

UCSF

UC San Francisco Previously Published Works

Title

The conserved autoimmune-disease risk gene TMEM39A regulates lysosome dynamics

Permalink

<https://escholarship.org/uc/item/7kf6z83h>

Journal

Proceedings of the National Academy of Sciences of the United States of America,
118(6)

ISSN

0027-8424

Authors

Luo, Shuo
Wang, Xin
Bai, Meirong
et al.

Publication Date

2021-02-09

DOI

10.1073/pnas.2011379118

Peer reviewed



The conserved autoimmune-disease risk gene *TMEM39A* regulates lysosome dynamics

Shuo Luo^a, Xin Wang^a, Meirong Bai^a, Wei Jiang^a, Zhe Zhang^a, Yifan Chen^b, and Dengke K. Ma^{a,c,1}

^aCardiovascular Research Institute, University of California, San Francisco, CA 94158; ^bDepartment of Molecular and Cell Biology, University of California, Berkeley, CA 94720; and ^cDepartment of Physiology, University of California, San Francisco, CA 94158

Edited by Victor R. Ambros, University of Massachusetts Medical School, Worcester, MA, and approved December 15, 2020 (received for review June 9, 2020)

***TMEM39A* encodes an evolutionarily conserved transmembrane protein and carries single-nucleotide polymorphisms associated with increased risk of major human autoimmune diseases, including multiple sclerosis. The exact cellular function of *TMEM39A* remains not well understood. Here, we report that *TMEM39*, the sole *Caenorhabditis elegans* (*C. elegans*) ortholog of *TMEM39A*, regulates lysosome distribution and accumulation. Elimination of *tmem-39* leads to lysosome tubularization and reduced lysosome mobility, as well as accumulation of the lysosome-associated membrane protein LMP-1. In mammalian cells, loss of *TMEM39A* leads to redistribution of lysosomes from the perinuclear region to cell periphery. Mechanistically, *TMEM39A* interacts with the dynein intermediate light chain *DYNC112* to maintain proper lysosome distribution. Deficiency of *tmem-39* or the *DYNC112* homolog in *C. elegans* impairs mTOR signaling and activates the downstream TFEB-like transcription factor *HLH-30*. We propose evolutionarily conserved roles of *TMEM39* family proteins in regulating lysosome distribution and lysosome-associated signaling, dysfunction of which in humans may underlie aspects of autoimmune diseases.**

TMEM39A | lysosome | dynein | autoimmunity | autoimmune disorders

Autoimmune diseases affect millions of people worldwide, while exact causes of the diseases remain not well understood, despite decades of research (1–3). Recently, genome-wide association studies have identified single-nucleotide polymorphisms (SNPs) in *TMEM39A* that are associated with enhanced risks of multiple sclerosis (MS), systemic lupus erythematosus (SLE), and autoimmune thyroid disease (4–8), indicating an important link between *TMEM39A* and autoimmune disease. *TMEM39A* is also overexpressed in glioma cell lines and glioma patient cells (9), implicating its possible role in glioma pathogenesis. *TMEM39A* encodes a predicted multitransmembrane protein, homologs of which are found in most multicellular organisms (Fig. 1 *A* and *B* and *SI Appendix*, Fig. *S1A*). A recent study reported that *TMEM39A* localizes to the endoplasmic reticulum (ER) and regulates autophagy through controlling trafficking of the PtdIns(4)P phosphatase *SAC1* (10). To investigate *TMEM39A*'s physiological roles and whether its cellular functions might be evolutionarily conserved, we generated *Caenorhabditis elegans* (*C. elegans*) strains and HEK 293T human cell lines that lack respective *TMEM39* homologs in each system. Phenotypic and protein-interactor analyses revealed conserved roles of *TMEM39* in regulating lysosome dynamics and signaling, providing a plausible link of human *TMEM39A* to pathogenesis of autoimmune diseases.

Results

We identified the uncharacterized *C. elegans* gene *D1007.5* (named as *tmem-39* hereafter) in a previous genome-wide RNA interference (RNAi) screen for cellular-stress regulators (11). *tmem-39* encodes the only homolog of mammalian *TMEM39A* in the *C. elegans* genome. Using CRISPR-Cas9, we either removed the entire coding region of *tmem-39* or inserted stop codons in the first exon of the gene (Fig. 1*C*). Mutant animals carrying *tmem-39* null alleles showed multiple developmental abnormalities, including slow growth, reduced body length, and a tendency to burst, indicating defects in cuticle development (Fig. 1*D* and *SI*

Appendix, Fig. *S1B*). In addition, *tmem-39* mutant animals also have markedly higher expression of ER stress reporter *hsp-4_p::gfp* in the hypoderm, supporting an important role of *TMEM39* in normal protein homeostasis (Fig. 1*D*). To investigate if disruption of *TMEM39* impairs the nervous system, the primary target of MS, we examined neuronal morphology of the mutant animals using transgenic strains that express cholinergic (*unc-17_p::gfp*), GABAergic (*unc-47_p::gfp*), or somatosensory PVD interneuron (*F49H12.4::gfp*) reporters. About 10 to 20% of the wild-type animals showed mild morphological abnormalities (e.g., short filopodia-like structures) when they aged (Table 1 and *SI Appendix*, Fig. *S1G*). Strikingly, disruption of *tmem-39* led to severe morphological defects of the nervous system in 50 to 90% of the mutant animals examined, including long ectopic branches, incorrect neurite projections, and beading of neuronal processes (Fig. 1*E* and Table 1). We also found that the dendritic structures of the PVD neurons were missing in L2 and L3 larva, despite the presence of the PVD soma, suggesting developmental defects of the neuronal processes (*SI Appendix*, Fig. *S1H*). Together, our observations indicate that *TMEM39* is important for maintaining normal ER homeostasis and neuronal morphology in the nervous system.

To determine the subcellular localization of *TMEM39* protein, we generated transgenic strains that express either a rescuing *tmem-39* genomic transgene with *mCherry* tagged to the N terminus or *N-mCherry*-tagged *tmem-39* complementary DNA (cDNA) under an *hsp-16.48* heat-shock promoter. The *tmem-39*

Significance

Autoimmune diseases are characterized by overactivation of an organism's immune system and subsequent tissue damage and pathogenesis. Although genome-wide association studies have identified genetic variations, such as single-nucleotide polymorphisms (SNPs), that are linked to autoimmunity, the exact function of the genes involved and molecular mechanisms underlying their contribution to autoimmunity remain not well understood. Here, we show that an evolutionarily conserved autoimmunity risk gene, *TMEM39A*, regulates lysosome dynamics through interacting and functioning with molecular motor dynein intermediate light chain. As lysosomes are implicated in various aspects of immune-system dysfunction, including abnormal cytokine secretion and antigen presentation, our study provides an important mechanistic insight suggesting how *TMEM39A* might contribute to autoimmunity like multiple sclerosis and lupus.

Author contributions: S.L. and D.K.M. designed research; S.L., X.W., M.B., W.J., Z.Z., Y.C., and D.K.M. performed research; S.L. and D.K.M. analyzed data; and S.L. and D.K.M. wrote the paper. The authors declare no competing interest.

This article is a PNAS Direct Submission.

Published under the PNAS license.

¹To whom correspondence may be addressed. Email: dengke.ma@ucsf.edu.

This article contains supporting information online at <https://www.pnas.org/lookup/suppl/doi:10.1073/pnas.2011379118/-DCSupplemental>.

Published February 2, 2021.

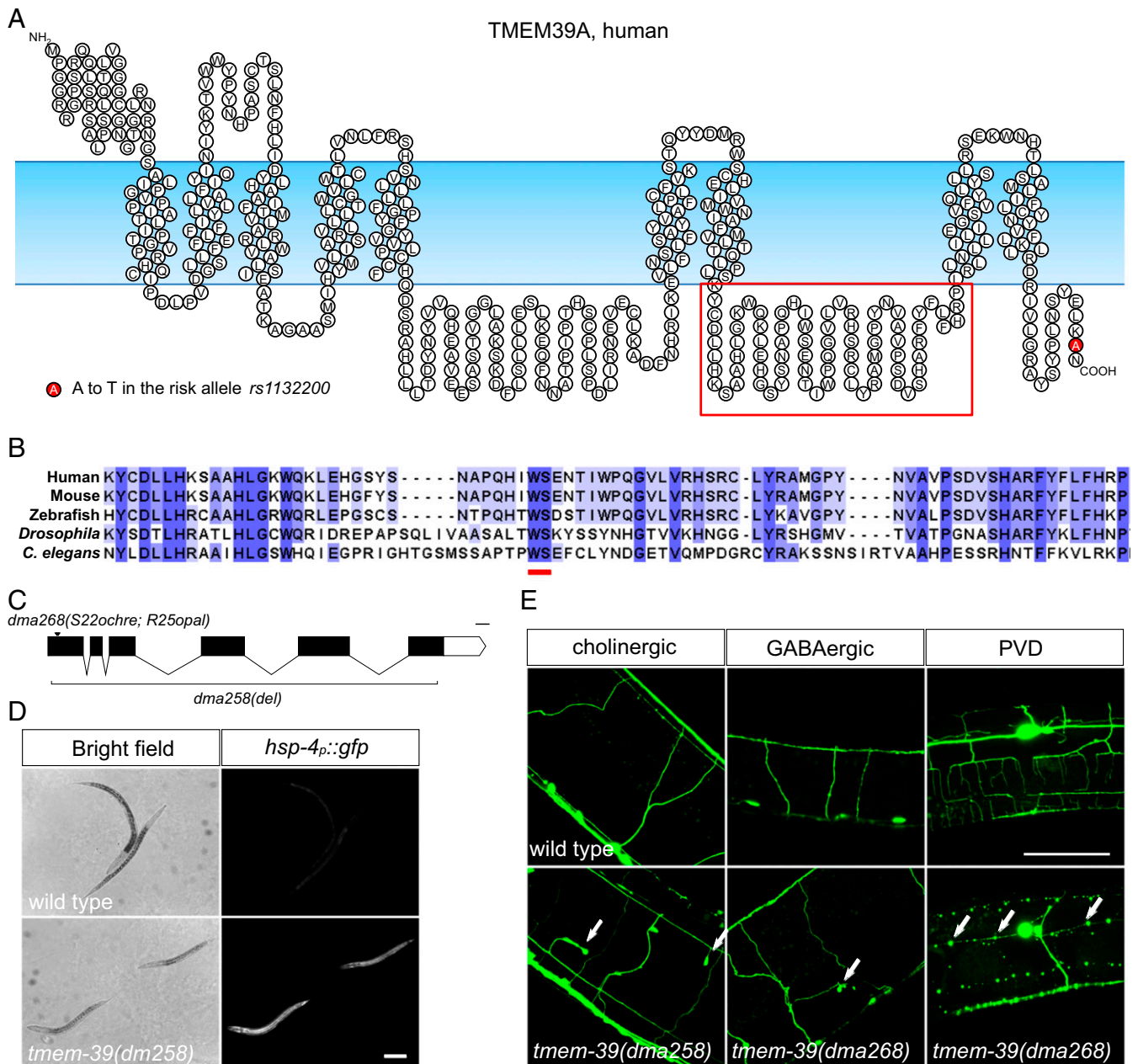


Fig. 1. TMEM-39 belongs to a conserved family of transmembrane proteins in metazoans and its role in maintaining normal neuronal morphology in *C. elegans*. (A) Schematic diagram showing the predicted structure of human TMEM39A protein, which contains nine transmembrane domains and two large loops (by SOSUI structural prediction server: https://harrier.nagahama-i-bio.ac.jp/sosui/sosui_submit.html). The autoimmune-risk allele *rs1132200* confers an A-to-T substitution in the C terminus of the protein, which is represented by a red circle. The red box highlights the loop domain aligned in B. (B) Sequence alignment of the second loop of the TMEM39A protein from the worm to human. The conserved WS residues within the loop are highlighted by a red line. (C) Schematic diagram showing the genomic structure and loss-of-function alleles of *C. elegans tmem-39*. *dma258* contains a deletion that removes most of the *tmem-39* coding sequences, while *dma268* contains two early stop codons in *tmem-39* that likely render the allele null. (D) Bright-field and fluorescence images showing wild-type and *tmem-39* mutant animals, with the ER stress reporter *hsp-4_p::gfp* markedly up-regulated in *tmem-39* mutants. (Scale bar, 100 μ m.) (E) The *tmem-39* mutants show broad morphological defects in the nervous system exemplified by ectopic branching (labeled by cholinergic reporter *unc-17_p::gfp* and GABAergic reporter *unc-47_p::gfp*; arrows) and neurodegeneration-like beading phenotypes (labeled by the PVD neuron reporter *F49H12.4::gfp*; arrows). (Scale bar, 40 μ m.)

genomic construct rescued developmental defects of *tmem-39* mutant animals, including dumpiness and elevated *hsp-4_p::gfp* ER stress reporter expression (SI Appendix, Fig. S1 D–F). We observed broad mCherry::TMEM-39 expression in multiple tissues, particularly the nervous system, hypoderm, and intestine (Fig. 2A). Both transgenic strains showed TMEM-39 localization to cytosolic puncta, largely colocalizing with GFP-tagged lysosome-associated membrane protein LMP-1 (Fig. 2B and

C). By contrast, TMEM-39 did not colocalize with a Golgi-targeted reporter *mans::gfp* (SI Appendix, Fig. S2 A and B), suggesting that TMEM-39 protein primarily localizes to compartments closely associated with lysosomes, at least in hypoderm. To investigate if disruption of TMEM-39 impairs lysosome morphology and/or function, we generated a compound mutant *tmem-39(null);dmaIs58[rpl-28_p::lmp-1::gfp]* and examined lysosome reporter expression (under the ubiquitous promoter *rpl-28*)

Table 1. Neuronal defects of adult (2 d after L4 stage) *tmem-39* mutants

Neuronal reporter and genotype	Abnormal, %	<i>n</i>	Morphological defects
Cholinergic*			
Wild type	26	54	Short filopodia-like structures
<i>tmem-39(dma258)</i>	96	50	Long ectopic branches; incorrect projections
<i>tmem-39(dma268)</i>	80	50	Long ectopic branches; incorrect projections
GABAergic†			
Wild type	16	32	Short filopodia-like structures
<i>tmem-39(dma258)</i>	56	50	Beading on neurites
<i>tmem-39(dma268)</i>	36	50	Long ectopic branches; incorrect projections
PVD‡			
Wild type	10	10	Mild beading on neurites
<i>tmem-39(dma258)</i>	73	11	Extensive degeneration-like beading on neurites
<i>tmem-39(dma268)</i>	77	13	Extensive degeneration-like beading on neurites

*Contains *vsls48[unc-17_p::gfp]*.

†Contains *oxls12[unc-47_p::gfp]*.

‡Contains *wds52[F49H12.4::gfp]*.

in *tmem-39* null mutant animals. We found that *tmem-39* mutant animals showed markedly higher accumulation of LMP-1::GFP in intestine, hypoderm, pharynx, and neuronal processes along the body, but to a lesser extent in the nerve ring, indicating that the level of lysosome reporter up-regulation is likely tissue-type dependent (Fig. 2D). Consistent with the observations, a second transgenic strain that expresses *lmp-1::gfp* specifically in neurons (under a pan-neuronal promoter *rab-3*) showed a similar pattern of lysosome reporter accumulation in nerve ring and nerve cords upon *tmem-39* disruption (Fig. 2D and E). By contrast, the *tmem-39* null mutants showed slightly decreased expression of a different reporter *mCherry::pdr-1* (which encodes the ortholog of human PRKN protein) driven by the ubiquitous promoter *rpl-28*, indicating that the lysosome reporter up-regulation is likely caused by changes in protein dynamics rather than transcriptional up-regulation of the *rpl-28* promoter (SI Appendix, Fig. S2C and D). Unlike the punctal morphology of LMP-1::GFP-labeled lysosomes in the hypoderm of wild-type animals, lysosomes in *tmem-39* mutants formed more tubular structures (Fig. 2F). Time-lapse confocal imaging revealed that the tubular lysosomes were more static compared with wild-type lysosomes (Fig. 2G and Movies S1 and S2). We conclude that TMEM-39 regulates lysosomal LAMP1 abundance and is important for maintaining normal morphology and motility of lysosomes.

Given the evolutionary conservation and homology of TMEM39 proteins between worms and humans, we next examined human TMEM39A protein localization in HEK 293T cells. Confocal imaging of cells transiently transfected with *mCherry::TMEM39A* and GFP-tagged reporters for various subcellular compartments revealed that TMEM39A partially colocalized with ER, Golgi apparatus, and Rab7-labeled late-endosomes/lysosomes (SI Appendix, Fig. S2E–G). To determine if knocking out TMEM39A disrupts any of the subcellular organelles, we generated TMEM39A knockout (KO) HEK 293T cells by CRISPR-Cas9 and examined subcellular organelle morphology and localization by confocal microscopy. We found that disruption of TMEM39A led to redistribution of lysosomes from perinuclear regions to cell periphery, while other subcellular organelles, including ER, Golgi apparatus, and mitochondria, remained largely unaffected (Fig. 2H and SI Appendix, Fig. S2H). The peripheral distribution of lysosomes in TMEM39A KO cells was rescued by a wild-type *mCherry::TMEM39A* transgene (Fig. 2I). Together, our findings indicate that mammalian TMEM39A has an evolutionarily conserved role in regulating lysosome distribution.

Lysosome distribution is highly regulated, and its positioning affects lysosomal functions in cellular material degradation, nutrient sensing, and autophagy signaling (12, 13). To determine if

redistribution of lysosomes in TMEM39A KO cells and animals is associated with changes in lysosomal function, we examined lysosomal luminal enzyme activity in wild-type and mutant cells or worms using the Magic Red Cathepsin B assay (14). We did not detect a noticeable difference in Magic Red staining between wild-type and *tmem-39* mutant worms or wild type and TMEM39A KO cells, suggesting that the degradative activity of lysosomal enzymes remains largely unimpaired in the absence of TMEM39A (SI Appendix, Fig. S2I–L). In addition to its degradative function, the cytosolic surface of lysosomes also serves as an important hub for nutrient and mTOR signaling (15–19). Using a GFP reporter under the promoter of *tts-1*, which is activated by mTOR inhibition or starvation through the *C. elegans* transcription factor HLH-30 (ref. 20), we found that disruption of TMEM-39 in *C. elegans* strongly induced *tts-1* reporter expression (Fig. 3A and B). This indicates that mTOR signaling in *tmem-39* mutant animals is reduced under nonstarved conditions in *C. elegans*.

The mTOR complex phosphorylates a variety of downstream substrates, including the transcription factor EB (TFEB), a master regulator of lysosome biogenesis (21). Starvation or pharmaceutical inhibition of mTOR induces dephosphorylation of TFEB, which then translocates to nucleus and activates lysosome biogenesis-related gene transcription (22). Similarly, the *C. elegans* TFEB ortholog HLH-30 responds to nutritional status and activates autophagic gene expression upon starvation (23, 24). We reasoned that HLH-30 might be activated in mutant animals lacking TMEM-39 and examined HLH-30 expression in wild-type and *tmem-39* mutant animals using a stably integrated *hlh-30::gfp* transgene. The HLH-30::GFP was primarily detected in the cytosol of neurons, muscles, and intestinal and hypodermal cells of wild-type animals (Fig. 3C). By contrast, HLH-30::GFP accumulated in nuclei of neuronal and nonneuronal cells of *tmem-39* mutants, indicating activation of the transcription factor (Fig. 3C and E). We then asked if HLH-30 is required for the strong *tts-1_p::gfp* mTOR reporter expression induced by *tmem-39* disruption. Feeding *tmem-39* mutant animals with *hlh-30* small interfering RNA (siRNA)-expressing bacteria, but not with *mxl-3* (another bHLH factor) siRNA-expressing or vector bacteria, nearly abolished the mTOR reporter expression (Fig. 3A and B). Therefore, we conclude that *tmem-39* disruption in *C. elegans* likely inhibits lysosome-associated mTOR signaling, which induces strong *tts-1_p::gfp* mTOR reporter expression through nuclear translocation and activation of HLH-30/TFEB.

Another cellular program regulated by mTOR signaling is autophagy, which is enhanced upon starvation or pharmaceutical inhibition of mTOR complex (17, 18). To find out if cells lacking TMEM39A have altered autophagy, we examined expression

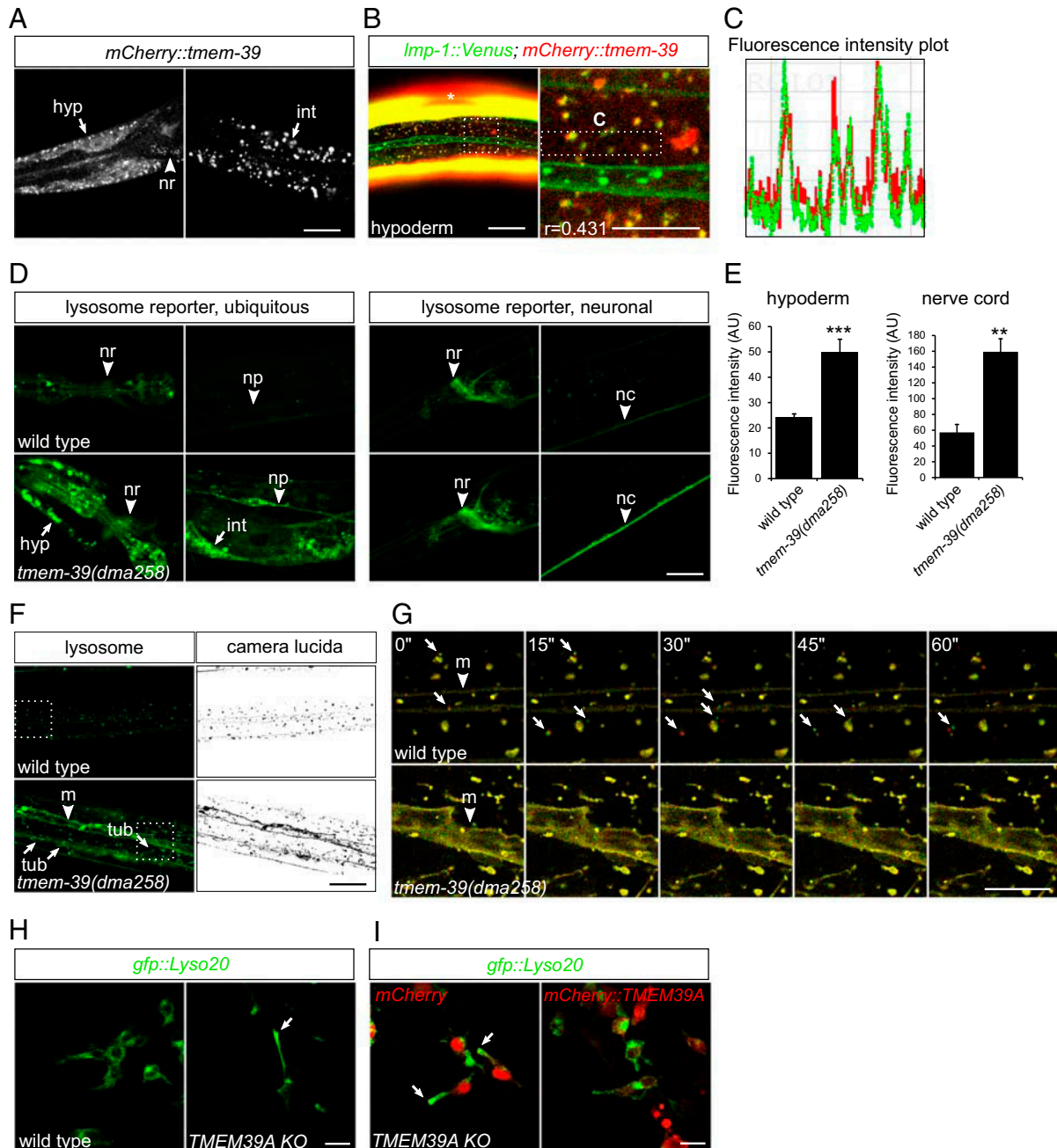


Fig. 2. TMEM-39 is associated with lysosomes and regulates lysosome positioning and morphology. (A) Confocal images showing expression and punctate distribution of the *C. elegans* mCherry::TMEM-39 fusion protein, detected in hypoderm (hyp), nerve ring (nr), and intestine (int). (Scale bar, 20 μ m.) (B) Confocal images showing partial colocalization of mCherry::TMEM-39 with lysosomal marker LMP-1::Venus in *C. elegans* hypoderm. Asterisks denote the expression of muscle GFP or mCherry from coinjection plasmids. The Pearson's correlation coefficient ($r = 0.431$) indicates moderate association of TMEM-39 with lysosomes. (Scale bar, 20 μ m [B, Left] and 10 μ m [B, Right].) (C) Intensity plot of mCherry::TMEM-39 and LMP-1::Venus fluorescence in the boxed region in B. (D) Ubiquitous lysosome reporter *rpl-28_p::Imp-1::Venus* is strongly up-regulated in *tmem-39* mutant animals, evident in hypoderm, intestine, nerve ring, and nerve cord (nc). The lysosome reporter induction in nerve ring is more modest compared with that in hypoderm or intestine, based on a neuronal lysosomal reporter *rab-3_p::Imp-1::gfp*. (Scale bar, 20 μ m.) (E) Quantification of lysosome reporter up-regulation in *tmem-39* mutants. Data represent mean \pm SEM. ** $P < 0.01$; *** $P < 0.001$. AU, arbitrary units. (F) The lysosome LMP-1::Venus reporter labels cytoplasmic or endosomal membrane (m; arrowhead) more strongly in *tmem-39* mutant hypoderm than does in the wild type and tends to form more tubular structures (tub; arrows). Dashed boxes represent areas enlarged in G. (Scale bar, 20 μ m.) (G) Time-lapse confocal images showing lysosome dynamics. Two consecutive time-lapse images are pseudocolored with red and green and are superposed to reveal mobile (nonyellow) lysosomes (arrows). Compared with lysosomes in wild type that are mobile, tubular lysosomes in *tmem-39* mutants are relatively static. The putative cytoplasmic or endosomal membrane (m) is labeled by arrowheads. (Scale bar, 10 μ m.) (H) Confocal images showing redistribution of lysosomes from perinuclear regions to cell periphery in *TMEM39A* KO HEK 293T cells (arrow). (Scale bar, 20 μ m.) (I) Confocal images showing the restoration of perinuclear localization of lysosomes in *TMEM39A* KO cells transfected with mCherry::TMEM39A, but not with mCherry plasmid (arrows). (Scale bar, 20 μ m.)

patterns of mCherry::gfp::LC3B, a tandem-fluorescence-tagged autophagosome protein in wild-type and *TMEM39A* KO HEK 293T cells (25). LC3B was primarily diffused in the cytoplasm of

wild-type cells, and 1 h of starvation of the cells in the serum-free Earle's Balanced Salt Solution medium induced strong formation of LC3B puncta that was indicative of autophagosome

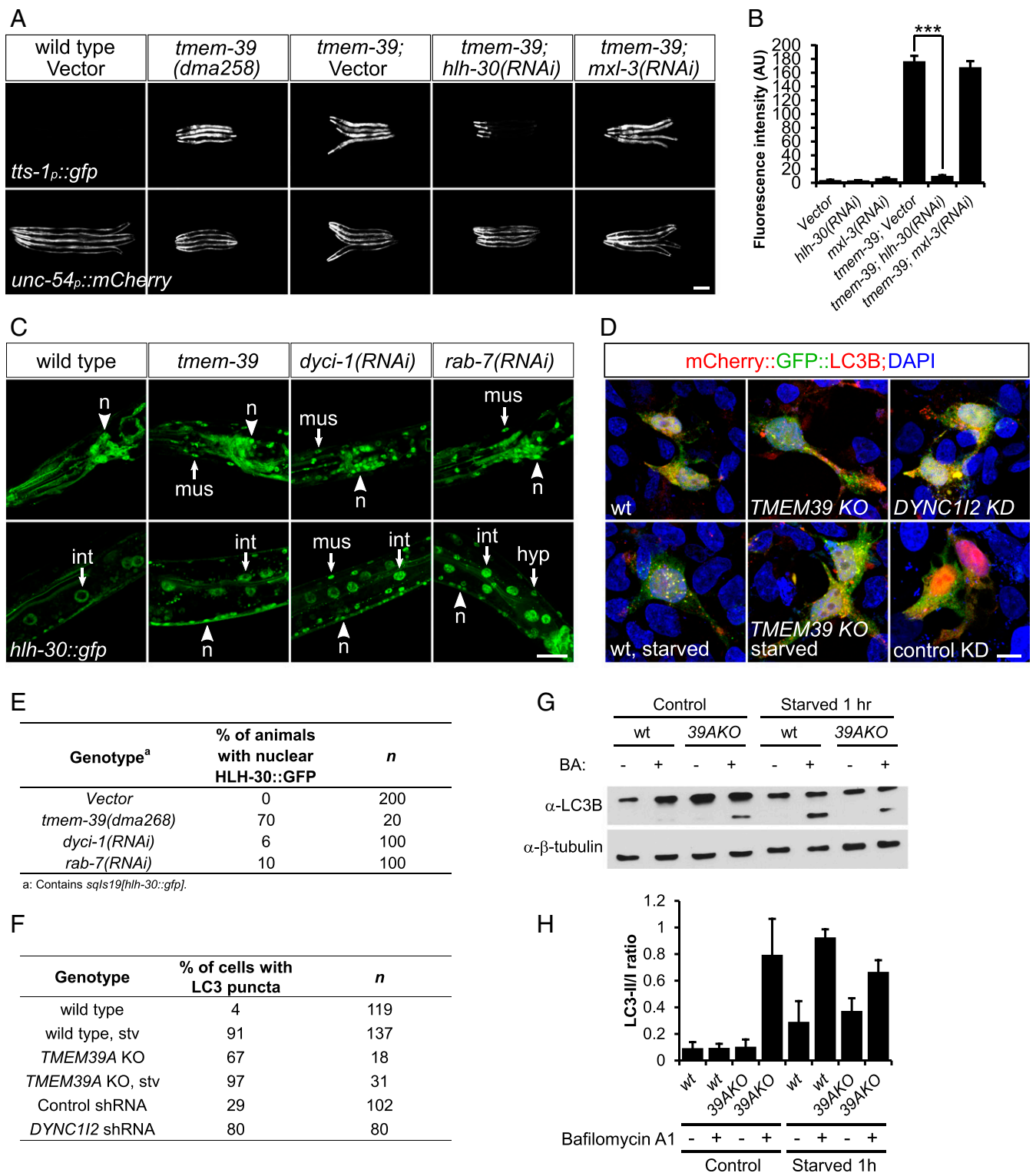


Fig. 3. *TMEM-39* regulates mTOR signaling through the master regulator of lysosome biogenesis HLH-30/TFEB. (A) Expression of an mTOR reporter *tts-1_p::gfp* is strongly up-regulated in *tmem-39* mutant animals, while disruption of *hlh-30/TFEB* by RNAi abolishes *tts-1_p::gfp* up-regulation. *unc-54_p::mCherry*, coinjection reporter; *mxl-3*, bHLH transcription factor that antagonizes HLH-30 function in lipolysis upon starvation. (Scale bar, 100 μ m.) (B) Quantification of *tts-1_p::gfp* fluorescence intensity in A. Data represent mean \pm SEM. ****P* < 0.001. AU, arbitrary units. (C) Confocal images showing nuclear localization of HLH-30 in neurons (n), hypoderm (hyp), intestine (int), and muscle (mus) cells of animals deficient in *tmem-39*, *dyci-1*, or *rab-7*, but not in wild type. (Scale bar, 20 μ m.) (D) Representative confocal images showing puncta formation of mCherry::GFP::LC3 in cells deficient in *TMEM39A*, *DYNC112* or following 1 h of starvation. The presence of more red, but not yellow, LC3 puncta in *TMEM39A* KO cells indicates lysosome-mediated quenching of the GFP signal. (Scale bar, 10 μ m.) (E) Quantification of HLH-30::GFP nuclear localization in C. (F) Quantification of mCherry::GFP::LC3 puncta formation in D. (G) Western blot analysis showing up-regulation of LC3-II/I ratio in nonstarved *TMEM39A* KO compared with wild-type HEK 293T cells, indicating enhanced autophagosome formation. BA, bafilomycin A1. (H) Quantification of the Western blot results. Data represent mean \pm SEM. wt, wild type.

formation (Fig. 3 D and F). Strikingly, in cells lacking TMEM39A, LC3B clustered in puncta even under nonstarved conditions, and starvation further enhanced puncta formation (Fig. 3 D and F). Moreover, Western blot analysis using an anti-LC3B antibody detected enhanced accumulation of LC3-II in nonstarved *TMEM39A* KO cells, indicating enhanced autophagosome formation in the absence of TMEM39A (Fig. 3 G and H). Consistent with the observations, phosphorylation of Ser15 on Beclin-1, a molecular event that is indicative of amino acid starvation and mTOR inhibition and required for autophagy initiation (26), was enhanced in nonstarved *TMEM39A* KO cells (SI Appendix, Fig. S3C). Together, these findings support an important role of TMEM39A in maintaining normal mTOR signaling and suppressing autophagy initiation, and disrupting TMEM39A enhances autophagy, at least in HEK 293T cells.

TMEM39A is predicted to contain nine transmembrane domains and two large loops (https://harrier.nagahama-i-bio.ac.jp/sosui/sosui_submit.html), with the second loop highly conserved from *C. elegans* to human (Fig. 1A). To identify molecular mechanisms of how TMEM39A regulates lysosome dynamics, we performed yeast two-hybrid (Y2H) screens to identify proteins that interact with the second loop of TMEM39A. From a normalized universal human cDNA library, we identified 20 candidate interacting proteins, including the dynein intermediate light chain DYNC1I2 (Fig. 4A and SI Appendix, Table S1). The interaction appeared to require the conserved tryptophan-serine (WS) residues within the loop, as mutations of these amino acids to alanines abolished the interaction (Fig. 4A). We further confirmed the interaction between TMEM39A and DYNC1I2 in mammalian cells using coimmunoprecipitation (Fig. 4B). To test whether the interaction is functionally important *in vivo*, we generated CRISPR-knockin *C. elegans* strains, in which the conserved WS residues were replaced by alanines. Strains that carry such mutations in *tmem-39* showed markedly stronger *hsp-4_p::gfp* ER stress-reporter expression and PVD neuronal morphological defects, closely resembling the abnormalities seen in *tmem-39* null mutant animals (Fig. 4 C and D).

Dynein is a family of cytoskeletal motor proteins responsible for cargo trafficking toward the minus end of microtubules and can be recruited to lysosomes via dynein light intermediate chain DYNC1LI1 and Rab-interacting lysosomal protein RILP (27, 28). To find out if dynein functions downstream of TMEM-39 to regulate lysosome distribution and biogenesis, we performed feeding RNAi to disrupt the expression of the sole homolog of dynein intermediate light chain *dyci-1* in *C. elegans*. Knockdown of *dyci-1* led to partial embryonic lethality and growth arrest, consistent with an essential role of the dynein subunit in early embryonic development (29). Strikingly, *dyci-1* deficiency induced a strong increase in mTOR reporter *tts-1_p::GFP* expression, as well as a moderate, but significant, increase in LMP-1::GFP, suggesting a starvation-like status in the RNAi-treated animals (Fig. 4 E and F and SI Appendix, Fig. S3 A and B). Disruption of *hlh-30* by RNAi in the *dyci-1*-deficient animals abolished mTOR reporter *tts-1_p::GFP* up-regulation, indicating that HLH-30 functions downstream of both TMEM-39 and DYCI-1 to activate mTOR reporter expression (Fig. 4 E and F). We further tested if RAB-7, RILP-1, or dynactin DNC-1 are required for regulating mTOR reporter expression and/or lysosome biogenesis, given their involvement in dynein–dynactin complex formation and recruitment of the complex to lysosomes (30). We found that disruption of the expression of *rab-7* by RNAi induced a marked increase in the mTOR reporter *tts-1_p::GFP* and lysosomal LMP-1::GFP expression, while deficiency in *rilp-1* or *dnc-1* caused only a moderate effect (Fig. 4 E and F and SI Appendix, Fig. S3 A and B).

We next asked if the role of dynein is also conserved in mammalian cells. The expression of *DYNC1I2* in HEK 293T cells

could be efficiently knocked down by using small-hairpin RNA (shRNA)-expressing plasmids (SI Appendix, Fig. S3 D and E). We found that disruption of *DYNC1I2* induced lysosome redistribution to cell periphery in $51.2 \pm 1.8\%$ of the transfected cells, similar to that observed in *TMEM39A* KO cells ($60.7 \pm 7.1\%$) (Fig. 4 G and H), while knocking down *DYNC1I2* in *TMEM39A* KO cells did not further enhance lysosome redistribution ($53.3 \pm 2.8\%$) (Fig. 4H). In addition, expression of shRNA against *DYNC1I2*, but not GFP control shRNA plasmid, induced strong puncta formation of autophagosome protein LC3B, indicating that *DYNC1I2* and *TMEM39A* function similarly to regulate lysosome positioning and autophagy initiation (Fig. 3 D and F). Interestingly, mCherry::*DYNC1I2* was redistributed to distal cell periphery, similar to that seen with lysosome reporter in *TMEM39A* KO cells, suggesting that TMEM39A promotes centripetal locomotion of both dynein and lysosomes (Fig. 4 I and J). Taken together, our findings support a model in which TMEM39A regulates lysosome positioning and associated signaling, at least in part through its interaction with the dynein intermediate light chain DYNC1I2 (Fig. 4K).

Discussion

TMEM39A is an evolutionarily conserved protein in which SNPs were previously identified in several human genetic studies with enhanced risks to MS and lupus. A recent study revealed a critical role of TMEM39A in regulating autophagy initiation (10). Here, we describe a key role of TMEM39A in regulating lysosome dynamics and mTOR signaling, in addition to autophagy initiation. We have also identified the dynein intermediate light chain DYNC1I2 as a mechanistic link between TMEM39A and downstream cellular processes. As MS is characterized by demyelination of neurons, owing to the immune-system attack of oligodendrocytes, the broad expression of TMEM39A and function in lysosomes indicate that it may contribute to MS pathology via lysosomal regulation in both a cell-autonomous (neurons) and/or nonautonomous manner (immune cells).

As an essential organelle in which antigens undergo processing and presentation through the major histocompatibility complex, lysosomes are essential for normal immune response (31). Dysregulation of lysosomes is also connected to pathological cytokine release, linking lysosome dysfunction to tissue destruction and autoimmunity (32, 33). In both SLE and MS, aberrant lysosomal activities have been postulated to underlie accumulation of autoantigens in lysosomes, which leads to aberrant immune responses and pathogenesis of the diseases (34, 35). In addition, dysregulated lysosomes also contribute to impaired cellular material and organelle degradation under conditions of aging, fertilization, and neurodegenerative diseases (18, 28, 36–40). Interestingly, *TMEM39A* RNAi has been shown to affect mitophagy (to degrade mitochondria by autophagy) mediated by the Parkinson's disease protein Parkin (41, 42), and inhibition of mitochondrial ribosomes together with impaired mitochondrial fission or fusion activates HLH-30 and increases *C. elegans* lifespan (43).

TMEM39A has been shown to promote ER-to-Golgi transport by interacting with the COPII complex subunits SEC23/SEC24 (10). From our Y2H screen, we also identified SEC23A as an interacting partner of TMEM39A (SI Appendix, Table S1), supporting an important role of TMEM39A in regulating ER-to-Golgi trafficking. This is also consistent with our observation that disruption of TMEM-39 in *C. elegans* induces strong activation of the ER stress reporter *hsp-4_p::gfp* (Figs. 1D and 4C). Interestingly, the C terminus of mammalian TMEM39A contains a putative COPI (which regulates Golgi-to-ER trafficking)-interacting lysine motif KAN, and the autoimmune risk allele *rs1132200* confers an A-to-T mutation in the lysine motif that is predicted to impair interaction with COPI and possibly causes a loss-of-function of the protein (44). We speculate that TMEM39A

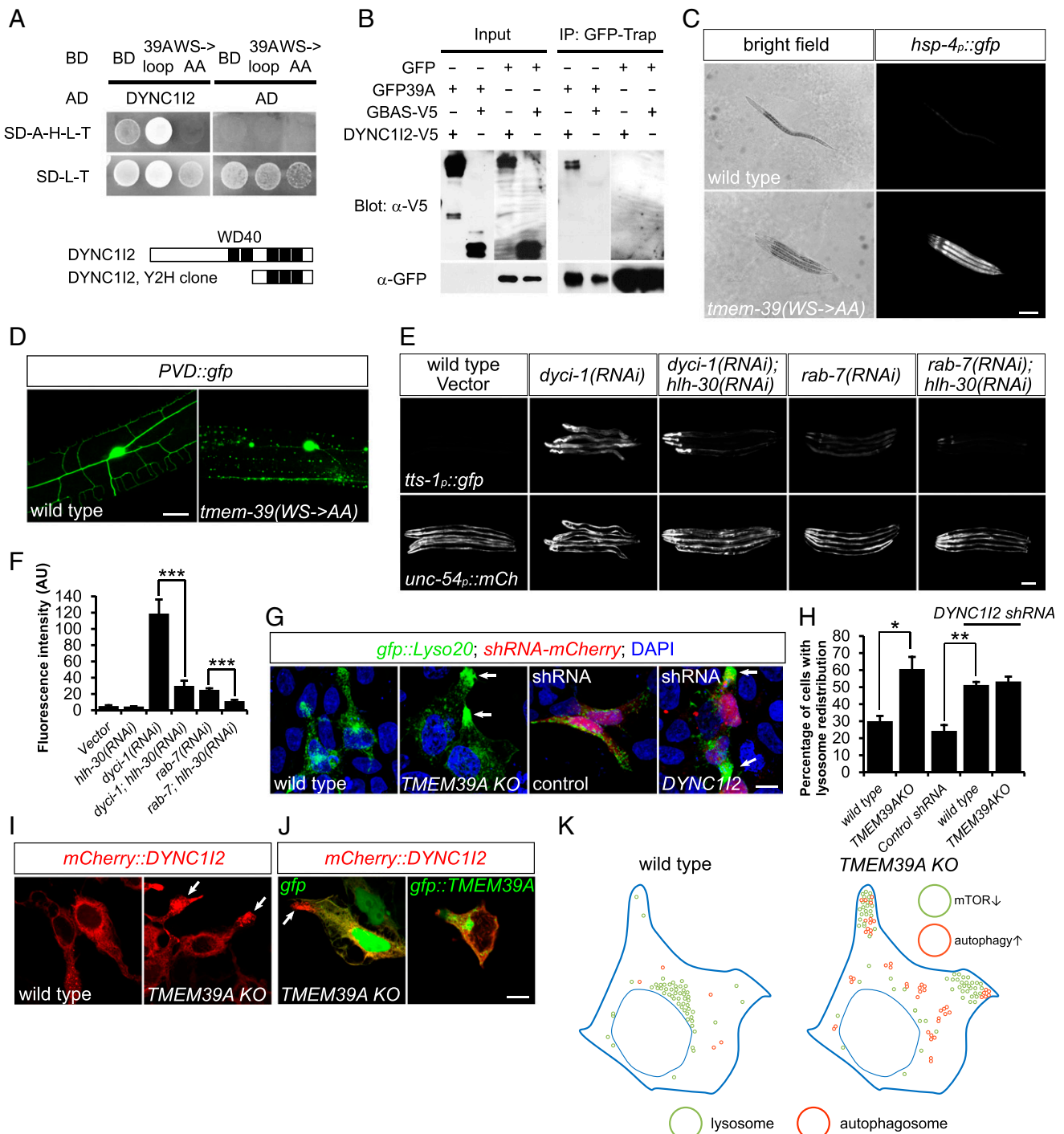


Fig. 4. TMEM39A regulates lysosome distribution and function through dynein intermediate light chain DYNC112. (A) Y2H assays showing the interaction between TMEM39A loop with DYNC112, which is dependent on the conserved WS residues within the TMEM39A loop. Schematic diagrams of the WD domain containing DYNC112 fragment recovered from the Y2H screen are shown. The 39A loop is the second cytosolic loop of TMEM39A. (B) Full-length DYNC112 was coimmunoprecipitated (IP) with GFP::TMEM39A, but not with GFP from transiently transfected HEK 293T cells. GBAS, another candidate protein identified from the Y2H screen, did not coimmunoprecipitate with GFP::TMEM39A. GFP::TMEM39A was detected in GFP-trap fraction, but not in total lysate, likely because of low expression levels. (C) Bright-field and compound fluorescence images showing the dumpy phenotype and elevated *hsp-4p::gfp* ER stress-reporter expression in *tmem-39(WS->AA)* CRISPR-knockin animals. (Scale bar, 100 μ m.) (D) Confocal images showing PVD degeneration-like beading phenotype in *tmem-39(WS->AA)* CRISPR-knockin animals. (Scale bar, 20 μ m.) (E) Disruption of *dyci-1* or *rab-7* by RNAi induces *tts-1p::gfp* mTOR reporter expression, which is abolished by *hlh-30/TFEB* deficiency. (Scale bar, 100 μ m.) (F) Quantification of *tts-1p::gfp* intensity in E. Data represent mean \pm SEM. *** P < 0.001. AU, arbitrary units. (G) Expression of DYNC112-targeting shRNA in HEK 293T cells induces redistribution of lysosomes to cell periphery, similar to that observed in *TMEM39A KO* cells (arrows). (Scale bar, 10 μ m.) (H) Quantification of percentage of cells with lysosome redistribution in G. Data represent mean \pm SEM. * P < 0.05; ** P < 0.01. (I) Confocal images showing abnormal accumulation of mCherry::DYNC112 in cell periphery of *TMEM39A KO* cells (arrows). (J) Expression of *gfp::TMEM39A*, but not *gfp*, rescued mCherry::DYNC112 localization in *TMEM39A KO* cells. (Scale bar, 10 μ m.) (K) Schematic diagram showing a model in which disruption of TMEM39A leads to lysosome redistribution to cell periphery, which affects lysosome-associated mTOR signaling and autophagy initiation.

is involved not only in anterograde (from ER to Golgi), but also in retrograde (from Golgi to ER), trafficking. These observations, together with our findings, support important roles of TMEM39A in regulating proper functions of multiple subcellular organelles, including the ER, lysosome, and autophagosome. Detailed mechanisms of how such multifaceted roles of TMEM39A are executed, coordinated, and regulated warrant further investigation.

To summarize, our results identify a previously uncharacterized function of TMEM39A in lysosome regulation. We propose a broadly evolutionarily conserved role of TMEM39A in promoting proper lysosome positioning and accumulation, which are important for regulation of mTOR signaling, autophagy initiation, and immune-cell responses under a variety of physiological and pathological conditions (28, 32, 36, 45–47).

Materials and Methods

All *C. elegans* strains were handled and maintained at 22 °C, and the N2 Bristol strain was used as the wild type. The *dma258* and *dma268* loss-of-function alleles and *dma308* (WS->AA) knockin allele of *tmem-39* were generated using the CRISPR-Cas9 system via germline transformation. The Ahringer and ORFeome RNAi libraries were used to knock down

target gene expression in *C. elegans*. For mammalian cell experiments, HEK 293T cells were maintained in Dulbecco's Modified Eagle Medium supplemented with 10% fetal bovine serum and were grown at 37 °C with 5% CO₂. The *TMEM39A* KO HEK 293T cells were generated using the CRISPR-Cas9 system via transfection and screening for deletion clones. The Sigma-Aldrich MISSION shRNA was used to knock down target gene expression in HEK 293T cells. Compound images were obtained using either an EVOS inverted microscope (Life Technologies) or a Leica CTR5000 compound microscope (Leica), and confocal images were obtained using a Zeiss LSM 700 or Leica SPE microscope. The processing, quantification and colocalization analysis of images were performed in Fiji software (NIH). Detailed materials and methods are provided in *SI Appendix*.

Data Availability. All study data are included in the article and/or supporting information.

ACKNOWLEDGMENTS. We thank the *Caenorhabditis* Genetics Center and National BioResource Project in Japan for *C. elegans* strains, and Drs. Hong Zhang, Zimu Li, Anthony Shum, Scott Zamvil, and Jorge Oksenberg for helpful discussions. The work was supported by NIH Grant R01GM117461, a Pew Scholar Award, and a Packard Fellowship in Science and Engineering (to D.K.M.).

- G. S. Cooper, M. L. K. Bynum, E. C. Somers, Recent insights in the epidemiology of autoimmune diseases: Improved prevalence estimates and understanding of clustering of diseases. *J. Autoimmun.* **33**, 197–207 (2009).
- M. H. Cheng, M. S. Anderson, Monogenic autoimmunity. *Annu. Rev. Immunol.* **30**, 393–427 (2012).
- M. Gutierrez-Arcelus, S. S. Rich, S. Raychaudhuri, Autoimmune diseases—connecting risk alleles with molecular traits of the immune system. *Nat. Rev. Genet.* **17**, 160–174 (2016).
- Q. Yao *et al.*, Genetic variants in *TMEM39A* gene are associated with autoimmune thyroid diseases. *DNA Cell Biol.* **38**, 1249–1256 (2019).
- J. Varadé *et al.*, Replication study of 10 genes showing evidence for association with multiple sclerosis: Validation of *TMEM39A*, *IL12B* and *CBLB* [correction of *CLBL*] genes. *Mult. Scler.* **18**, 959–965 (2012).
- X. Cai, W. Huang, X. Liu, L. Wang, Y. Jiang, Association of novel polymorphisms in *TMEM39A* gene with systemic lupus erythematosus in a Chinese Han population. *BMC Med. Genet.* **18**, 43 (2017).
- J. L. McCauley, J. P. Hussman; International Multiple Sclerosis Genetics Consortium (IMSGC), Comprehensive follow-up of the first genome-wide association study of multiple sclerosis identifies *KIF21B* and *TMEM39A* as susceptibility loci. *Hum. Mol. Genet.* **19**, 953–962 (2010).
- C. J. Lessard *et al.*; BIOLUPUS Network; GENLES Network, Identification of *IRF8*, *TMEM39A*, and *IKZF3-ZBP2* as susceptibility loci for systemic lupus erythematosus in a large-scale multiracial replication study. *Am. J. Hum. Genet.* **90**, 648–660 (2012).
- J. Park *et al.*, Recognition of transmembrane protein 39A as a tumor-specific marker in brain tumor. *Toxicol. Res.* **33**, 63–69 (2017).
- G. Miao, Y. Zhang, D. Chen, H. Zhang, The ER-localized transmembrane protein *TMEM39A/SUSR2* regulates autophagy by controlling the trafficking of the PtdIns(4)P phosphatase *SAC1*. *Mol. Cell* **77**, 618–632.e5 (2020).
- Z. Zhang *et al.*, Broadly conserved roles of *TMEM131* family proteins in intracellular collagen assembly and secretory cargo trafficking. *Sci. Adv.* **6**, eaay7667 (2020).
- V. I. Korolchuk *et al.*, Lysosomal positioning coordinates cellular nutrient responses. *Nat. Cell Biol.* **13**, 453–460 (2011).
- X. Li *et al.*, A molecular mechanism to regulate lysosome motility for lysosome positioning and tubulation. *Nat. Cell Biol.* **18**, 404–417 (2016).
- J. Chen *et al.*, Metformin extends *C. elegans* lifespan through lysosomal pathway. *eLife* **6**, e31268 (2017).
- J. Chen *et al.*, *KLHL22* activates amino-acid-dependent mTORC1 signalling to promote tumorigenesis and ageing. *Nature* **557**, 585–589 (2018).
- R. L. Wolfson *et al.*, *KICSTOR* recruits *GATOR1* to the lysosome and is necessary for nutrients to regulate mTORC1. *Nature* **543**, 438–442 (2017).
- R. E. Lawrence, R. Zoncu, The lysosome as a cellular centre for signalling, metabolism and quality control. *Nat. Cell Biol.* **21**, 133–142 (2019).
- A. Efeyan, R. Zoncu, D. M. Sabatini, Amino acids and mTORC1: From lysosomes to disease. *Trends Mol. Med.* **18**, 524–533 (2012).
- K. H. Mak *et al.*, Lysosomal nucleotide metabolism regulates ER proteostasis through mTOR signaling. *bioRxiv* [Preprint] (2020) <https://doi.org/10.1101/2020.04.18.048561> (Accessed 10 June 2020).
- S. Nakamura *et al.*, Mondo complexes regulate TFEB via TOR inhibition to promote longevity in response to gonadal signals. *Nat. Commun.* **7**, 10944 (2016).
- C. Settembre *et al.*, TFEB links autophagy to lysosomal biogenesis. *Science* **332**, 1429–1433 (2011).
- C. Settembre *et al.*, A lysosome-to-nucleus signalling mechanism senses and regulates the lysosome via mTOR and TFEB. *EMBO J.* **31**, 1095–1108 (2012).
- L. R. Lapierre *et al.*, The TFEB orthologue *HLH-30* regulates autophagy and modulates longevity in *Caenorhabditis elegans*. *Nat. Commun.* **4**, 2267 (2013).
- E. J. O'Rourke, G. Ruvkun, *MXL-3* and *HLH-30* transcriptionally link lipolysis and autophagy to nutrient availability. *Nat. Cell Biol.* **15**, 668–676 (2013).
- S. Kimura, T. Noda, T. Yoshimori, Dissection of the autophagosome maturation process by a novel reporter protein, tandem fluorescently-tagged LC3. *Autophagy* **3**, 452–460 (2007).
- R. C. Russell *et al.*, *ULK1* induces autophagy by phosphorylating *Beclin-1* and activating *VP53A* lipid kinase. *Nat. Cell Biol.* **15**, 741–750 (2013).
- S. C. Tan, J. Scherer, R. B. Vallee, Recruitment of dynein to late endosomes and lysosomes through light intermediate chains. *Mol. Biol. Cell* **22**, 467–477 (2011).
- P. P. Y. Lie, R. A. Nixon, Lysosome trafficking and signaling in health and neurodegenerative diseases. *Neurobiol. Dis.* **122**, 94–105 (2019).
- S. M. O'Rourke, M. D. Dorfman, J. C. Carter, B. Bowerman, Dynein modifiers in *C. elegans*: Light chains suppress conditional heavy chain mutants. *PLoS Genet.* **3**, e128 (2007).
- J. Pu, C. M. Guardia, T. Keren-Kaplan, J. S. Bonifacio, Mechanisms and functions of lysosome positioning. *J. Cell Sci.* **129**, 4329–4339 (2016).
- E. S. Trombetta, I. Mellman, Cell biology of antigen processing *in vitro* and *in vivo*. *Annu. Rev. Immunol.* **23**, 975–1028 (2005).
- W. Ge, D. Li, Y. Gao, X. Cao, The roles of lysosomes in inflammation and autoimmune diseases. *Int. Rev. Immunol.* **34**, 415–431 (2015).
- D. Rigante, C. Cipolla, U. Basile, F. Gulli, M. C. Savastano, Overview of immune abnormalities in lysosomal storage disorders. *Immunol. Lett.* **188**, 79–85 (2017).
- S. Nagata, R. Hanayama, K. Kawane, Autoimmunity and the clearance of dead cells. *Cell* **140**, 619–630 (2010).
- D. Haves-Zburuf *et al.*, Cathepsins and their endogenous inhibitors cystatins: Expression and modulation in multiple sclerosis. *J. Cell. Mol. Med.* **15**, 2421–2429 (2011).
- A. Fraldi, A. D. Klein, D. L. Medina, C. Settembre, Brain disorders due to lysosomal dysfunction. *Annu. Rev. Neurosci.* **39**, 277–295 (2016).
- Q. Zhou, H. Li, D. Xue, Elimination of paternal mitochondria through the lysosomal degradation pathway in *C. elegans*. *Cell Res.* **21**, 1662–1669 (2011).
- K. A. Bohnert, C. Kenyon, A lysosomal switch triggers proteostasis renewal in the immortal *C. elegans* germ lineage. *Nature* **551**, 629–633 (2017).
- M. Savini, Q. Zhao, M. C. Wang, Lysosomes: Signaling hubs for metabolic sensing and longevity. *Trends Cell Biol.* **29**, 876–887 (2019).
- Y. Sun *et al.*, Lysosome activity is modulated by multiple longevity pathways and is important for lifespan extension in *C. elegans*. *eLife* **9**, e55745 (2020).
- A. Orvedahl *et al.*, Image-based genome-wide siRNA screen identifies selective autophagy factors. *Nature* **480**, 113–117 (2011).
- Q. Tran *et al.*, *TMEM39A* and human diseases: A brief review. *Toxicol. Res.* **33**, 205–209 (2017).
- L. Ali, C. M. Haynes, Mitochondrial translation, dynamics, and lysosomes combine to extend lifespan. *J. Cell Biol.* **219**, 1–2 (2020).
- W. Ma, J. Goldberg, Rules for the recognition of dilysine retrieval motifs by coatamer. *EMBO J.* **32**, 926–937 (2013).
- S. R. Bonam, F. Wang, S. Muller, Lysosomes as a therapeutic target. *Nat. Rev. Drug Discov.* **18**, 923–948 (2019).
- V. I. Korolchuk, D. C. Rubinsztein, Regulation of autophagy by lysosomal positioning. *Autophagy* **7**, 927–928 (2011).
- Y. G. Zhao, H. Zhang, Core autophagy genes and human diseases. *Curr. Opin. Cell Biol.* **61**, 117–125 (2019).