RNA into lysosomes for degradation. Hereinafter, we refer to this pathway as RNautophagy.

Finally, we assessed whether LAMP2C mediates RNA degradation by RNautophagy. Cells overexpressing LAMP2C exhibited a significant enhancement in the overall levels of RNA degradation, which was not observed in those overexpressing LAMP2A or LAMP2B (Fig. 4A; Fig. S8). Lysosomes isolated from cells overexpressing LAMP2C also showed elevated levels of RNautophagy (Fig. 4B). Additionally, the levels of RNautophagy were significantly reduced in lysosomes derived from LAMP2-deficient mouse brains (Fig. 4C). Moreover, a significant accumulation of total RNA was observed in the brains of LAMP2 knockout mice (Fig. 4D). Collectively, these results indicate that LAMP2C functions as a receptor for RNautophagy. The brains of the knockout mice, RBPs such as ribosomal protein S6 (RPS6), RPL8 and HNRNPA1 were upregulated (Fig. 4E). RBPs may be upregulated as a response to the accumulation of RNA, since RPL8 was not directly taken up by lysosomes (Fig. S5). Also, the upregulation of RBPs did not seem to be a result of CMA impairment, because representative substrates for CMA, GAPDH and SNCA/α-synuclein, were not accumulated in the brains of the knockout mice (Fig. S5). Consistent with this notion, RPL8, RPS6 and HNRNPA1 do not contain a KFERQ motif, a sequence required for targeting to CMA. The constancy of CMA substrates in the knockout brains may be due to compensation by other degradation systems.13

RNA degradation mechanisms in the cytosol independent of lysosomes have long been studied.19 On the other hand, an autophagic pathway selectively targeting RNA has not been elucidated, even though lysosomal RNases and exonucleases have long been recognized.20-22 We have shown here a novel, direct type of autophagy for RNA degradation, which we term RNautophagy. Our data suggest that RNautophagy can contribute to at least 10% to 20% of the total amount of RNA degradation in living cells (Fig. 4A). RNautophagy activity was not completely abolished in lysosomes derived from LAMP2-deficient mouse (Fig. 4C).

Figure 3. Uptake and degradation of RNA by isolated lysosomes. (A and B) Uptake of RNA by isolated lysosomes. Isolated lysosomes were incubated with purified total RNA (5 μg) in the presence or absence of ATP (energy regeneration system) and/or HSPA8. Levels of RNA remaining in solution outside of lysosomes (A) and levels of RNA resistant to exogenous RNase A (B) were analyzed (n = 3). (C) Lysosomes were incubated with 10 μg of RNA in the presence or absence of ATP. At the indicated times, the levels of RNA remaining outside of lysosomes were analyzed (n = 3). (D) Degradation of RNA by isolated lysosomes. Lysosomes and RNA were incubated with or without ATP. Total levels of RNA in the incubated samples were analyzed (n = 3). (E) Immunogold labeling of RNA in isolated lysosomes incubated with ATP and RNA. Immunogold labeling was performed using an anti-rRNA antibody followed by anti-mouse IgG coupled with 10 nm of gold particles. Gold particles were observed in the lysosomes.

Figure 4 (See opposite page). LAMP2C mediates lysosomal degradation of RNA. (A) RNA turnover in HeLa cells transfected with each LAMP2 isoform or empty vector (n = 3). (B and C) Uptake of RNA into lysosomes isolated from HeLa cells transfected with LAMP2C or empty vector (n = 3) (B). Uptake of RNA into lysosomes isolated from the brains of wild-type (WT) and LAMP2 knockout (KO) mice (n = 3) (C). Levels of RNA uptake were measured by subtracting the levels of RNA remaining in solution outside of lysosomes from the levels of input RNA. (D) Relative levels of total RNA in the brains of WT and LAMP2 KO mice (KO: n = 3, WT: n = 4). (E) Levels of RBPs and CMA substrate proteins in the brains of WT and LAMP2 KO mice were analyzed by immunoblotting (n = 3/group). (F) Model for the possible mechanism of RNautophagy. LAMP2C on the lysosomal membrane recognizes RNA, and the RNA is then imported into the lysosomal lumen in an ATP-dependent manner, followed by its degradation. The requirement for ATP in RNautophagy suggests an involvement of ATPases such as RNA helicases in the process on the lysosomal membrane.
Figure 4. For figure legend, see page 406.
Thus, we cannot rule out the possibility of LAMP2-independent pathway(s) in RNautophagy.

The RNA-binding capacities of the cytosolic sequences of LAMP family proteins are widely conserved in animals from nematodes to humans (Fig. 2). Together with the fact that a LAMP ortholog has not been found in yeast, RNautophagy likely contributes to RNA metabolism in higher eukaryotes. In the mouse, the expression levels of Lamp2c vary from tissue to tissue, showing notable levels in brain, eye, heart, liver, kidney and skeletal muscle, while the gene was scarcely expressed in other tissues (Fig. 1D). The different levels of Lamp2c expression among tissues could be related to requirements for RNA metabolism among tissues. RNautophagy may have an important role in RNA homeostasis in tissues, especially in those expressing LAMP2C at high levels. From this point of view, the high expression level of Lamp2c in neurons is reasonable, because neurons are rich in RNA. 25 RNautophagy may play an important role in the maintenance and quality control of neuronal function.

A possible model for the process of RNautophagy is shown in Figure 4F. First, a substrate RNA binds to the cytosolic tail of LAMP2C on the surface of the lysosomal membrane. This process may be regulated by other RNA-interacting proteins in the cytosol, because the cytosolic sequence of LAMP2C itself does not seem to have selectivity in its binding to RNA. Second, RNA is directly imported into the lysosomal lumen. LAMP2C and other proteins may be involved in this translocation process, which is ATP-dependent. RNA-dependent ATPases such as RNA helicases may operate in this process to unfold RNA, similar to the function of HSPA8 in CMA. Third, RNA taken up into lysosomes is degraded by lysosomal enzymes such as acid RNases and exonucleases.

We speculate that DNA could bind to the cytosolic tail of LAMP2C, because LAMP2C peptide interacted with DNA-binding proteins in addition to RBPs (Fig. 1C and E). The interaction between DNA and LAMP2C is currently the subject of ongoing research.

Because a lack of LAMP2 causes Danon disease, RNautophagy dysfunction may contribute to the pathogenesis of that disease. In addition, aberrant expression, function and accumulation of RNA have been considered as key factors in the pathogenesis of various diseases such as macular degeneration, myotonic dystrophies and some neurodegenerative disorders. 24 In the immune system, antiviral signaling is thought to be initiated intracellularly, within endosomes/lysosomes containing viral nucleic acid. 25 LAMP2 has been used as a macrophage marker. 26,27 Synthesis and expression of LAMP2 increase when macrophages are activated. 26,27 Given that Lamp2c is highly expressed in a macrophage cell line (Fig. S1), RNautophagy is possibly involved in the immune response. For instance, in certain types of cells, viral RNA could be taken up into lysosomes by RNautophagy and then be involved in antiviral responses via toll-like receptors on the endosomal/lysosomal membrane. The possible involvement of RNautophagy in the immune system and in lysosome- or RNA-related disorders warrants investigation.

**Materials and Methods**

Uptake and degradation of RNA by isolated lysosomes. Lysosomes were isolated from 10 to 12-week-old mouse brains or from ~1.4 × 10⁸ HeLa cells by density gradient centrifugation. Isolated lysosomes (25 to 50 µg of protein) were then incubated with 5 or 10 µg of purified total RNA at 37°C in 30 µl of 0.3 M sucrose containing 10 mM MOPS buffer (pH 8.0) with or without energy regeneration system (10 mM ATP, 10 mM MgCl₂, 2 mM phosphocreatine, and 50 µg/ml creatine phosphokinase). After the incubation, lysosomes were precipitated by centrifugation and the levels of RNA remaining in the supernatant were analyzed. To assess the presence of uptaken RNA inside lysosomes, RNA outside of lysosomes was degraded by 200 µg of RNase A after the incubation, and the levels of RNase A-resistant RNA were analyzed. Degradation of RNA by isolated lysosomes was investigated by analyzing the total levels of RNA in the samples.

Immunogold electron microscopy. Isolated lysosomes were fixed in 0.1% glutaraldehyde and 4% paraformaldehyde in PBS for 16 h at 4°C. Samples were then dehydrated in a series of water/ethanol mixtures to 100% ethanol, and embedded in LR White (Nisshin EM Co., Ltd., 3962). The embedded samples were sectioned at 100 nm, and collected on 400-mesh nickel grids. Immunogold labeling was performed using an anti-rRNA antibody followed by anti-mouse IgG coupled with 10 nm of gold particles, and samples were viewed using a Tecnai Spirit transmission electron microscope (FEI) at 80 kV.

**Statistical analyses.** For comparisons between two groups, the statistical significance of differences was determined by Student’s t test. For comparisons of more than two groups, Dunnett’s multiple comparison test was used. * and ** indicate p < 0.05 and p < 0.01, respectively.

Further details on materials and methods are provided in Supplemental Materials and Methods.

**Disclosure of Potential Conflicts of Interest**

No potential conflicts of interest were disclosed.

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**Supplemental Materials**

Supplemental materials may be found here: www.landebioscience.com/journals/autophagy/article/23002
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