Mice deficient in Epg5 exhibit selective neuronal vulnerability to degeneration

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Introduction

Autophagy is a degradation system involving the enclosure of cytosolic components in a double-membrane autophagosome and their subsequent delivery to the vacuole/lysosome for degradation (Nakatogawa et al., 2009). In yeast, autophagosomes directly fuse with the vacuole, whereas nascent autophagosomes in higher eukaryotes undergo maturation processes, including fusion with endosomes, before fusion with lysosomes to produce degradative autolysosomes (Longatti and Tooze, 2009). The more elaborate autophagic machinery in higher eukaryotes involves highly conserved Atg (autophagy related) proteins, factors involved in the endolysosomal pathway, and also metazoan-specific autophagy proteins (Longatti and Tooze, 2009; Tian et al., 2010).

Basal autophagy removes misfolded and toxic aggregate-prone proteins and damaged organelles (Levine and Kroemer, 2008). Neural-specific depletion of genes essential for autophagosome formation, including Atg5, Atg7, and Ei24, causes accumulation of p62 aggregates and ubiquitin-positive inclusions in neurons and glial cells. Epg5 knockdown also impaired endocytic trafficking. Our study establishes Epg5-deficient mice as a model for investigating the pathogenesis of ALS and indicates that dysfunction of the autophagic–endolysosomal system causes selective damage of neurons associated with neurodegenerative diseases.

The molecular mechanism underlying the selective vulnerability of certain neuronal populations associated with neurodegenerative diseases remains poorly understood. Basal autophagy is important for maintaining axonal homeostasis and preventing neurodegeneration. In this paper, we demonstrate that mice deficient in the metazoan-specific autophagy gene Epg5/epg-5 exhibit selective damage of cortical layer 5 pyramidal neurons and spinal cord motor neurons. Pathologically, Epg5 knockout mice suffered muscle denervation, myofiber atrophy, late-onset progressive hindquarter paralysis, and dramatically reduced survival, recapitulating key features of amyotrophic lateral sclerosis (ALS). Epg5 deficiency impaired autophagic flux by blocking the maturation of autophagosomes into degradative autolysosomes, leading to accumulation of p62 aggregates and ubiquitin-positive inclusions and leads to axonal degeneration and massive nonselective neuronal loss (Hara et al., 2006; Komatsu et al., 2006; Zhao et al., 2012). In contrast, neurodegenerative diseases, such as amyotrophic lateral sclerosis (ALS), Huntington’s disease, and Alzheimer’s disease, involve selective damage of certain neuronal populations (Hardy and Gwinn-Hardy, 1998). For example, ALS is characterized by the selective loss of motor neurons in cortical layer 5 and in the spinal cord and manifests as muscle weakness, also...
Epg5\textsuperscript{−/−} mice develop behavioral and motor abnormalities

We generated Epg5 knockout mice as shown in Fig. 1 (A and B). Epg5 heterozygous mouse (Epg5\textsuperscript{+/-}) had no noticeable pathological defects over a 2-yr period. Epg5 homozygous offspring (Epg5\textsuperscript{−/−}) were born at Mendelian frequency and appeared normal at birth but showed growth retardation and reduced survival (Fig. 1 C). Epg5\textsuperscript{−/−} males developed progressive behavioral and motor deficits. They showed poor motor coordination in a rotarod test at 4 mo (Fig. 1 D) and gradually exhibited limb-clasping reflexes (Fig. 1 E). The hindquarters became completely paralyzed and stiff in Epg5\textsuperscript{−/−} males at 10 mo of age, accompanied by poor grooming, rough coat, flank muscle wasting, and notable kyphosis (Fig. 1, F and G). Epg5\textsuperscript{−/−} males started to die by 10 mo after birth. Epg5\textsuperscript{−/−} females were similarly affected, but the phenotype occurred later than in males (Fig. 1 H). Three out of four females died at 8–10 mo with slight hind-limb paralysis and muscle wasting, and one female survived to 12 mo and showed complete hind-limb paralysis.
Selective vulnerability of cerebral cortex layer 5 and hippocampal pyramidal neurons and spinal cord motor neurons in Epg5−/− mice

Histological analyses were performed on cerebrum, cerebel- lum, and spinocerebellar tract motor neurons from control and mutant mice at 10 mo of age. In Epg5−/− males, total brain size was comparable to controls. The number of neurons in cortical layers 1–4 remained unchanged (Fig. S1 A), but the number of pyramidal neurons in layer 5 of the motor and sensory cortices was reduced by 28 and 37%, respectively (Figs. 2, A and B; and S1 B). The thickness and cell number of the hippocam- pal pyramidal cell layer was reduced (Fig. 2, C and D). The pattern of nerve fibers in the alveus of the hippocampus was slightly irregular (Fig. S1 C).

Epg5−/− mutant cerebella appeared normal: the size, foliation, and fissuration were comparable with controls as was the thickness of the molecular layer (Fig. 2, E and F). Purkinje cell numbers and the distribution of Purkinje cell axon terminals in the deep cerebellar nuclei (DCN) of Epg5-deficient mice were similar to controls (Figs. 2, G and H; and S1 D). Epg5−/− mice had fewer motor neurons in the anterior horn of the cervical, thoracic, and lumbar spinal cord (Fig. 2, I and J). Many of the remaining motor neurons were pyknotic (Fig. 2 I). The number of interneurons was normal (Fig. 2 K). Eosino- philic spheroids accumulated in the dorsal corticospinal tract of the cervical, thoracic, and lumbar spinal cord in Epg5−/− mice (Fig. 2 L). The eosinophilic spheroids were stained by the axon marker β-tubulin III and were surrounded by discontinuous myelin basic protein–stained myelin (Fig. 2 M), indicating that these spheroids were swollen degenerating axons. Ultrastructurally, Epg5−/− mice brains contain many atrophic motor neurons in the cortex (Fig. S1 F). Epg5−/− mice also showed many degenerating axons and swollen mitochondria in the spinal cord (Figs. 2 N and S1 E). Expression of the glial marker glial fibrillary acidic protein (GFAP) increased in various regions of the cerebellum, cerebral cortex, and spinal cord of mutant mice compared with control littermates, indicating that Epg5−/− mice exhibit reactive astrogliosis (Figs. 2, O and P; and S1, G and H).

Amplification of p62 aggregates and ubiquitin-positive inclusions in Epg5-deficient mice

We next examined whether Epg5 is essential for basal autophagy activity by examining the level and distribution of p62, polyubiquitinated proteins, and LC3 (mammalian Atg8 homologue; Mizushima et al., 2010). Brain and spinal cord extracts from Epg5−/− mice at 10 mo showed dramatic accumulation of p62, high molecular mass polyubiquitinated proteins, and LC3-II, a lipidated form of LC3 that associates with autophagic vacuoles (Fig. 3, A and B). p62 mRNA levels and proteasome activity were comparable in controls and mutants (Figs. S1, I and J). Cytoplasmic p62 aggregates and ubiquitin-positive inclusions accumulated in various regions of the cerebrum, cerebellum, and spinal cord of Epg5−/− but not Epg5+/+ mice (Figs. 3 C and S1, K–N). p62 colocalized with neuronal nuclei–labeled neurons and anti-CNPass (2’,3’-cyclic-nucleotide 3’-phosphodiesterase)–labeled oligodendrocytes but absent from GFAP-stained astroglia (Fig. 3, D–F). Ubiquitin-positive inclusions colocalized with p62 aggregates, but they were fewer in number than p62 aggregates (Figs. 3 C and S1 N). LC3 was diffusely localized with very few puncta detected in wild-type brain and spinal cord (Fig. S1, P–R). In Epg5−/− mice, LC3 foci dramatically accumulated in SMI 32–labeled motor neurons and oligoden- droglia (Fig. S1 P). LC3 immunoreactivity largely colocalized with p62 aggregates (Fig. S1, O, Q, and R). Thus, Epg5 deficiency impairs autophagic flux, leading to accumulation of LC3 puncta, p62 aggregates, and ubiquitin-positive inclusions in neurons and glial cells.

Epg5 deficiency causes accumulation of cytoplasmic TDP-43 aggregates

Accumulation of cytoplasmic TDP-43 aggregates in vulnerable neurons is a common feature of ALS and frontotemporal lobar degeneration (Neumann et al., 2006). In Epg5−/− mice, TDP-43 showed nuclear staining in neurons in various regions of brain and spinal cord (Fig. 3 G). However, a large number of TDP-43 aggregates accumulated in the cytoplasm in motor neurons in the spinal cord and the fifth layer of cortices and, to a lesser extent, in neurons in other brain regions of Epg5−/− mice (Figs. 3, G and H; and S1 T). Protein levels of TDP-43 remained unchanged in mutant mice (Fig. S1 S). Cytoplasmic TDP-43 aggregates were absent in motor neurons in E124H-deficient mice (Fig. S1 U).

Epg5-deficient mice show muscle denervation and myofiber degeneration

Electromyography was performed to determine whether Epg5 deficiency causes muscle denervation, characteristic of the early stage of motor neuron diseases (Boilée et al., 2006). Fibri- llation and positive sharp waves were found in the resting gastroc- nemius (lower hind limb) muscle of 3-mo-old Epg5−/− mice (presymptomatic), indicative of active denervation of muscle fibers (Figs. 4, A and B). At 7 mo (early symptomatic), the duration of the motor unit action potential (MUAP) increased 44% in Epg5−/− mice compared with controls, indicating reinnervation of denervated muscle fibers by the sprout from relatively normal axons. At 10 mo, MUAP duration increased 78% in Epg5−/− mice with occasional high-amplitude and long-duration action potentials (Fig. 4 C), but no fibrillation or positive sharp waves were found, indicating no obvious active denervation in end-stage Epg5−/− mice. More- over, the bicipital (upper forelimb) muscle started to suffer de- nervation changes at 10 mo, indicated by the appearance of positive sharp waves. The evoked motor response (compound muscle action potential [CMAP]) of the gastrocnemius muscle elicited by distal stimulation of the sciatic nerve in Epg5−/− mice at 7 and 10 mo exhibited increased latency and decreased amplitude (Fig. S2 A), indicating axonal degeneration. Thus, Epg5−/− mice showed neurogenic muscle damage progressing from hind limb to forelimb.

Histological analysis of the gastrocnemius muscle of Epg5−/− mice at 10 mo revealed myofiber degeneration, including...
Figure 2. Defects in brain and spinal cord in Epg5<sup>−/−</sup> mice. (A) Nissl staining of the fifth layer of the cortex in 10-mo-old Epg5<sup>+/−</sup> and Epg5<sup>−/−</sup> mice. (B) The number of pyramidal cells in the fifth layer per millimeters squared. (C and D) H&E staining of the hippocampus shows reduced pyramidal cell numbers in Epg5<sup>−/−</sup> mice. Means ± SEM of three mice are shown in B and D. (E) The normally developed cerebellum in Epg5<sup>−/−</sup> mice. (F) The thickness of the molecular layer in the cerebellum in mutant and control mice. (G and H) Purkinje cells, stained by calbindin, are not reduced in number in Epg5<sup>−/−</sup> mice. Means ± SEM of five mice are shown in F and H. (I) Arrows indicate pyknotic neurons in the anterior horn of Epg5<sup>−/−</sup> mice. (J) The number of motor neurons in the spinal cord (SC) in Epg5<sup>+/−</sup> and Epg5<sup>−/−</sup> mice. (K) The numbers of Nissl-stained interneurons in the spinal cord are similar in control and Epg5<sup>−/−</sup> mice. Means ± SEM of three mice are shown in J and K. (L) Eosinophilic spheroids (arrows) accumulate in the thoracic spinal cord of Epg5<sup>−/−</sup> mice. (M) Eosinophilic spheroids are swollen degenerating axons (arrow). (N) EM pictures showing degenerated axons (arrows) in the spinal cord of Epg5<sup>−/−</sup> mice. (O and P) GFAP staining in Epg5<sup>+/−</sup> and Epg5<sup>−/−</sup> mice. DCST, dorsal corticospinal tract. Bars: (A, C, L, and O) 50 µm; (E) 500 µm; (G) 20 µm; (I) 100 µm; (M and P) 10 µm; (N) 2 µm.
centrally nucleated fibers, vacuolated fibers, irregularly shaped atrophic fibers, and decreased myofiber size (Fig. 4, D–F). Consistent with this, the atrophy-related genes were up-regulated in Epg5−/− muscles (Fig. 4 G). Ultrastructurally, Epg5−/− muscles exhibited misaligned Z lines and accumulation of abnormal enlarged mitochondria (Fig. 4, H and I). Autophagic flux was also impaired in atrophic muscles (gastrocnemius myofibers) but not in adductor longus and vastus medialis of Epg5−/− mice (Figs. 4, J–L; and S2, D and E).
Epg5 knockout mice show muscle atrophy. (A and B) Electromyography of the gastrocnemius muscle from a 3-mo-old Epg5−/− mouse showing fibrillations (A) and positive sharp waves (B). These defects were absent in control mice. (C) At 10 mo, Epg5−/− mice showed high-amplitude and long-duration action potentials when conducting MUAP tests. The blue lines show the start and end of an action potential. Dur, duration; Amp, amplitude. (D–F) H&E staining of gastrocnemius muscles showed features of muscle degeneration in Epg5−/− mice. The arrow in E indicates centrally nucleated fibers. (G) The transcription levels of the atrophy-related genes were up-regulated in gastrocnemius muscles of Epg5−/− mice. (H and I) Electron micrographs of Epg5−/− and Epg5+/− muscles. The arrows indicate abnormal enlarged mitochondria. (J) Levels of LC3-II and p62 in muscle extracts from Epg5+/− and Epg5−/− mice. (K) Percentage of p62 aggregate-positive myofibers in Epg5−/− and Epg5+/− mice. Means ± SEM of three mice are shown. (L) Accumulation and colocalization of p62 and ubiquitin aggregates in the gastrocnemius muscles of Epg5−/− mice but not controls. Bars: (D–F and I) 10 µm; (H and I) 1 µm.

Epg5−/− mice exhibit reduced adipose mass Epg5−/− mice at 10–11 mo of age showed dramatic reduced mass of white adipose tissue. Wild-type adipocytes are morphologically homogeneous, and the entire cytoplasm is occupied by a single large unilocular lipid droplet (Fig. S2 F). However, mutant adipocytes were much smaller, and a large population of cells contained smaller multilocular lipid droplets (Fig. S2 C, F, and G). Epg5−/− livers at 10 mo appeared grossly normal, and the hepatic lobular structure was unchanged (Fig. S2 H). Levels of LC3-II and p62 were increased in Epg5−/− adipose tissues and liver fractions at 10 mo (Fig. S2 B), indicating that autophagic flux is impaired in these tissues.

Epg5 deficiency impairs autophagosome maturation We further characterized the autophagy pathway in Epg5−/− primary mouse embryonic fibroblasts (MEFs). Under nutrient repletion conditions, LC3 puncta were largely absent from control MEFs but accumulated in Epg5−/− MEFs (Figs. 5, A and E; and S3, A–F). LC3-II levels were also elevated (Fig. 5 B). In control MEFs, the number of LC3 puncta dramatically increased 4 h after starvation and decreased 8 h after starvation, whereas in mutant MEFs, LC3 puncta continued to increase 8 h after starvation (Fig. 5, A and E; and S3, A–D). Upon bafilomycin A1 treatment, which inhibits lysosomal degradation, control MEFs showed increased LC3-II levels, whereas mutant
Figure 5. *Epg5* deficiency causes a defect in autophagosome maturation and impairs endocytic trafficking. (A) Number of LC3 puncta in *Epg5*+/− (control) and *Epg5*−/− MEFs. (B) Western blot showing levels of LC3-I and LC3-II in control and *Epg5*−/− MEFs upon indicated treatment (Starv., starvation; Rapa., rapamycin; Bafilo., bafilomycin A1). (C) Proteinase K (ProK) protection assay in different MEFs. PNS, postnuclear supernatant; P, pellet fractions. (D) Colocalization
MEFs exhibited no further elevation (Fig. 5 B), indicating that autophagic flux is impaired in Epg5<sup>−/−</sup> MEFs. Consistent with this, compared with control MEFs, Epg5<sup>−/−</sup> MEFs contained more p62 aggregates that largely colocalized with LC3 puncta (Fig. S3, E–H).

We performed a protease protection assay to determine whether autophagosome formation is completed in Epg5<sup>−/−</sup> MEFs. p62 sequestered into autophagosomes is resistant to proteinase K digestion, whereas treatment with Triton X-100 permeabilizes the membrane, making p62 accessible to digestion (Velikkakath et al., 2012). After proteinase K treatment in the absence of Triton X-100, p62 was digested in Atg7 (autophagosome defective) knockout MEFs but was protected in Epg5<sup>−/−</sup> and control MEFs, indicating that autophagosomes were sealed (Fig. 5 C). In control MEFs, the extensive colocalization of LC3 with LAMP-1 and LysoTracker-stained acidic lysosomes 4 h after starvation had decreased by 8 h, whereas in Epg5<sup>−/−</sup> MEFs, the colocalization continued to increase 8 h after starvation (Figs. 5, D and E; and S3, A–D, I, K, and L). p62 immunoreactivity also partially colocalized with LAMP-1–labeled structures in Epg5<sup>−/−</sup> MEFs (Fig. S3 J). However, lysosomal degradation of the self-quenched fluorescent dye DQ-BSA was much lower in Epg5<sup>−/−</sup> MEFs than control MEFs (Fig. S3, M and N). These results indicate that autophagosomes fuse with late endosomes/lysosomes in Epg5<sup>−/−</sup> MEFs, but the proteolysis activity of autolysosomes is greatly impaired.

No obvious autophagic structures were observed by EM in unstarved control MEFs (Fig. 5 F). However, Epg5<sup>−/−</sup> MEFs contained numerous autophagic vacuoles at all stages, including autophagosomes, possible amphisomes, and early autolysosomes (aAV-I [advanced autophagic vacuole type I], containing recognizable cytoplasmic content) or late autolysosomes (aAV-II, containing digested, recognizable content; Figs. 5, G–I; and S3 O). Epg5<sup>−/−</sup> MEFs had many complex vacuoles containing multiple membrane layers even under food repletion conditions (Fig. 5 H), suggesting that fusion of autophagosomes with lysosomes is hindered or the degradative activities of autolysosomes are impaired. After 4-h starvation, aAV-I and aAV-II autophagic vacuoles were increased in control and even more so in Epg5<sup>−/−</sup> MEFs (Fig. S3, P–S). Final-stage autolysosomal vacuoles (aAV-III), which contain a lipid-like area, were also detected in Epg5<sup>−/−</sup> MEFs under basal and starvation conditions (Figs. 5 I and S3, R and S), suggesting that autophagic degradation still proceeds slowly in Epg5-depleted cells.

Epg5 knockdown impairs endosomal trafficking

We next investigated the role of Epg5 in the endocytic pathway. Compared with control cells, early endosomes and late endosomes were significantly enlarged in Epg5 siRNA-treated cells (Figs. 5 J and S3 T). Degradation of EGF receptor (EGFRI) was significantly slowed in Epg5 siRNA-treated cells (Fig. 5, K and L), indicating that the endocytic degradation pathway is defective in Epg5 knockdown cells. We further investigated endocytic recycling by following the transferrin transport, which is internalized into early endosomes and eventually returned to the surface and exported from the cell. In control cells, fluorescently conjugated transferrin was first localized on the plasma membrane, mostly recycled to the cell surface and released within 30 min, and completely disappeared after 1 h (Fig. 5 M). In Epg5 siRNA-treated cells, internalized transferrin accumulated in the cytoplasm (Fig. 5 N), which partially colocalized with enlarged early and late endosomes but not with LysoTracker-stained acidified lysosomes (Fig. S3, V and W). After 30-min chasing, a large amount of transferrin was retained in the cytoplasm (Fig. 5 N). After 1 h, most transferrin was recycled to the surface and released (Fig. 5 N). These results indicate that loss of Epg5 function slows endocytic degradation and delays endocytic recycling.

Conclusion

Here, we demonstrated that mice deficient in Epg5 exhibit key features of ALS, including selective vulnerability of cortical layer 5 pyramidal neurons and spinal cord motor neurons, accumulation of cytoplasmic TDP-43 aggregates in motor neurons, progressive muscle weakness and atrophy, muscle denervation, and premature death. Although autophagy flux is systematically impaired and p62 aggregates accumulate in many regions of brain and spinal cord in Epg5 knockout mice, only specific neuronal populations are degenerated. Selective neuronal damage in Epg5 knockout mice is unlikely caused by the fact that motor neurons are more vulnerable to the autophagic defect. Ei24 deficiency causes dramatically reduced interneurons, whereas motor neurons are largely normal, and there is no accumulation of eosinophilic spheroids in spinal cord (Zhao et al., 2012). The massive neuron degeneration in Atg5<sup>−/−</sup>, Atg7<sup>−/−</sup>, and Ei24-deficient mice could be caused by the fact that these genes have autophagy-independent function. For example, Atg5 and Atg7 also participate in cell secretion (DeSelm et al., 2011). Accumulation of nondegradative autophagic vacuoles may contribute to the pathogenesis of ALS in Epg5-deficient mice.
Functional impairment of the endosomal sorting complex required for transport or valosin-containing protein, which causes accumulation of nondegradative autophagic vacuoles, has been linked with familial ALS (Parkinson et al., 2006; Filimonenko et al., 2007; Lee et al., 2007; Ju et al., 2009; Johnson et al., 2010; Tresse et al., 2010). The ALS3 locus is mapped to chromosome 18q21, where EPG5 is located (Hand et al., 2002). Our study implies that human EPG5 might define an ALS susceptibility locus, and the animal model established here will help us to understand the molecular pathogenesis of ALS and to develop new therapeutic treatments.

Materials and methods

Generation of knockout mice

The Epg5 targeting vector was constructed by substituting exons 18 and 19 of Epg5 genomic DNA with a neomycin cassette. The Epg5 conventional knockout mice were generated in the National Institute of Biological Sciences Transgenic Research Center. The detailed procedure is as follows: After electroporation of the targeting vector into 129 R1 embryonic stem cells, colonies resistant to neomycin were chosen for homologous recombination by Southern blotting with probes 5’ and 3’ of the genomic sequence in the targeting vector. Heterozygous Epg5+/− embryonic stem cell clones were used to microinject C57BL/6N blastocysts. Chimeric offspring were mated with wild-type C57BL/6N mice. Heterozygotes from this cross were mated to obtain homozygous mutant mice. Germine transmission was confirmed by PCR analysis of tail DNA. The following primers were used to detect wild-type and deleted Epg5 alleles: 5’-AGAAAGGTGGTGAGGACCTTC-3’ [forward wild-type allele], 5’-ATGGGAAAGACATAGCCCCGATG-3’ [forward deleted allele], and 5’-TCTCTTACCACAGGGCTGCAAG-3’ [reverse common]. The expected sizes are 531 and 273 bp, respectively.

E14foxfox, nestin-Cre mice were on the C57BL/6N background, and exon 3 of E14 was flanked by two fox sequences. E14foxfox mice were crossed with nestin-Cre to generate neural-specific E14 deficient mice. All mice were kept under specific pathogen-free conditions in the animal facility at the National Institute of Biological Sciences, Beijing. All animal experiments were approved by the institutional committee of the National Institute of Biological Sciences, Beijing.

Antibodies

The following antibodies were used: rabbit anti-p62 (PM045; MBL International), mouse anti-p62 (ab66416; Abcam), mouse antibiubiquitin (3936; Cell Signaling Technology), rabbit anti-GEAP (bs-0199R; Bioss, Inc.), rabbit anti-AT3 (for Western blotting: PM046; MBL International), rabbit anti-LC3 (for immunostaining: 2775; Cell Signaling Technology), mouse anti-neuronal nuclei (MAB3877; EMD Millipore), mouse anti-β-tubulin III (ab7751; Abcam), mouse anti-calbindin (C9848; Sigma-Aldrich), mouse anti-CNPase (ab3619; Abcam), rabbit anti-myelin basic protein (ab40390; Abcam), rabbit anti-α-tubulin (3470; Cell Signaling Technology), rabbit anti-Rab7 (3547; Cell Signaling Technology), rabbit anti-β-actin (60008-1-lg; Proteintech). Rabbit anti-Epg5 antibody was developed against a fragment of mouse Epg5 protein (amino acids 1,061–1,169).

Behavioral analysis

To monitor motor function, an accelerating rotorod (YIS-4C; Beijing Zhongshidichuang Science and Technology Development Co., Ltd.) was used. After training for 2 d, mice were placed on the rolling rod with autoacceleration ranging from 5 to 20 rpm within 60 s for a maximum of 4 min. The time the animal stayed on the rod was measured.

Histology and immunohistochemistry

Mice were transcardially perfused with 10% neutral buffered formalin (Sigma-Aldrich). Tissues were postfixed and embedded in paraffin. For histological analysis, 5-µm sections were stained with hematoxylin and eosin (H&E) and examined by light microscopy (Imager.A1; Carl Zeiss) with a fluorescence microscope or a confocal microscope (LSM 510 Meta plus Axiovert zoom; Carl Zeiss) with 63x/1.40 NA oil immersion objective lens (Plan-Apochromat; Carl Zeiss) and a camera (AxioCam HRm; Carl Zeiss) at RT. Images were processed and viewed using LSM Image Browser software. Sections of the lumbar spinal cord were stained in this study unless otherwise noted.

Cell counting

To count the number of spinal cord motor neurons, fixed spinal cords were sequentially sectioned at 5 µm, and every sixth section was stained with hematoxylin and eosin (Sigma-Aldrich). Tissues were postfixed and embedded in paraffin. For his-
with 2.5 µl of each substrate (final concentration of 50 nM) for 30 min at 37°C in the dark. After adding 252.5 µl of precooled 96% ethanol solution to stop the reaction, proteasome activity of the mixture was measured by detecting fluorescence from AMC hydrolysis (380-nm excitation and 460-nm emission).

Transmission EM
Mice were perfused with 2% PFA/2% glutaraldehyde. Tissues were dissected and postfixed in 2.5% glutaraldehyde followed by fixation in 1% OsO4 for 2 h. After dehydration with graded ethanol solutions, tissues were embedded in EMBed812 (Electron Microscope Sciences). 80-nm ultrathin sections were stained with 2% uranyl acetate for 30 min and lead citrate for 10 min. Cells were fixed with 2% PFA and 0.25% glutaraldehyde in PBS, pH 7.4, at 37°C for 2 h and then dehydrated in a graded ethanol series and embedded. 70-nm ultrathin sections were mounted, stained, and dried at RT. The samples were visualized using a 120-kV electron microscope (H-7650B; Hitachi) at 80 kV, and images were captured by a charge-coupled device camera (XR-41; Advanced Microscopy Techniques) using DigitalMicrograph software (Gatan, Inc.) at RT.

Cell culture, transfection, and immunostaining
Primary MEFs generated from day 13.5 embryos from Epg5−/− crosses or HeLa cells were cultured in DMEM supplemented with 10% FBS, 100 µg/ml penicillin, 100 µg/ml streptomycin, and 2 mM glutamine. Epg5 siRNA and nonsense (NS) siRNA were purchased from GenePharma. The sequences were as follows: NS siRNA, 5′-UUCUCCGAACGUGUCACGU-3′; and Epg5 siRNA, 5′-GAAACGUGUAACCCAGUCU-3′. Cells were transfected with siRNAs using Lipofectamine 2000 (Invitrogen) for 72 h according to the manufacturer’s instructions.

For immunostaining, cells were plated on glass-bottomed dishes. After overnight recovery, cells were starved in serum-free medium for the indicated time. Cells were washed with PBS, fixed in 4% PFA for 10 min, and permeabilized in cold methanol for 10 min. After blocking with 10% PBS in PBS for 30 min at RT, cells were incubated with antibodies overnight at 4°C, washed with PBS three times, and then stained with fluorescein isothiocyanate labeled secondary antibodies. Coverslips were mounted on microscope slides with mounting medium with DAPI. Images were acquired with a confocal microscope (LSM 510 Meta). Dot numbers were analyzed by ImageJ software (National Institutes of Health), and colocalization masks and ratios were analyzed by Imaris software (Bitplane).

DQ-BSA degradation assay
72 h after transfection with Epg5 siRNA or NS siRNA, cells were serum starved for 2 h and then treated with 10 µg/ml DQ-BSA (Life Technologies/Invitrogen) for 30 min. Cells were then washed with PBS, fixed, permeabilized, and stained with DAPI. The fluorescent signal from lysosomal proteolysis of DQ-BSA was recorded by confocal microscopy (LSM 510 Meta).

Transferin recycling
Cells were cultured in 3.5-cm dishes with a 1-cm microwell and transfected with NS siRNA or Epg5 siRNA. 72 h after transfection, 0.1 mg/ml Alexa Fluor 488–conjugated transferrin (Invitrogen) was added into each plate and incubated for 30 min on ice. Cells were washed with PBS and incubated with regular medium at 37°C for the indicated times before transferrin distribution was examined. For examining the colocalization of transferrin with various endosome compartments, LC3 puncta, and p62 aggregates, cells transfected with Epg5 siRNA or NS siRNA were incubated with 0.1 mg/ml Alexa Fluor 488–conjugated transferrin in regular medium for 1 h at 37°C. Then, the cells were fixed, permeabilized, and incubated with anti-Rab5, Rab7, -LC3, or -p62 antibodies and secondary antibodies.

EGRF degradation assay
Control and Epg5 siRNA-treated HeLa cells were serum starved overnight and treated with 100 ng/ml EGF and 25 µg/ml cycloheximide for the indicated period. Cells were then washed with PBS and lysed. Degradation of EGRF was examined using an immunoblotting assay with the anti-EGRF antibody.

Protease protection assay
Cultured MEF cells were starved, treated with bafilomycin A1, harvested, and suspended in homogenization buffer (20 mM Hepes/KOH, pH 7.4, 0.22 M mannitol, 0.07 M sucrose, and protease inhibitors). Atg7 knockout MEFs were provided by M. Komatsu (Tokyo Metropolitan Institute of Medical Science, Setagaya-ku, Tokyo, Japan). Cells were then passed 10 times through a 27-gauge needle using a 1-ml syringe and centrifuged at 2,000 rpm for 5 min. The supernatant (as postnuclear supernatant) was centrifuged again at 10,000 rpm to obtain the pellet fraction. The pellet was resuspended in homogenization buffer and then treated with 100 µg/ml proteinase K with or without 0.5% Triton X-100. After 20-min incubation on ice, 10% TCA was added, and the samples were centrifuged at 10,000 rpm for 5 min. The pellet (as pellet fraction) was washed with ice-cold acetone, resuspended in SDS-PAGE sample buffer, and boiled immediately. Proteinase K digestion was detected by Western blotting.

Electromyography
Evoked CMAP, MUAP, and resting potentials were evaluated with an electromyogram apparatus (Keypoint; Dantec) and analyzed by software (Keypoint.NET; Dantec). For evaluating CMAP, we stimulated the sciatic nerve at the tuberosity of the ischium by a single 0.1-ms, 1-Hz, and 20-nA supramaximal pulse and recorded CMAPs from the medial part of the gastrocnemius with a 26-gauge concentric needle electrode [37 × 0.64 mm, recording area of 0.07 mm²] (Alpine). The MUAP of the gastrocnemius and bicipital muscles was recorded with the same type of needle electrode from a single motor unit at moderate contractions. Resting potentials were recorded with the same needle electrode.

Statistical analysis
Data from at least three sets of samples were used for statistical analysis. Statistical significance was calculated by Student’s t test. P < 0.05 was considered significant.

Online supplemental material
Fig. S1 shows defects in the brain and spinal cord in Epg5−/− mice. Fig. S2 shows muscle atrophy in Epg5−/− mice. Fig. S3 shows a defect in the maturation of autophagosomes into degradative autolysosomes in Epg5−/− MEFs. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201211014/DC1.

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