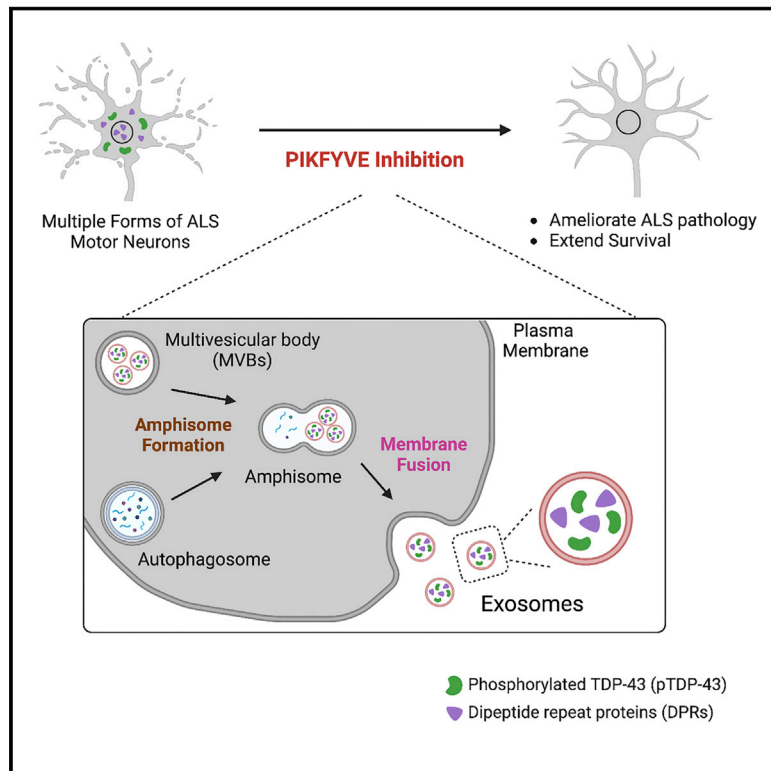


# PIKFYVE inhibition mitigates disease in models of diverse forms of ALS

## Graphical abstract



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## In brief

Pharmacological reduction of PIKFYVE kinase activity activates an unconventional protein clearance mechanism involving exocytosis of aggregation-prone proteins that mitigates disease in diverse models of ALS.

## Highlights

- PIKFYVE inhibition mitigates disease in iPSC and animal models of diverse forms of ALS
- PIKFYVE inhibition clears aggregation-prone proteins via exocytosis
- *Pikfyve* suppression reduces pathology and extends survival of ALS mouse models
- Chronic reduction of PIKFYVE activity is well tolerated and efficacious in TDP-43 mice



Article

# PIKFYVE inhibition mitigates disease in models of diverse forms of ALS

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## SUMMARY

Amyotrophic lateral sclerosis (ALS) is a fatal neurodegenerative disease that results from many diverse genetic causes. Although therapeutics specifically targeting known causal mutations may rescue individual types of ALS, these approaches cannot treat most cases since they have unknown genetic etiology. Thus, there is a pressing need for therapeutic strategies that rescue multiple forms of ALS. Here, we show that pharmacological inhibition of PIKFYVE kinase activates an unconventional protein clearance mechanism involving exocytosis of aggregation-prone proteins. Reducing PIKFYVE activity ameliorates ALS pathology and extends survival of animal models and patient-derived motor neurons representing diverse forms of ALS including *C9ORF72*, *TARDBP*, *FUS*, and sporadic. These findings highlight a potential approach for mitigating ALS pathogenesis that does not require stimulating macroautophagy or the ubiquitin-proteasome system.

## INTRODUCTION

Amyotrophic lateral sclerosis (ALS) is a fast-progressing neurodegenerative disease characterized by motor neuron loss, paralysis, and death within 2–5 years of onset.<sup>1</sup> Similar to other prominent neurodegenerative diseases such as Alzheimer's disease and frontotemporal dementia (FTD), ALS has many diverse genetic etiologies.<sup>2</sup> Since each genetic form is rare and the etiology is unknown for most cases, a central goal is the identification of pathways that can treat multiple forms of ALS.

The accumulation of misfolded proteins can induce neuron death and is a common feature of neurodegenerative diseases.<sup>1,3,4</sup> Canonical proteostasis mechanisms including the ubiquitin-proteasome system and autophagy decline with age, but recent studies suggest that neurons can utilize a third, unconventional protein clearance pathway involving exocytosis of aggregation-prone proteins.<sup>5–7</sup> However, it remains unclear if

stimulating exocytosis of aggregation-prone proteins could be used to treat neurodegenerative diseases.

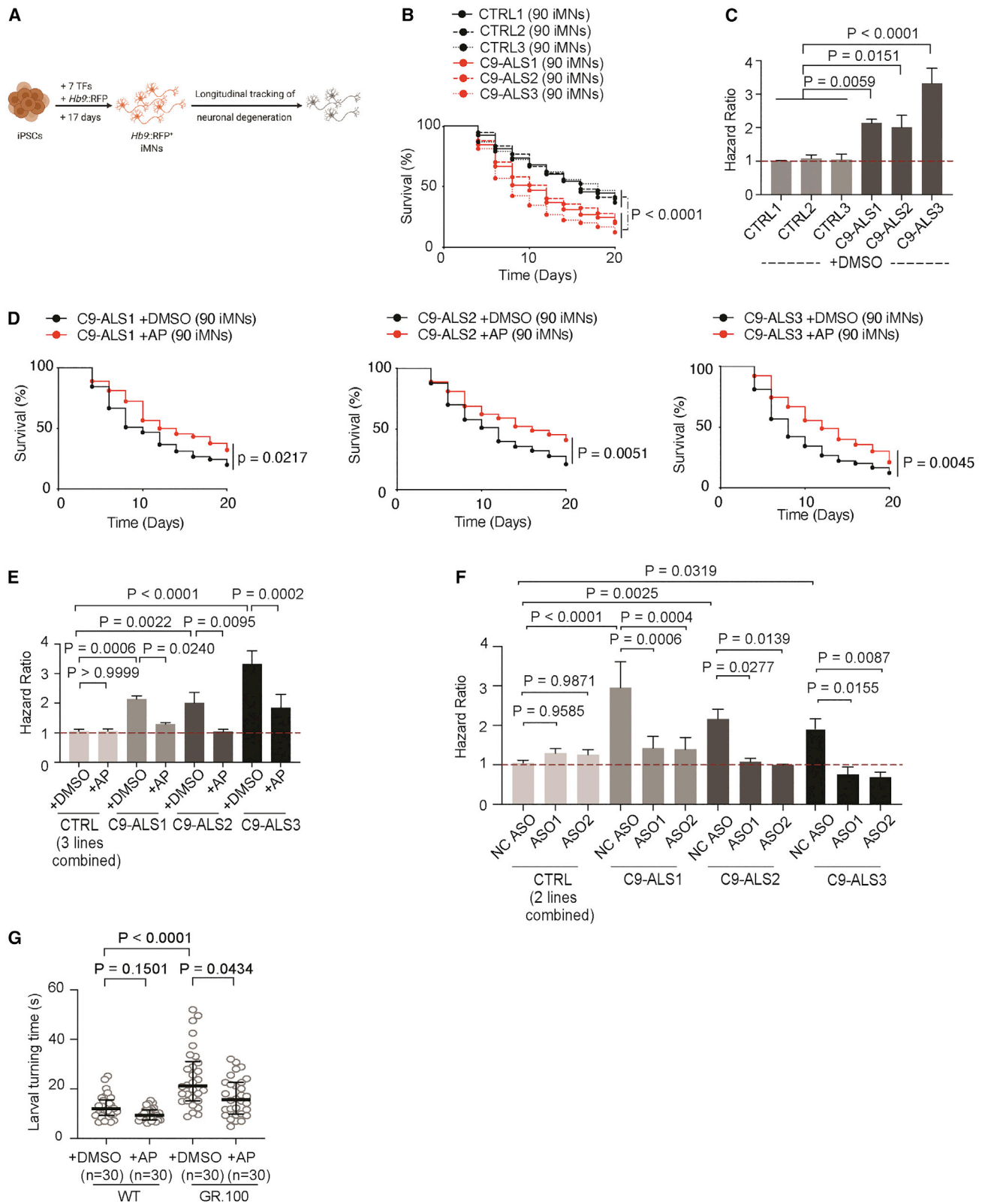
Here, we show that PIKFYVE kinase inhibition is broadly efficacious in models of diverse forms of ALS. PIKFYVE inhibition stimulates the exocytosis of neurotoxic proteins from induced motor neurons (iMNs) of patients with ALS, reduces intracellular pathology, and increases TAR DNA-binding protein 43 (TDP-43) splicing activity. Thus, activating the exocytosis of pathological proteins from neurons mitigates disease in models representing diverse forms of ALS.

## RESULTS

### PIKFYVE inhibition ameliorates *C9ORF72* ALS/FTD neurodegeneration

In a phenotypic screen to identify targets that extend *C9ORF72* ALS/FTD iMN survival, we previously found that PIKFYVE kinase





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inhibitors were among the most efficacious compounds.<sup>8</sup> PIKFYVE converts phosphatidylinositol-3-phosphate (PI3P) to phosphatidylinositol-3,5-bisphosphate (PI(3,5)P<sub>2</sub>), which resides on autophagosomal, endosomal, and lysosomal membranes and regulates vesicle fusion.<sup>9,10</sup> However, how PIKFYVE inhibition ameliorates *C9ORF72* processes, if it can address other forms of ALS, and its efficacy in chronic *in vivo* models remains unknown.

To extend our analyses to additional *C9ORF72* dipeptide repeat proteins (DPRs) and TDP-43 loss-of-function phenotypes, we expressed *Ngn2*, *Isl1*, *Lhx3*, *Ascl1*, *Bm2*, *Myt1l*, *NeuroD1*, and a lentiviral *Hb9::RFP* reporter to convert control and *C9ORF72* ALS/FTD-induced pluripotent stem cells (iPSCs) into *Hb9::RFP*+ iMNs (Figure 1A; Figure S1A).<sup>8,11</sup> Control and patient lines generated iMNs with similar efficiency (Figure S1B; Table S1). *C9ORF72* ALS/FTD iMNs displayed prominent poly(GR)+ and poly(PR)+ punctae compared with isogenic and non-isogenic controls and showed TDP-43 mislocalization into the cytoplasm (Figures S1C–S1I).

Similar to our previous studies, we used longitudinal tracking to determine iMN survival.<sup>8,11–14</sup> When stressed by neurotrophic factor withdrawal, *C9ORF72* ALS/FTD iMNs degenerated faster than controls (Figures 1B and 1C; Figures S1J and S1K). To facilitate quantitative comparisons between multiple groups, we used the Mantel-Haenszel method to assess iMN survival data and calculate a hazard rate or likelihood of neuron death during a specified time interval.<sup>15</sup> We normalized the hazard rate for each group to a control condition to generate a hazard ratio. For *C9ORF72* ALS/FTD iMNs relative to control iMNs, this resulted in a hazard ratio > 1 (Figure 1C). The small-molecule PIKFYVE inhibitor apilimod and multiple PIKFYVE-suppressing antisense oligonucleotides (ASOs) improved *C9ORF72* ALS/FTD iMN survival in a dose-dependent manner (Figures 1D–1F; Figures S1L–S1R). Apilimod also increased motor function and the number of neuromuscular junction (NMJ) active zones in poly(GR)-expressing *Drosophila* larvae (Figure 1G; Figures S1S and S1T).<sup>16</sup>

### PIKFYVE inhibition ameliorates disease pathology in iMNs

RNA-seq analysis of DMSO- and apilimod-treated *C9ORF72* ALS/FTD iMNs revealed differences in autophagy- and vesicle-related gene categories, suggesting that PIKFYVE inhibition might affect proteostasis (Figure S2A; Table S2). Indeed, apilimod reduced poly(GR)+, poly(PR)+, and poly(GA)+ punctae, lowered cytoplasmic TDP-43, increased the nuclear:cytoplas-

mic TDP-43 ratio, reduced *STMN2* cryptic exon inclusion, and increased full-length *STMN2* and *UNC13A* transcripts in *C9ORF72* ALS/FTD iMNs (Figures 2A–2F; Figures S2B–S2D).

### PIKFYVE inhibition stimulates exocytosis of aggregation-prone proteins from iMNs

Since apilimod induced changes in autophagy-related gene sets, we determined if it increased autophagy (Figure S2A). We transduced *C9ORF72* ALS/FTD iMNs with an mRFP-GFP-LC3 construct that labels autophagosomes as GFP+/mRFP+ and lysosomes as mRFP+ only (Figure S3A).<sup>11,17</sup> In the presence of bafilomycin, apilimod increased mRFP+/GFP+ autophagosome production by 3-fold (Figures S3A and S3B).<sup>18</sup> Although apilimod induced gene expression changes resembling transcription factor EB (TFEB) overexpression, it did not increase nuclear TFEB, phosphorylated TFEB, or TFEB target gene activation in *C9ORF72* ALS/FTD iMNs (Figures S3C and S3D; Table S2; Data S1). In addition, apilimod increased the ratio of mRFP+/GFP+ autophagosomes:total mRFP+ vesicles and LC3-II:LC3-I by a slightly greater amount without bafilomycin than with bafilomycin, suggesting that it modestly decreased autophagic flux (Figures S3A, S3B, S3E, and S3F). Moreover, apilimod decreased mRFP+/GFP– autolysosomes without bafilomycin even though LysoSensor Green detected more lysosomes (Figures S3G and S3H). Importantly, bafilomycin did not prevent PIKFYVE inhibition from increasing *C9ORF72* ALS/FTD iMN survival (Figure S3I; Data S1). Thus, PIKFYVE inhibition did not increase macroautophagy, nor was macroautophagy required for apilimod's efficacy.

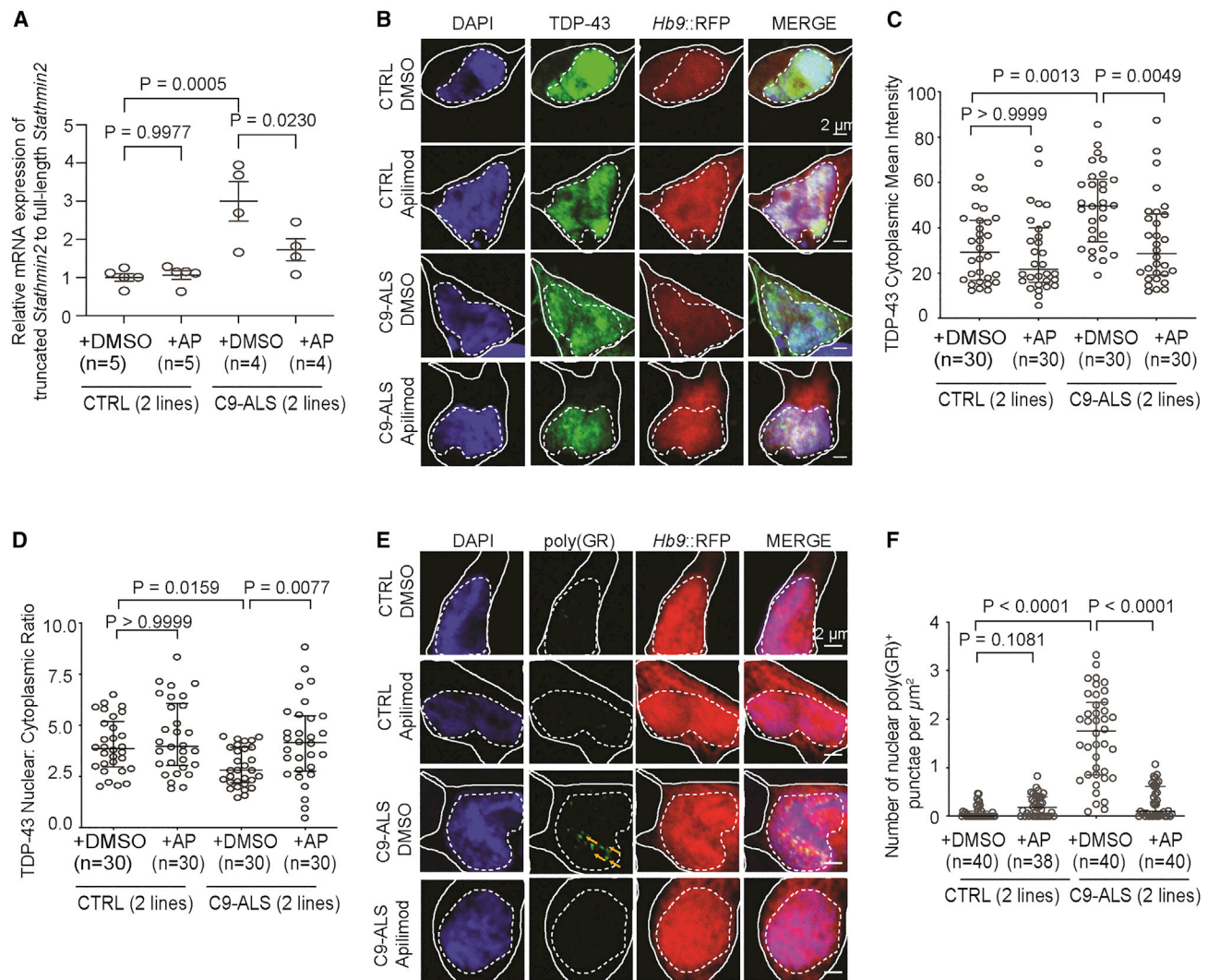
Surprisingly, apilimod did not increase the total number of LC3-mRFP+ vesicles in iMNs, suggesting that autophagosomes might leave the cell or be degraded through a macroautophagy-independent route (Figure S3J). PIKFYVE inhibition can activate an unconventional form of secretion in which autophagosome-associated proteins, cytosolic aggregation-prone proteins, and leaderless proteins are preferentially exocytosed (Figure 3A).<sup>19–21</sup> We harvested extracellular vesicles from iMN supernatants using polyethylene glycol/ultracentrifugation, and after normalizing by total protein content and controlling for differences in neuron death using TUJ1 levels in the cell pellet fraction, immunoblotting showed that apilimod increased levels of the exosomal marker TSG101 in extracellular vesicles from iMN supernatants (Figures 3B and 3C; Figure S3K; Data S1).<sup>22,23</sup> Co-treatment with GW4869, a small-molecule neutral sphingomyelinase 2 inhibitor that blocks exosome release, abolished the ability of apilimod to increase supernatant TSG101

### Figure 1. PIKFYVE inhibition ameliorates *C9ORF72* ALS/FTD neurodegeneration

- Production of *Hb9::RFP*+ iMNs and survival tracking by longitudinal microscopy.
- iMN survival for three control (CTRL) and three *C9ORF72* ALS/FTD lines. Log rank test.
- Hazard ratios for (B). Mean of independent iMN conversions ± SEM. One-way ANOVA.
- Survival of *C9*-ALS/FTD iMNs (three lines) in DMSO/apilimod (AP). Log rank test.
- Hazard ratios for CTRL (three lines in aggregate) and *C9*-ALS/FTD iMNs treated with DMSO/apilimod. Mean of independent iMN conversions ± SEM. One-way ANOVA.
- Hazard ratios from CTRL (two lines in aggregate) and *C9*-ALS/FTD iMNs treated with negative control (NC) ASO, PIKFYVE ASO1, or PIKFYVE ASO2. Mean of independent iMN conversions ± SEM. One-way ANOVA.
- Locomotor function in *Drosophila* larvae overexpressing *C9ORF72* GR.100 with/without apilimod treatment. n = 30 larvae/group. Kruskal-Wallis test. Mean ± SEM.

See also Figure S1.





**Figure 2. PIKFYVE inhibition ameliorates disease pathology in C9ORF72 ALS/FTD iMNs**

(A) Truncated *Stathmin2* RNA levels (normalized to *MAP2*) relative to full-length *Stathmin2* in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = independent iMN conversions. One-way ANOVA. Mean  $\pm$  SEM.

(B–D) Immunostaining/quantification of total TDP-43 in CTRL/C9-ALS/FTD iMNs with DMSO/apilimod treatment. Dots = (C) the mean cytoplasmic TDP-43 intensity or (D) the average nuclear:cytoplasmic TDP-43 ratio in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.

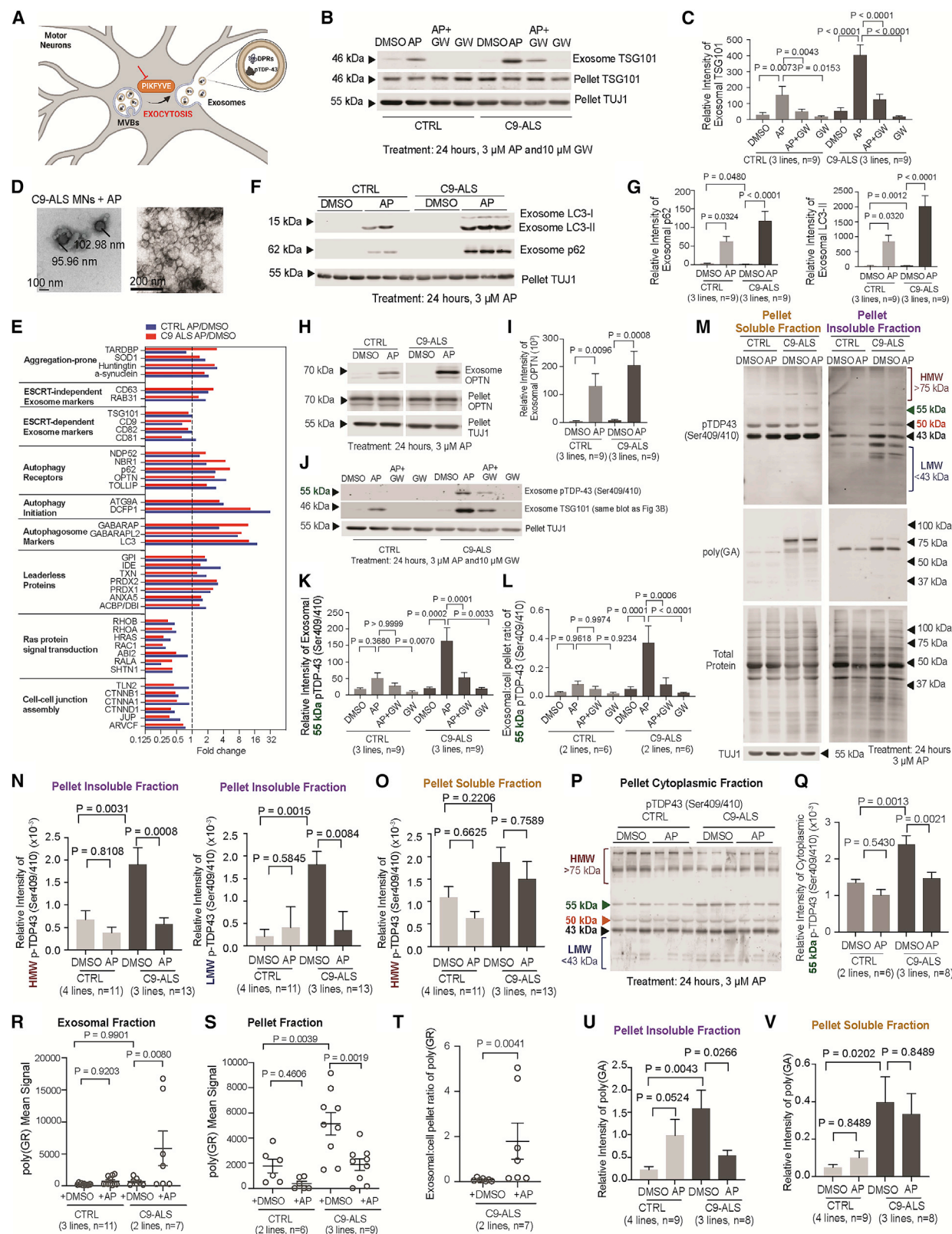
(E and F) Immunostaining/quantification of endogenous poly(GR)+ punctae in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of nuclear poly(GR)+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively. See also Figure S2.

levels (Figures 3B and 3C; Data S1).<sup>6,24</sup> To visualize exosome release, we overexpressed GFP in iMNs, isolated and labeled exosomes with carboxyfluorescein succinimidyl ester (CFSE) far red dye, and performed flow cytometry.<sup>25</sup> Apilimod increased the number of GFP+/CFSE+ particles in the iMN supernatant (Figure S3L). Electron microscopy confirmed that the particles secreted by C9ORF72 ALS/FTD iMNs upon apilimod treatment were exosomes of  $\sim 100$  nm in diameter (Figure 3D).<sup>26</sup>

Mass spectrometry and immunoblotting showed that apilimod-induced exosomes showed enrichment of autophagosome-associated proteins and aggregation-prone proteins

linked to neurodegenerative disease while showing de-enrichment for Ras signaling and cell-cell junction proteins (Figures 3E–3I; Figures S3K–S3N; Table S3). Apilimod treatment increased the exosome:cell pellet ratios of TSG101 and the autophagosome-associated protein optineurin (OPTN) to  $\sim 2:1$  and  $\sim 0.4:1$ , respectively, suggesting that large fractions of these proteins were exocytosed (Figures 3B and 3H; Figure S3O; Data S1). Importantly, apilimod increased secretion of pTDP-43 (Ser409/410) from C9ORF72 ALS/FTD iMNs but not controls (Figures 3J and 3K; Data S1). Apilimod also increased total TDP-43 release from C9ORF72 ALS/FTD iMNs to a greater





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extent than from control iMNs (Figure S3P; Data S1). Co-administration of GW4869 severely reduced pTDP-43 and total TDP-43 release, confirming that their secretion was exocytosis dependent (Figures 3J and 3K; Figure S3P; Data S1). Apilimod treatment resulted in a pTDP-43 exosome:cell pellet ratio of  $\sim 0.4:1$ , a 10-fold increase over the DMSO-treated condition, and reduced insoluble high-molecular weight (HMW) and low-molecular weight (LMW) pTDP-43 species in *C9ORF72* ALS/FTD iMNs (Figures 3L–3N; Figure S3Q; Data S1). PIKFYVE inhibition did not reduce soluble pTDP-43 in whole-cell *C9ORF72* ALS/FTD iMN lysates or cytoplasmic extracts that were verified for purity using fibrillarin and HSP90, except for a 55-kDa pTDP-43 species in cytoplasmic extracts (Figures 3M and 3O–3Q; Figures S3R–S3T; Data S1). Apilimod did not significantly alter soluble or insoluble total TDP-43 in *C9ORF72* ALS/FTD iMNs, indicating that it caused the largest percent reduction in insoluble pTDP-43 rather than in soluble pTDP-43 or total TDP-43 (Figures S3U and S3V; Data S1). Thus, PIKFYVE inhibition triggers the secretion and clearance of insoluble pTDP-43 from *C9ORF72* ALS/FTD iMNs.

Our previously validated ELISA assay showed that apilimod increased poly(GR) in exosomes from *C9ORF72* ALS/FTD but not control iMNs (Figure 3R).<sup>27,28</sup> Apilimod increased the poly(GR) exosomal marker:pellet ratio to  $\sim 2:1$  and reduced poly(GR) to background levels in *C9ORF72* ALS/FTD iMNs (Figures 3R–3T). Apilimod also reduced cellular levels of insoluble but not those of soluble poly(GA), suggesting that it caused the clearance of oligomeric/aggregated poly(GA) (Figures 3M, 3U, and 3V; Data S1). Thus, proteins that drive ALS pathology are substrates for PIKFYVE inhibition-induced exocytosis in iMNs.

### PIKFYVE inhibition clears pTDP-43 through amphisome and multivesicular body exocytosis

While the exocytosis inhibitor GW4869 did not alter control or *C9ORF72* ALS/FTD iMN survival on its own, it blocked the ability

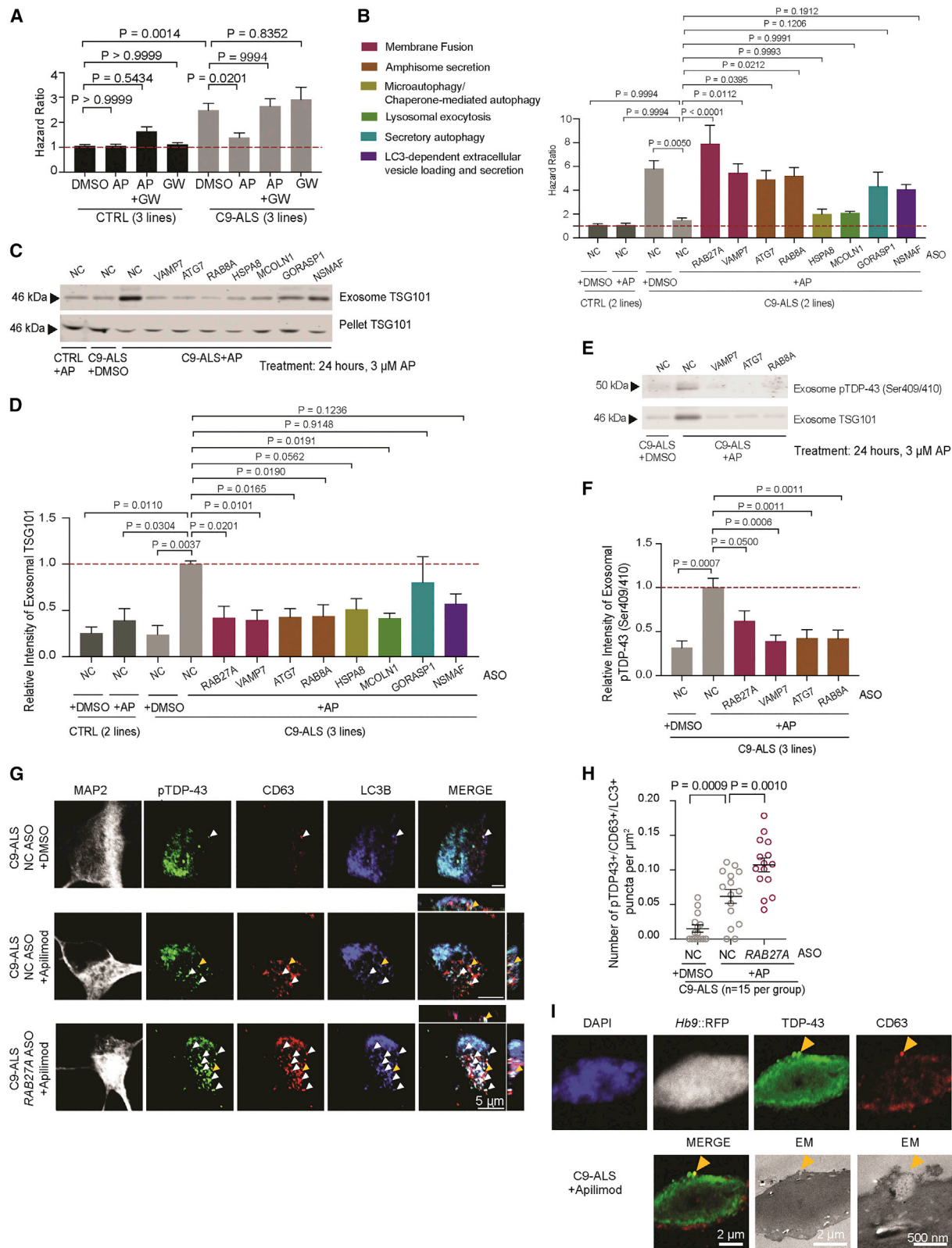
of apilimod to increase *C9ORF72* ALS/FTD iMN survival (Figure 4A; Figure S4A). We used ASOs to selectively suppress different secretory and protein clearance pathways in *C9ORF72* ALS/FTD iMNs to determine which were required for apilimod's efficacy. We suppressed the genes encoding VAMP7 and RAB27A, which are critical for plasma membrane fusion of multivesicular bodies and amphisomes, ATG7 and RAB8A, which promote amphisome exocytosis by stimulating autophagosome biogenesis and fusion with the plasma membrane, respectively, HSPA8, which is required for microautophagy and chaperone-mediated autophagy, MCOLN1, a protein critical for lysosomal exocytosis, GORASP1, an important component of the secretory autophagy pathway, and neutral sphingomyelinase activation associated factor (NSMAF), which is required for LC3-dependent extracellular vesicle loading and secretion (Figure S4B).<sup>29–39</sup> Suppression of VAMP7, RAB27A, ATG7, and RAB8A prevented apilimod from increasing *C9ORF72* ALS/FTD iMN survival and reduced apilimod-induced pTDP-43 and TSG101+ exosome secretion from *C9ORF72* ALS/FTD iMNs, suggesting that amphisome and multivesicular body exocytosis are most critical for the efficacy of PIKFYVE inhibition (Figures 4B–4F; Figures S4C and S4D; Data S1).

To confirm these results, we performed immunofluorescence and electron microscopy. When we blocked exocytosis by suppressing RAB27A, PIKFYVE inhibition increased the total number of LC3+ vesicles in *C9ORF72* ALS/FTD iMNs (Figure S4E). Apilimod increased LC3+/LAMP1+ and LC3+/CD63+ vesicles in *C9ORF72* ALS/FTD iMNs, reflecting more autophagosome fusion with late endosomes/multivesicular bodies to form amphisomes (Figures S4F and S4G). RAB27A suppression further increased the number of LC3+/CD63+ vesicles within apilimod-treated but not vehicle-treated *C9ORF72* ALS/FTD iMNs, suggesting that PIKFYVE inhibition normally caused secretion of these vesicles (Figure S4G). Apilimod increased RAB7+ late

### Figure 3. PIKFYVE inhibition increases exocytosis of aggregation-prone proteins from iMNs

(A) PIKFYVE's putative role in the exocytosis of aggregation-prone proteins.  
(B and C) Immunoblots/quantification of TSG101 in exosomal or pellet fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO, apilimod, apilimod + GW4869, or GW4869. Pellet TUJ1 controlled for cell death. One-way ANOVA. Mean  $\pm$  SEM.  
(D) Electron microscopic images of secreted vesicles in the exosomal fraction of apilimod-treated C9-ALS/FTD iMN cultures.  
(E) Fold-change of selected protein levels in exosomal fractions after apilimod treatment versus DMSO from CTRL/C9-ALS/FTD iMNs.  
(F and G) Immunoblots/quantification of LC3 and p62 in the exosomal fraction from CTRL/C9-ALS/FTD iMN cultures treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.  
(H and I) Immunoblots/quantification of OPTN in exosomal and pellet fractions from C9-ALS/FTD iMNs treated with DMSO/apilimod. Unpaired t test. Mean  $\pm$  SEM.  
(J–L) Immunoblots/quantification of pTDP-43 in exosomal and pellet fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO, apilimod, apilimod + GW4869, or GW4869. The same blot from Figure 3B was used to facilitate comparisons between markers. Exosome TSG101 is the same as Figure 3B. (K) Exosomal pTDP-43 levels. (L) pTDP-43 exosome:cell pellet ratio for CTRL/C9-ALS/FTD iMNs. One-way ANOVA. Mean  $\pm$  SEM.  
(M) pTDP43 and poly(GA) immunoblots on soluble/insoluble fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod.  
(N and O) Quantification of high-molecular weight (HMW, >75 kDa) and low-molecular weight (LMW, <43 kDa) pTDP-43 in the pellet insoluble (N) or soluble (O) fraction from (M). For HMW, one-way ANOVA. Mean  $\pm$  SEM. For LMW, Kruskal-Wallis test, median  $\pm$  interquartile range (N). One-way ANOVA, mean  $\pm$  SEM (O).  
(P and Q) Immunoblots/quantification of pTDP-43 (55 kDa) in cytoplasmic fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.  
(R and S) Immunoassay of poly(GR) levels in exosomal (R) or cell pellet (S) fractions of CTRL/C9-ALS/FTD iMNs. Gray circles = one biological replicate. One-way ANOVA. Mean  $\pm$  SEM.  
(T) Ratio of poly(GR) in exosome:cell pellet fractions of CTRL/C9-ALS/FTD iMNs. Mann-Whitney test. Mean  $\pm$  SEM.  
(U and V) Quantification of poly(GA) in the pellet insoluble (U) and soluble (V) fraction from (M). Values = relative intensity of insoluble poly(GA) (37–100 kDa) normalized to total protein. One-way ANOVA. Mean  $\pm$  SEM.  
See also Figure S3.





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endosomes in *C9ORF72* ALS/FTD iMNs, which possibly contributed to the elevated amphisome formation (Figure S4H). Increased amphisome formation did not result from fewer available primary lysosomes since PIKFYVE inhibition slightly increased the number of small, primary lysosomes and decreased the abundance of large, secondary lysosomes that had been likely fused with endosomes (Figure S4I). Lysosomes also possess high LAMP1 levels, and the LC3/LAMP1 co-labeling results could, in principle, reflect increased autophagosome-lysosome fusion. However, our experiments with the LC3-RFP-GFP construct showed that apilimod decreased the number of LC3-RFP+/GFP– autolysosomes (Figure S3G). Thus, PIKFYVE inhibition increased amphisome formation and promoted their secretion.

We next determined if amphisomes mediated pTDP-43 clearance. Indeed, apilimod increased the number of pTDP-43+/LC3+/CD63+ vesicles in *C9ORF72* ALS/FTD iMNs (Figures 4G and 4H). Overlaying immunofluorescence and electron microscopy imaging enabled direct visualization of the presence and secretion of pTDP-43+/CD63+ vesicles from apilimod-treated *C9ORF72* ALS/FTD iMNs (Figure 4I). Blocking exocytosis with *RAB27A* suppression increased pTDP-43+/LC3+/CD63+ vesicles within apilimod-treated *C9ORF72* ALS/FTD iMNs, providing further evidence that PIKFYVE inhibition induced the secretion of pTDP-43+ exosomes (Figures 4G and 4H). Apilimod reduced the number and size of cytoplasmic pTDP-43+ punctae without affecting their localization or intensity in *C9ORF72* ALS/FTD iMNs (Figure S4J). Thus, PIKFYVE inhibition mitigates pTDP-43 pathology in *C9ORF72* ALS/FTD iMNs by increasing the formation and secretion of pTDP-43+ amphisomes and multivesicular bodies.

### PIKFYVE inhibition improves iMN proteostasis and survival for diverse forms of ALS

Since the accumulation of misfolded proteins is common to all patients with ALS, we hypothesized that activating exocytosis might increase iMN survival for diverse forms of ALS. For iMNs from 8 sporadic ALS (sALS) lines without known ALS mutations, apilimod and PIKFYVE ASO treatment ameliorated TDP-43 mislocalization and neurodegeneration in a dose-dependent manner (Figures 5A–5E; Figures S5A and S5B; Table S1).

Although wild-type TDP-43 aggregates in most ALS cases, some patients harbor ALS-causing mutations in TDP-43, such as G298S, that promote its aggregation.<sup>40,41</sup> TDP-43<sup>G298S</sup> iMNs degenerated significantly faster than controls, and apilimod potently increased their survival (Figure 5F; Figure S5F). A small percentage of patients with ALS harbor disease-causing fused in sarcoma (*FUS*) mutations and develop neurotoxic *FUS* pathology.<sup>42</sup> We generated iMNs from two patients with ALS who carried causal *FUS* mutations (H517Q, R522R) that drive *FUS* mislocalization in iMNs.<sup>43</sup> Indeed, *FUS* ALS iMNs displayed decreased survival and nuclear:cytoplasmic *FUS* ratios, and apilimod ameliorated these phenotypes (Figure 5F; Figures S5F and S5G). Suppression of *RAB27A*, *VAMP7*, or *ATG7* or GW4869 treatment blocked the ability of apilimod to increase sporadic, *TARDBP*, and *FUS* ALS iMN survival (Figures 5F and 5G; Figures S5A and S5F–S5M; Data S1). Similar to iMNs with TDP-43-mediated disease processes, blocking autophagic flux with bafilomycin did not affect apilimod's efficacy in *FUS* ALS iMNs (Figures S3I and S5N). Thus, PIKFYVE inhibition extends the survival of *C9ORF72*, *TARDBP*, *FUS*, and sALS iMNs in an exocytosis-dependent manner.

### PIKFYVE inhibition increases motor function in vivo

Apilimod and *Pikfyve* RNAi both ameliorated motor defects in TDP-43<sup>G298S</sup>-expressing *Drosophila* larvae as determined by larval turning assay (Figures 5H and 5I).<sup>44</sup> Apilimod also reduced the percentage of TDP-43<sup>A315T</sup>-expressing *C. elegans* displaying paralysis and neurodegeneration (Figures S5O–S5Q).<sup>45</sup> In mice, intrathecal injection of apilimod increased cerebrospinal fluid (CSF) levels of PIKFYVE inhibition-induced exosome markers such as OPTN, LC3, and WIPI2 (autophagosome markers), Huntingtin and superoxide dismutase type 1 (SOD1) (aggregation-prone proteins), and ANXA2 (leaderless protein), suggesting that PIKFYVE inhibition can induce exocytosis *in vivo* (Figure S6A). To chronically and stably activate exocytosis, we genetically suppressed *Pikfyve* by two approaches—an ASO targeting *Pikfyve* or genetic deletion of exon 6 in one copy of *Pikfyve* using a CMV-Cre driver line crossed with a published *Pikfyve*-flox line (Figures S6B–S6G; Data S1).<sup>46,47</sup> Deletion of exon 6 in one copy of *Pikfyve* or intracerebroventricular *Pikfyve* ASO administration reduced PIKFYVE mRNA and protein levels by

### Figure 4. PIKFYVE inhibition clears pTDP-43 through amphisome and multivesicular body exocytosis

(A) Hazard ratios of CTRL/*C9*-ALS/FTD iMNs treated with DMSO, apilimod, apilimod + GW4869, or GW4869. Mean of independent iMN conversions ± SEM. One-way ANOVA.  
(B) Hazard ratios of CTRL (2 lines in aggregate) or *C9*-ALS/FTD iMNs (2 lines in aggregate) treated with the ASOs indicated plus DMSO/apilimod. Mean of independent iMN conversions ± SEM. One-way ANOVA.  
(C and D) Immunoblots/quantification of TSG101 in exosomal or pellet fractions from CTRL/*C9*-ALS/FTD iMNs. Cells were treated with the ASOs indicated and DMSO/apilimod. One-way ANOVA. Mean ± SEM.  
(E and F) Immunoblots/quantification of exosomal pTDP-43 (50 kDa) from *C9*-ALS/FTD iMNs treated with the ASOs indicated and DMSO/apilimod. Exosomal TSG101 controlled for exosome secretion/collection. One-way ANOVA. Mean ± SEM.  
(G and H) Immunostaining/quantification of pTDP-43+/CD63+/LC3B+ punctae in *C9*-ALS/FTD iMNs treated with negative control or *RAB27A* ASO and DMSO/apilimod. (G) Confocal z axis scanning. Yellow arrows denote pTDP-43 colocalized with CD63 and LC3B in 3D space. White arrows denote other colocalized pTDP-43+/CD63+/LC3B+ punctae. (H) Gray circles = the average number of pTDP-43+/CD63+/LC3B+ punctae/μm<sup>2</sup> in one *C9*-ALS/FTD iMN. One-way ANOVA. Mean ± SEM.  
(I) Correlative light and electron microscopic images of secreted CD63+ vesicles containing TDP-43 on the membrane of a *C9*-ALS/FTD iMN treated with apilimod. Top row: light microscopy images. Bottom row: electron microscopy images of the same cell. Arrows mark a CD63+/TDP-43+ vesicle in the process of being secreted.  
See also Figure S4.







50% (Figures S6B–S6G; Data S1). *Pikfyve* ASO treatment increased CSF OPTN levels, suggesting that suppressing *Pikfyve* by ~50% activated exocytosis *in vivo* (Figure 6A; Data S1). Single-allele deletion of *Pikfyve* or *Pikfyve* ASO treatment ameliorated motor impairment in the TAR4/4 *Thy1::TDP-43* mouse model and extended survival by 25%–30% (Figures 6B–6E; Figures S6H–S6K; Video S1).<sup>48</sup> GW4869 blocked stimulation of exocytosis by the ASO and impaired its ability to increase TDP-43 mouse motor function (Figures 6F and 6G; Figures S6L and S6M; Video S2; Data S1). A 5-fold lower ASO dose (5  $\mu$ g) significantly improved motor function in TDP-43 mice, suggesting a therapeutic index of at least 5 $\times$  (Figures S6N–S6P). Thus, *Pikfyve* suppression improves motor function and survival of TDP-43 mice in an exocytosis-dependent manner.

We next examined *C9ORF72* disease processes using an AAV-GR100-GFP mouse model.<sup>49</sup> AAV-GR100-GFP-infected mice displayed GFP+/poly(GR)+ cells in the cortex, whereas AAV-GFP-infected mice only possessed GFP+/poly(GR)– cells (Figure S6Q). *Pikfyve* ASO treatment at P4 and P30 reduced hindlimb clasping and extended median survival of AAV-GR100-GFP mice by at least 60% beyond day 57, i.e., the last time point tested (Figures 6H and 6I).

### ***Pikfyve* suppression reduces TDP-43 and *C9ORF72* pathology and neurodegeneration *in vivo***

*Pikfyve* ASO treatment lowered cytoplasmic pTDP-43+ (Ser403/404) punctae and pTDP-43 levels, reduced TDP-43 mislocalization, and fully prevented the loss of large neurons in the spinal cord ventral horn at day 21, and GW4869 blocked these therapeutic effects (Figures 7A–7F; Figures S7A and S7B; Data S1).

Beyond the ventral spinal cord, *Pikfyve* ASO administration reduced pTDP-43+ punctae in dorsal spinal cord neurons and CTIP2+ layer V neurons in the motor cortex in TDP-43 mice (Figures S7C and S7D). In the caudal putamen, neurons did not display elevated pTDP-43+ punctae in TDP-43 mice, and *Pikfyve* ASO treatment did not alter neuronal pTDP-43+ punctae levels (Figure S7E). Therefore, *Pikfyve* suppression ameliorates TDP-43 pathology in multiple regions of the CNS without spreading pathology to neurons in surrounding areas.

In contrast to neurons, microglia displayed more pTDP-43+ punctae after *Pikfyve* ASO treatment, suggesting that they received the pTDP-43 exocytosed from neighboring neurons (Figures 7I and 7J). Consistent with this, iPSC-derived microglia exposed to exosomes harvested from apilimod-treated *C9ORF72* ALS/FTD iMNs displayed a large increase in pTDP-

43+ (Ser409/410) punctae and amoeboid morphology within 24 h (Figures S7F–S7H). By 96 h, the pTDP-43+ punctae and amoeboid morphology decreased significantly, suggesting that microglia can take up and clear the pTDP-43 secreted from ALS neurons and that changes in microglial morphology are transient and reversible (Figures S7F–S7H). *Pikfyve* ASO treatment did not increase the number of microglia or astrocytes in control or TDP-43 mice and in fact reduced microgliosis in TDP-43 mice, indicating that activation of exocytosis did not stimulate gliosis (Figures S7I and S7J).

To examine efficacy over a longer disease course, we employed TAR4 mice, which express less human TDP-43 and develop neurodegeneration at 10 months instead of 1 month.<sup>48</sup> To reduce *Pikfyve* expression by 50%, we crossed TAR4 mice with *Pikfyve*-flox mice harboring a ubiquitous Cre driver.<sup>46</sup> At 10 months, *Pikfyve*<sup>+/-</sup>;TAR4 mice possessed more large ventral horn neurons and fewer pTDP-43+ (Ser409/410) punctae in these neurons than *Pikfyve*<sup>+/-</sup>;TAR4 mice (Figures 7K–7N). With respect to *C9ORF72* disease processes, *Pikfyve* ASO treatment ameliorated the loss of large ventral horn neurons in AAV-GR100-GFP mice at day 26 (Figures 7O and 7P).

Severe reduction in PIKFYVE activity can lead to enlarged LAMP1+ vacuoles that might drive cytotoxicity.<sup>10,50</sup> However, we did not observe abnormalities in the number of RAB7+ late endosomes, percentage of large LAMP1+ lysosomes, or number of LC3+/LAMP1+ vesicles in *Pikfyve* ASO-treated mice with ~50% reduced PIKFYVE levels or 10-month-old *Pikfyve*<sup>+/-</sup> mice (Figures S6G, S7K–S7M, and S7O). *Pikfyve* ASO treatment did not affect TFEB localization in the brain (Figure S7N; Data S1). Moreover, apilimod-treated iMNs in the middle of the *in vitro* survival assay (day 8) did not display increased LAMP1+ punctae (Figure S7P).

## **DISCUSSION**

Further investigation could increase the therapeutic implications of our findings. PIKFYVE inhibition provides one means to stimulate exocytosis of pathological proteins, but other mechanisms that stimulate exosome formation and release may provide unforeseen advantages. It will also be important to determine if PIKFYVE inhibition-induced exocytosis can clear other disease-associated proteins. With regard to safety, *Pikfyve* heterozygous mice are healthy, and reduction of PIKFYVE activity does not become cytotoxic until it is more severely reduced.<sup>10,50</sup> A recent study suggests that prions induce vacuolization in neurons by reducing PIKFYVE

### **Figure 5. PIKFYVE inhibition improves iMN proteostasis and survival for diverse forms of ALS**

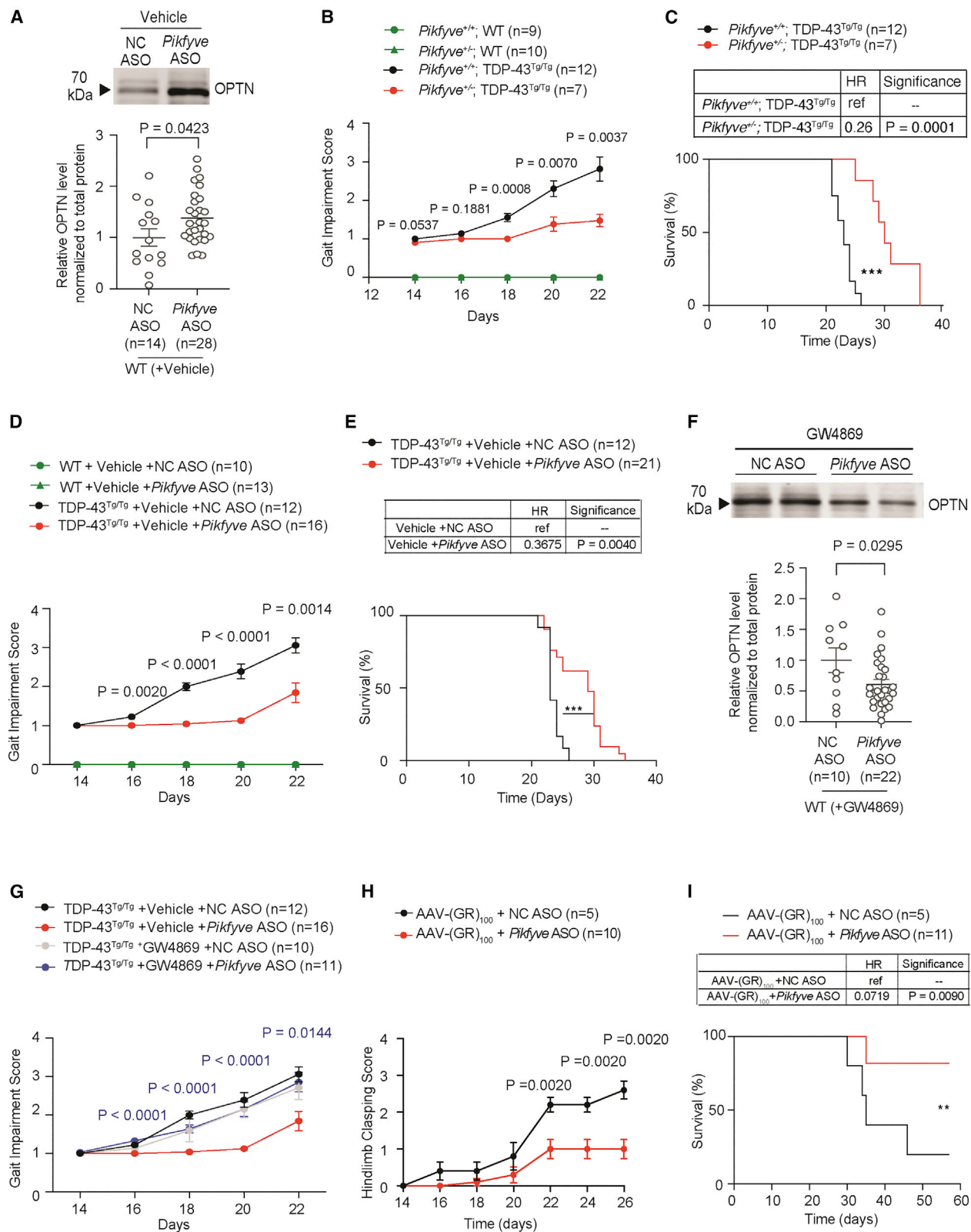
(A–D) Immunostaining/quantification of total TDP-43 in CTRL/sporadic ALS (sALS) iMNs (A and B) or in sporadic ALS iMNs (C and D) treated with DMSO/apilimod. Solid and dotted lines outline the cell body and nucleus, respectively. Gray circles = the nuclear:cytoplasmic TDP-43 ratio from one iMN. Mann-Whitney test. Median  $\pm$  interquartile range.

(E and F) Hazard ratios of iMNs from CTRL/sporadic ALS lines (E) or CTRL, two *FUS* ALS (H517Q, R522R mutation), and *TARDBP* ALS (G298S mutation) lines (F) treated with DMSO/apilimod. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.

(G) Hazard ratios of iMNs from CTRL, two sporadic ALS, and one *FUS* ALS (R522R mutation) line treated with negative control or *RAB27A* ASO plus DMSO/apilimod. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.

(H and I) Larval turning time of TDP-43<sup>G298S</sup> *Drosophila* larvae with *Pikfyve* RNAi (H) or apilimod treatment (I). *n* = 33 larvae/group. One-way ANOVA. Mean  $\pm$  SEM. See also Figure S5.





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activity.<sup>50</sup> Although vacuolization only emerged after severe reduction of PIKFYVE and the contribution of vacuolization to neuron viability was not determined, it would be important to consider these data during therapeutic development.<sup>50</sup> The unusually broad efficacy we observed across models of different forms of ALS merits further investigation into this unconventional protein clearance mechanism and its potential as a therapeutic approach.

### Limitations of the study

It will be important to determine if stable *PIKFYVE* suppression or transient *PIKFYVE* inhibition, such as that induced by ASO or small molecule treatment, respectively, is most efficacious. Stable *PIKFYVE* suppression may provide better protection against spreading of pathology after secretion if it prevents uptake of extracellular TDP-43 or DPRs by other neurons, which it does for tau and alpha-synuclein.<sup>51,52</sup>

### STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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### SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.cell.2023.01.005>.

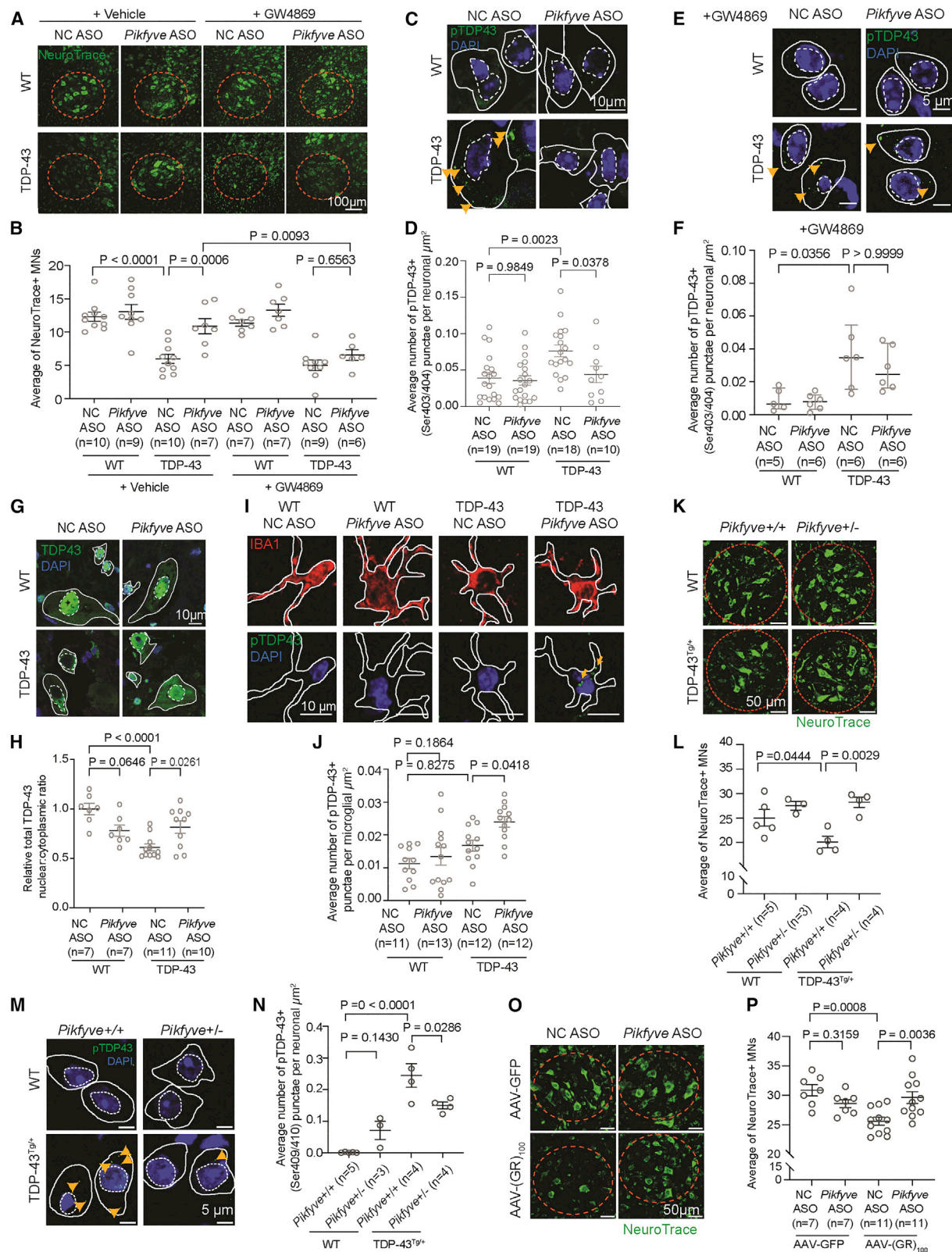
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### Figure 6. *Pikfyve* suppression improves motor function and extends survival of TDP-43 and *C9ORF72* mice

- (A) Immunoblot/quantification of OPTN in CSF with negative control/*Pikfyve* ASO intracerebroventricular administration in neonatal mice. Unpaired t test. Mean ± SEM. Data points = one mouse.
- (B) Gait impairment scores of *Pikfyve*<sup>+/+</sup>;WT, *Pikfyve*<sup>-/-</sup>;WT, *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/Tg</sup>, and *Pikfyve*<sup>-/-</sup>;TDP-43<sup>Tg/Tg</sup> mice. Unpaired t test at each time point. Mean ± SEM.
- (C) Survival of *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/Tg</sup> and *Pikfyve*<sup>-/-</sup>;TDP-43<sup>Tg/Tg</sup> mice. Log rank test.
- (D) Gait impairment scores of WT or TDP-43<sup>Tg/Tg</sup> mice treated with vehicle and negative control/*Pikfyve* ASO. Unpaired t test at each time point. Mean ± SEM.
- (E) Survival of TDP-43<sup>Tg/Tg</sup> mice treated with vehicle and negative control/*Pikfyve* ASO. Log rank test.
- (F) Immunoblot/quantification of OPTN in CSF after administration of GW4869 and negative control/*Pikfyve* ASO. Unpaired t test. Mean ± SEM. Data points = one mouse.
- (G) Gait impairment scores of TDP-43<sup>Tg/Tg</sup> mice treated with vehicle/GW4869 and negative control/*Pikfyve* ASO. Unpaired t test at each time point. Mean ± SEM.
- (H) Hindlimb clasping of AAV-eGFP-(GR)<sub>100</sub> mice treated with negative control/*Pikfyve* ASO. Unpaired t test at each time point. Mean ± SEM.
- (I) Survival of AAV-(GR)<sub>100</sub> mice treated with negative control/*Pikfyve* ASO. Log rank test.
- See also Figure S6.





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#### AUTHOR CONTRIBUTIONS

S.-T.H., G.R.L., W.-H.C., and J.K.I. conceived the project. S.-T.H., G.R.L., W.-H.C., Y.S., M.S., C.K., S.P., S.D., C.M., S.V.A., Y.W., B.V.Z., D.K.D., J.A.P., D.C.Z., F.-B.G., and J.K.I. designed the experiments. S.-T.H., G.R.L., W.-H.C., Y.E., S.M., S.H., Y.S., M.S., C.K., S.M.-I., S.P., S.D., C.M., J.C., J.P., A.C., J.L., Y.L., G.K., E.H., Y.W., B.V.Z., D.K.D., J.A.P., D.C.Z., and J.K.I. performed experiments and interpreted data. S.-T.H. and J.K.I. prepared the manuscript. All authors discussed the results and commented on the manuscript.

#### DECLARATION OF INTERESTS

J.K.I. and S.-T.A. are co-founders of AcuraStem, Inc. S.-T.A., W.-H.C., S.M., and S.H. are employees of AcuraStem, Inc. J.K.I. is a co-founder of Modulo Bio, serves on the scientific advisory boards of AcuraStem, Spinogenix, Synapticure, and Vesalius Therapeutics, and is employed at BioMarin Pharmaceutical. B.V.Z. is a co-founder of ZZ Biotech and chairman of its scientific advisory board.

J.A.P. is a co-founder of Modelis. F.-B.G. receives research funding from Stealth BioTherapeutics.

#### INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a gender minority in their field of research. One or more of the authors of this paper self-identifies as a member of the LGBTQIA+ community. One or more of the authors of this paper received support from a program designed to increase minority representation in their field of research.

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#### Figure 7. *Pikfyve* suppression reduces TDP-43 and C9ORF72 pathology and neurodegeneration in vivo

(A and B) Images/quantification of large, NeuroTrace (Nissl)+ lateral motor column (LMC) neurons in the lumbar spinal cord in WT/TDP-43<sup>Tg/Tg</sup> mice treated with vehicle/GW4869 and negative control/*Pikfyve* ASO. Data points = the average number of NeuroTrace+ LMC neurons/ventral horn hemicord section for one mouse. One-way ANOVA. Mean ± SEM. Red dotted line = the LMC region.

(C–F) Immunostaining/quantification of pTDP-43+ (Ser403/404) punctae in large TUJ1+ neurons in the spinal cord ventral horn from WT/TDP-43<sup>Tg/Tg</sup> mice treated with vehicle (C and D) or GW4869 (E and F) and negative control/*Pikfyve* ASO. Data points = the average number of pTDP-43+ punctae in TUJ1+ neurons/μm<sup>2</sup> for one mouse. One-way ANOVA. Mean ± SEM. Solid and dotted lines outline the cell body and nucleus, respectively. Arrows mark pTDP-43+ punctae.

(G and H) Immunostaining of total TDP-43 in large TUJ1+ neurons in the ventral horn of WT/TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the average nuclear:cytoplasmic TDP-43 ratio in TUJ1+ neurons for one mouse. One-way ANOVA. Mean ± SEM. Solid and dotted lines outline the cell body and nucleus, respectively.

(I and J) Immunostaining/quantification of pTDP-43+ (Ser403/404) punctae in IBA1+ microglia in the spinal cord ventral horn from WT/TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the average number of pTDP-43+ punctae in IBA1+ microglia/μm<sup>2</sup> for one mouse. One-way ANOVA. Mean ± SEM. Solid lines outline the cell body. Arrows mark pTDP-43+ punctae.

(K, L, O, and P) Representative images and quantification of large NeuroTrace+ LMC neurons in the lumbar spinal cord in 10-month-old *Pikfyve*<sup>+/+</sup>;WT, *Pikfyve*<sup>+/−</sup>;WT, *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/+</sup>, and *Pikfyve*<sup>+/−</sup>;TDP-43<sup>Tg/+</sup> mice (K and L) or day 26 AAV-eGFP or AAV-eGFP-(GR)<sub>100</sub> mice treated with negative control/*Pikfyve* ASO (O and P). Data points = the average number of large NeuroTrace+ LMC neurons/ventral horn hemicord section for one mouse. One-way ANOVA. Mean ± SEM. Red dotted line = the LMC region.

(M and N) Immunostaining/quantification of pTDP-43+ (Ser409/410) punctae in large TUJ1+ neurons in spinal cord ventral horn from 10-month-old *Pikfyve*<sup>+/+</sup>;WT, *Pikfyve*<sup>+/−</sup>;WT, *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/+</sup>, and *Pikfyve*<sup>+/−</sup>;TDP-43<sup>Tg/+</sup> mice. Data points = the average number of pTDP-43+ punctae/μm<sup>2</sup> in TUJ1+ neurons for one mouse. One-way ANOVA. Mean ± SEM. Solid and dotted lines outline the cell body and nucleus, respectively. Arrows mark pTDP-43+ punctae.

See also Figure S7.



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## STAR★METHODS

### KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
TSG101 (mouse IgG1)	BD Transduction Laboratories™	Cat# 612697; RRID: AB_399937
TUJ1 (Mouse IgG2a, κ)	Biolegend	Cat# 801213; RRID: AB_2728521
LC3 (Rabbit IgG)	MBL	Cat# PM036; RRID: AB_2274121
P62 (Mouse IgG1, κ)	BD Transduction	Cat# 610832; RRID: AB_398151
OPTN (Rabbit IgG)	Proteintech	Cat# 10837-1-AP; RRID: AB_2156665
Phospho-TDP43(Ser409/410) (Rabbit IgG)	Proteintech	Cat# 22309-1-AP; RRID: AB_11182943
PIP5K3 (Rabbit IgG)	Proteintech	Cat# 13361-1-AP; RRID: AB_10638310
TFEB (Rabbit IgG)	Bethyl Laboratories®	Cat# A303-673A; RRID: AB_11204751
Phospho-TFEB (Ser142) (Rabbit IgG)	Sigma-Aldrich	Cat# ABE1971-I
Huntingtin (Rabbit IgG)	Abcam	Cat# ab109115; RRID: AB_10863082
alpha Synuclein (Rabbit IgG)	Thermo Fisher Scientific	Cat# PA5-85791; RRID: AB_2792927
Fibrillarin (Rabbit IgG)	Abcam	Cat# ab166630; RRID: AB_2927645
HSP90 (Rabbit IgG)	Proteintech	Cat# 13171-1-AP; RRID: AB_2120924
GR repeat (Rabbit IgG)	Proteintech	Cat# 23978-1-AP; RRID: AB_2879387
PR repeat (Rabbit IgG)	Proteintech	Cat# 23979-1-AP; RRID: AB_2879388
GA repeat (Rabbit IgG)	Proteintech	Cat# 24492-1-AP; RRID: AB_2879571
TDP43 (Rabbit IgG)	ProteinTech	Cat# 10782-2-AP; RRID: AB_615042
Poly-GA (Mouse IgG)	Millipore	Cat# MABN889; RRID: AB_2728663
LysoSensor™ Green DND-189	Invitrogen	L7535
LysoTracker™ Deep Red	Invitrogen	L12492
LC3B (Rabbit IgG)	Abcam	Cat# ab192890; RRID: AB_2827794
LAMP1 (Mouse IgG)	Abcam	Cat# ab25630; RRID: AB_470708
CD63 (mouse IgG1, κ)	BD Transduction Laboratories™	Cat# 556019; RRID: AB_396297
RAB7 (Rabbit IgG)	Abcam	Cat# ab137029; RRID: AB_2629474
Phospho-TDP43(Ser409/410) (Rabbit IgG)	Cosmo Bio	Cat# CAC-TIP-PTD-P01; RRID: AB_1961899
Alexa Fluor® 647 Anti-LC3B antibody	Abcam	Cat# ab225383; RRID: AB_2928102
FUS (Goat IgG)	Bethyl Laboratories®	Cat# A303-839A; RRID: AB_2620190
IBA1 (Goat IgG)	Abcam	Cat# ab5076; RRID: AB_2224402
MAP2 (Chicken IgY)	Abcam	Cat# ab5392; RRID: AB_2138153
TUJ1/Tubb3 (Mouse IgG)	Neuromics	Cat# CH23005; RRID: AB_2210684
NeuN (Mouse IgG)	Millipore	Cat# MAB377; RRID: AB_2298772
ChAT (Goat polyclonal)	Millipore	Cat# AB144P; RRID: AB_2079751
Phospho TDP43(Ser403/404) (Mouse IgG2a)	ProteinTech	Cat# 66079-1-Ig; RRID: AB_11183767
LAMP1 (Rabbit IgG)	Abcam	Cat# ab24170; RRID: AB_775978
RAB7 (Mouse IgG1 κ)	Santa Cruz	Cat# sc-376362; RRID: AB_10987863
LC3B (Mouse IgG2a)	GeneTex	Cat# GTX632501; RRID: AB_2888317
GFP (Chicken IgY)	Aves Lab	Cat# GFP-1010; RRID: AB_2307313
GFAP (Rabbit IgG)	Abcam	Cat# ab7260; RRID: AB_305808
Iba1 (Rabbit IgG)	GeneTex	Cat# GTX101495; RRID: AB_1240433
DAPI	Thermo Fisher	62248

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
NeuroTrace™ 500/525 Green Fluorescent Nissl Stain	Thermo Fisher	N21480
Goat anti-Chicken IgY (H+L) Cross-Adsorbed Secondary Antibody, Alexa Fluor™ Plus 405	Thermo Fisher	A48260
Donkey anti-Rabbit, IgG (H+L) Highly Cross-Adsorbed, Alexa Fluor® 488	Invitrogen	A-21206
Donkey anti-Mouse, IgG (H+L) Highly Cross-Adsorbed, Alexa Fluor® 555	Invitrogen	A31570
Donkey anti-Mouse, IgG (H+L) Highly Cross-Adsorbed, Alexa Fluor® 488	Invitrogen	A21202
Donkey anti-Mouse, IgG (H+L) Highly Cross-Adsorbed, Alexa Fluor® 647	Invitrogen	A32787
Donkey Anti-Chicken IgY (IgG) (H+L) Alexa Fluor® 647	Jackson ImmunoResearch	703-605-155
Goat anti-Chicken IgY (H+L) Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488	Invitrogen	A32931TR
Donkey anti-Goat, IgG (H+L) Highly Cross-Adsorbed, Alexa Fluor 555	Invitrogen	A32816
IRDye® 680RD Donkey anti-Rabbit IgG Secondary Antibody	LI-COR Biosciences	926-68073
IRDye® 800CW Donkey anti-Mouse IgG Secondary Antibody	LI-COR Biosciences	926-32212
CFSE dye	Thermo	C34554
<b>Chemicals, peptides, and recombinant proteins</b>		
DMEM	Gibco	11995-065
Opti-MEM	Gibco	31985-070
FBS	GenClone	25-514
Trypsin	GenClone	25-510
Polybrene (Hexadimethrine bromide)	Sigma	H9268-5G
DMEM/F12	Corning	10-090-CV
Laminin	Invitrogen	23017015
Glutamax	Gibco	35050-061
N2	Gibco	17502048
B27	Gibco	17504044
Human FGF-beta	Peptotech	100-18B
Repsox	Selleckchem	S7223
human CNTF	R&D	257-NT
human BDNF	R&D	248BDB
human GDNF	R&D	212-GD
Matrigel	Corning	354277
mTeSR	StemCell	85851
Accutase	Innovative Cell Technologies	AT4104
Y-27632 2HCl	selleckchem	S1049
Lenti-X concentrator	TaKaRa	631232
Polybrene (Hexadimethrine bromide)	Sigma	H9268-5G
Neurobasal-A	Life Technologies	10888-022
Apilimod	achemblock	O33822
GW 4869 (hydrochloride hydrate)	Cayman Chemicals	13127
SB431542	Cayman Chemical	13031
CHIR99021	Cayman Chemical	13122
Retinoic Acid	Millipore Sigma	R2625

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
DMH1	selleckchem	S7146
Purmorphamine	Cayman Chemical	10009634
Donor Equine Serum	HyClone	16777-030
BMP4	Invitrogen	PHC9534
VEGF	Invitrogen	PHC9394
SCF	Miltenyi	130-096-692
X-VIVO15	SLS (Lonza)	LZBE02-060F
M-CSF	Invitrogen	PHC9501
IL-3	Invitrogen	PHC0033
2-mercaptoethanol	Gibco	21985023
IL-34	Biolegend	577906
Bafilomycin A1	Cayman	11038
HEPES Stock Solution (1 M)	Cayman	600212
Citrate Buffer pH 6.0	Electron Microscopy Sciences (EMS)	64142-07
Normal Donkey Serum	VWR	102644-006
cOmplete™, EDTA-free Protease Inhibitor Cocktail	Roche	4693132001
PhosSTOP™	Roche	4906845001
Chameleon® Duo Pre-stained Protein Ladder	Licor	928-60000
Odyssey® One-Color Protein Molecular Weight Marker	Licor	928-40000
MEM media	Life Technologies	10370088
Urea	Sigma	U5378-100G
Thiourea	Sigma	T8656-50G
CHAPS	Sigma	10810118001
Tris-HCl 1.5M, pH=8.8	Teknova	T1588
Penicillin-streptomycin 50X	Corning	30-001-CI
Compound E	Cayman Chemicals	15579
PEG 6000	Sigma-Aldrich	81253
<b>Experimental models: Cell lines</b>		
ND03231 (CTRL1)	NINDS Biorepository	3231
ND05280 (CTRL2)	NINDS Biorepository	5280
ND03719 (CTRL3)	NINDS Biorepository	3719
ND00184 (CTRL4)	NINDS Biorepository	00184
ND41865 (CTRL5)	NINDS Biorepository	41865
ND06769 (C9-ALS 1)	NINDS Biorepository	6769
ND10689 (C9-ALS 2)	NINDS Biorepository	10689
ND12099 (C9-ALS 3)	NINDS Biorepository	12099
ND50000 (C9-ALS 4)	NINDS Biorepository	50000
ND10739 (sALS1)	NINDS Biorepository	10739
ND13454 (sALS2)	NINDS Biorepository	13454
ND11813 (sALS3)	NINDS Biorepository	11813
ND09711 (sALS4)	NINDS Biorepository	9711
ND14185 (sALS5)	NINDS Biorepository	14185
ND08705 (sALS6)	NINDS Biorepository	8705
ND09292 (sALS7)	NINDS Biorepository	9292
ND09329 (sALS8)	NINDS Biorepository	9329
ALS93E (sALS9)	This Manuscript	N/A
ND35663 (FUS H517Q)	NINDS Biorepository	35663

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
ND39034 (FUS R522R)	NINDS Biorepository	39034
ND50007 (TDP-43 G298S)	NINDS Biorepository	50007
<b>Experimental models: Animal Models</b>		
<i>C. elegans</i> : TDP-43A315T	Parker lab	N/A
<i>Drosophila</i> : Wild type ( $w^{1118}$ )	Zarnescu and Dickman labs	N/A
<i>Drosophila</i> : GR.100 larvae ( $w$ ;OK6-Gal4/UAS-100xGR)	Dickman lab	N/A
<i>Drosophila</i> : $w^{1118}$ ; UAS-TDP-43 <sup>G298S</sup> -YFP	Zarnescu lab	N/A
<i>Drosophila</i> : $y[1]v[1]$ ; $P\{Y[+(7.7) = GSYP]attP2$	Bloomington Drosophila Stock Center	BDSC # 8622
<i>Drosophila</i> : $y[1]sc[*]v[1]$ ; $P\{y[+t7.7]v[+t1.8]=TRIIP.GL00246\}attP2$	Bloomington Drosophila Stock Center	BDSC # 35793
<i>Drosophila</i> : $P\{UAS-poly-GR.PO-100\}attP40$	Bloomington Drosophila Stock Center	BDSC#58696
Mice: TDP-43 transgenic mice	Jackson Laboratory	012836
Mice: B6.Cg-PIKFYVE<tm1.1Ashi>/J	Jackson Laboratory	029331
Mice: B6.C-Tg(CMV-Cre)1Cgn/J	Jackson Laboratory	006054
Mice: C57BL/6J mice	Jackson Laboratory	000664
<b>Oligonucleotides</b>		
sgRNA-1 targeting upstream of repeat expansion	This Manuscript	GUAACCUACGGUGUCCCGCU
sgRNA-2b targeting downstream of repeat expansion	This Manuscript	ACCCCAAACAGCCACCCGCC
Primer 1 for repeat primed PCR	This Manuscript	FAM-tgtaaaacgacggccagtCAAGGA GGGAAACAACCGCAGCC
Primer 2 for repeat primed PCR	This Manuscript	caggaaacagctatgaccGGGCCCGCCCC GACCACGCCCCGGCCCCGGCCCCGG
Primer 3 for repeat primed PCR	This Manuscript	caggaaacagctatgacc
Forward primer for Southern Probe	This Manuscript	AGAACAGGACAAGTTGCC
Reverse primer for Southern Probe	This Manuscript	AACACACACCTCCTAAACC
qPCR primer for <i>RAB27A</i> , Forward	This Manuscript	AGAGGCCAGAGAATCCACCT
qPCR primer for <i>RAB27A</i> , Reverse	This Manuscript	CACACCGTTCCATTGCTTC
qPCR primer for <i>HPRT</i> , Forward	This Manuscript	GACTTTGCTTCTTGCTGTCAG
qPCR primer for <i>HPRT</i> , Reverse	This Manuscript	GGCTTATATCCAACACTTCGTGGG
qPCR primer for <i>VAMP7</i> , Forward	This Manuscript	CGGTTCAAGAGCACAGACAGCA
qPCR primer for <i>VAMP7</i> , Reverse	This Manuscript	ATCCACTTGGGCTTGAGTCTCC
qPCR primer for <i>RAB8A</i> , Forward	This Manuscript	TCAGGAACGGTTTCGGACGATC
qPCR primer for <i>RAB8A</i> , Reverse	This Manuscript	GCTCCTCAATGTTGCGAATCCAG
qPCR primer for <i>HSP8A</i> , Forward	This Manuscript	TCCTACCAAGCAGACACAGACC
qPCR primer for <i>HSP8A</i> , Reverse	This Manuscript	CAGGAGGTATGCCTGTGAGTTC
qPCR primer for <i>NSMAF</i> , Forward	This Manuscript	GTCTGAACACCTTCACGAGTGG
qPCR primer for <i>NSMAF</i> , Reverse	This Manuscript	CTGTTCAAGTCTACACCTCCTTC
qPCR primer for <i>ATG7</i> , Forward	This Manuscript	CGTTGCCACAGCATCATCTTC
qPCR primer for <i>ATG7</i> , Reverse	This Manuscript	CACTGAGGTTCAACATCCTTGG
qPCR primer for <i>MCOLN1</i> , Forward	This Manuscript	CTGATGCTGCAAGTGGTCAAG
qPCR primer for <i>MCOLN1</i> , Reverse	This Manuscript	GGTGTCTCTTCCCGGAATGTC
qPCR primer for <i>GORASP1</i> , Forward	This Manuscript	CACGGGGATGTGGAACCATC
qPCR primer for <i>GORASP1</i> , Reverse	This Manuscript	ATCAGCTTCAAGGGCTTCCC
qPCR primer for <i>SMPD3</i> , Forward	This Manuscript	GAAGCACACCTCAGGACCAAAAG
qPCR primer for <i>SMPD3</i> , Reverse	This Manuscript	CAGCCAGTCCTGAAGCAGGTC

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**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
qPCR primer for <i>PIKFYVE</i> , Forward	This Manuscript	GGCTCACGCCTATAATCCTATC
qPCR primer for <i>PIKFYVE</i> , Reverse	This Manuscript	GCTGGAATGCAATGGTGAATC
qPCR primer for <i>GAPDH</i> , Forward	This Manuscript	CGAGATCCCTCCAAAATCAA
qPCR primer for <i>GAPDH</i> , Reverse	This Manuscript	GTCTTCTGGGTGGCAGTGAT
qPCR primer for <i>MAP2</i> , Forward	This Manuscript	GGAACCAACTCTCTCTGGATT
qPCR primer for <i>MAP2</i> , Reverse	This Manuscript	GCATTCTCTCTCAGCCTTCT
qPCR primer for truncated <i>STMN2</i> , Forward	This Manuscript	GGACTCGGCAGAAGACCTTC
qPCR primer for truncated <i>STMN2</i> , Reverse	This Manuscript	GCAGGCTGTCTGTCTCTCTC
qPCR primer for <i>Pikfyve</i> , Forward (exon 6)	This Manuscript	GAAGCAATATTGGATGCCAGA
qPCR primer for <i>Pikfyve</i> , Reverse (exon 6)	This Manuscript	CCCATAAAATTTCCAGGGATT
qPCR primer for <i>Pikfyve</i> , Forward	This Manuscript	ACAGTGCTGAAGAAGGGCTC
qPCR primer for <i>Pikfyve</i> , Reverse	This Manuscript	ACAGATCAGGGCTCTTCCCT
qPCR primer for <i>PPIA</i> , Forward	This Manuscript	TCCTGGACCCAAAACGCTCC
qPCR primer for <i>PPIA</i> , Reverse	This Manuscript	CCATGGCAAATGCTGGACCA
DNA primer for S0 genotyping, Forward	This Manuscript	GTGACTAGTGAACCTAGCACATTG
DNA Common primer, TDP-43 genotyping	This Manuscript	TGAAATCCGGGT GGTATTGG
DNA WT primer, TDP-43 genotyping	This Manuscript	GGTGAGTTTAACCTTCAAGGGCT
DNA Transgene primer, TDP-43 genotyping	This Manuscript	AGCTTGCTAGCGGATCCAGAC
DNA primer for S1 genotyping, Forward	This Manuscript	GCCTGAGTTCTGAGAGTGAGTG
DNA primer for S2 genotyping, Forward	This Manuscript	CTGAAGCAATATTGGATGCCAGATAGC
DNA primer for S3 genotyping, Forward	This Manuscript	GTAAAGGTTGACTTGACTTACATGATAG
DNA primer for A1 genotyping, Reverse	This Manuscript	CTATGATGGGAAGAGTGGACAGTC
DNA primer for Cre Control, Forward	This Manuscript	CTAGGCCACAGAATTGAAAGATCT
DNA primer for Cre Control, Reverse	This Manuscript	GTAGGTGGAAATTCTAGCATCATCC
DNA primer for Cre Transgene, Forward	This Manuscript	GCGGTCTGGCAGTAAAACTATC
DNA primer for Cre Transgene, Reverse	This Manuscript	GTGAAACAGCATTGCTGTCACTT
Negative control ASO	This Manuscript	mG*mC*mG*mA*mC*T*A*T*A*C*G*C*G* C*A*mA*mU*mA*mU*mG
<i>PIKFYVE</i> ASO1	This Manuscript	mU*mG*mG*mC*mC*T*C*T*T*C*T*G* C*T*mC*mU*mC*mU*mC
<i>PIKFYVE</i> ASO2	This Manuscript	mG*mC*mU*mG*mG*T*C*A*A*C*T*T* C*C*mA*mC*mU*mC*mA
<i>RAB27A</i> ASO	This Manuscript	mG*mA*mA*mA*mC*C*T*C*T*C*T*G* C*C*mC*mU*mG*mC*mU
<i>VAMP7</i> ASO	This Manuscript	mC*mU*mA*mG*mG*C*T*A*A*A*C*A*G* G*T*mG*mG*mC*mU*mA
<i>RAB8A</i> ASO	This Manuscript	mU*mG*mG*mU*mU*G*A*C*C*T*G*G* T*C*C*mC*mA*mG*mU*mC
<i>HSPA8</i> ASO	This Manuscript	mC*mA*mA*mG*mG*A*A*G*G*T*A*G*T* T*G*mC*mC*mA*mA*mC
<i>NSMAF</i> ASO	This Manuscript	mA*mU*mC*mU*mG*C*C*T*A*A*G*A* G*A*mA*mU*mA*mG*mC
<i>ATG7</i> ASO	This Manuscript	mG*mU*mU*mG*mA*G*T*G*C*A*T*A* C*C*mA*mG*mU*mA*mG
<i>MCOLN1</i> ASO	This Manuscript	mG*mG*mU*mU*mA*G*A*T*G*A*C*C* T*T*mC*mA*mC*mA*mU
<i>GORASP1</i> ASO	This Manuscript	mG*mG*mA*mU*mA*G*A*C*C*T*A*G*T* C*A*mG*mG*mU*mA*mG
<i>SMPD3</i> ASO	This Manuscript	mG*mG*mA*mU*mU*G*T*C*A*A*A*A* C*A*mG*mU*mC*mC*mC

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## Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Negative control ASO ( <i>in vivo</i> )	This Manuscript	/52MOErC/*i2MOErC/ /i2MOErT/ /i2MOErA/ /i2MOErT/ A*G*G*A*C*T*A*T*C*/i2MOErA/ /i2MOErG/ /i2MOErG/* /i2MOErA*/32MOErA/
Pikfyve ASO ( <i>in vivo</i> )	This Manuscript	/52MOErG/*i2MOErC/ /i2MOErA/ /i2MOErT/ /i2MOErC/T*T*G*T*G*G*A*A*C*T*/i2MOErC/ /i2MOErT/ /i2MOErC/* /i2MOErC*/32MOErC/
Deposited data		
Bulk RNA-seq data (DMSO or AP treatment)	This Manuscript	GEO: GSE221015
Bulk RNA-seq data (Negative control or PIKFYVE ASO treatment)	This Manuscript	GEO: GSE221016, GEO: GSE221017
Software and algorithms		
ImageJ	Schneider et al. <sup>53</sup>	<a href="https://imagej.nih.gov/ij/index.html">https://imagej.nih.gov/ij/index.html</a>
Prism	Graphpad	<a href="https://www.graphpad.com/scientific-software/prism/">https://www.graphpad.com/scientific-software/prism/</a>
Enrichr	Kuleshov et al. <sup>54</sup>	<a href="https://maayanlab.cloud/Enrichr/">https://maayanlab.cloud/Enrichr/</a>

## RESOURCE AVAILABILITY

### Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Justin Ichida ([ichida@usc.edu](mailto:ichida@usc.edu)).

### Materials availability

Human iPSC lines used in this study are available from the Ichida lab at the University of Southern California (<https://ichidalab.usc.edu/>).

### Data and code availability

- The bulk RNA-sequencing data generated as part of this study are available at the Gene Expression Omnibus. Accession numbers are listed in the [key resources table](#) and the output of the analyses are available upon request. Original western blot images are included in [Data S1](#). The mass spectrometry raw data is provided in [Table S3](#).
- All code used in this study is available from the [lead contact](#) upon request.
- Any additional information required to reanalyze the data reported in this work paper is available from the [lead contact](#) upon request.

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

### Cell lines

Human iPSC lines were generated by the Ichida lab from lymphoblastoid lines obtained from the NINDS Biorepository ([Table S1](#)). Both male and female iPSC lines were used. The iPSCs were maintained in mTeSR1 medium using a feeder-free culture protocol in six-well plates coated with growth factor-reduced Matrigel and iPSCs below passage 25 were used to generate iMNs. iPSCs were cultured at 37 C and 5% CO<sub>2</sub> with daily feeding of 2mL mTeSR per well. To ensure consistency, cultures were kept below 90% confluency. Passaging of iPSC colonies was carried out using Accutase (Innovative Cell Technologies) to the desired split ratio.

### C. elegans

*C. elegans* mutants were developed and maintained by the Parker lab as previously described.<sup>45</sup> Both male and female worms were used. For motor and histological assessment, age-synchronized L4 worms were transferred to NGM plates and scored daily for paralysis, from day 1 to day 12 of adulthood.

### Drosophila

The wild-type, GR.100, and TDP-43 strains were obtained from the Bloomington Drosophila Stock Center and maintained in the Dickman and Zarnescu labs as described in the [STAR Methods](#). Both male and female larvae were used at the third-instar larvae stage for both turning and NMJ assays.



### Mouse models

FVB/NJ (Stock No: 001800), and TAR4/4 (Stock No: 012836) mice were obtained from Jackson Laboratory. Mice were housed in standard conditions with food and water *ad libitum* in the conventional vivarium at the University of Southern California. All animal use and care were in accordance with local institution guidelines of the University of Southern California and the IACUC board of the University of Southern California with the protocol numbers 21099 and 11938.

### METHOD DETAILS

#### iPSC reprogramming

Human lymphocytes from healthy subjects and ALS patients were obtained from the NINDS Biorepository at the Coriell Institute for Medical Research and reprogrammed into iPSCs as previously described.<sup>8,11</sup>

#### Molecular cloning and viral production

The complementary DNA (cDNA) for each iMN factor (*Ngn2*, *Lhx3*, *Isl1*, *NeuroD1*, *Ascl1*, *Brn2* and *Myt1l*) and mRFP-GFP-LC3 construct was purchased from Addgene and cloned into the pMXs retroviral expression vector using Gateway cloning technology (Invitrogen). The *Hb9::RFP* lentiviral vector was also purchased from Addgene (ID: 37081). Viruses were produced as follows. HEK293T cells were transfected in a 10-cm dish at 80–90% confluence with viral vectors containing each iMN factor and viral packaging plasmids (PIK-MLV-gp and pHDM for retrovirus, pPAX2 and VSVG for lentivirus) using polyethylenimine (PEI) (Sigma-Aldrich). The medium was changed 24 hours after transfection. Viruses were harvested at 48 hours and 72 hours after transfection. Viral supernatants were filtered with 0.45- $\mu$ m filters, incubated with Lenti-X concentrator (Clontech) for 24 hours at 4°C, and centrifuged at 1,500 g at 4°C for 45 minutes. Pellets were resuspended in DMEM plus 10% FBS (200  $\mu$ L per 10-cm dish of HEK293T) and stored at –80°C.

#### Conversion of iPSCs into induced motor neurons

iPSCs were first differentiated into fibroblast-like cells to enable efficient retroviral transduction as described.<sup>11</sup> Reprogramming of fibroblast-like cells was performed in 96-well plates (5 x 10<sup>3</sup> cells/ well) or 13-mm plastic coverslip (3 x 10<sup>4</sup> cells/ coverslip) that had been pre-coated with 0.1% gelatin (1 hour, room temperature) and laminin (4°C, overnight). Seven iMN factors were added in 100  $\mu$ L of fibroblast medium (DMEM plus 10% FBS) per 96-well or 500  $\mu$ L per coverslip with 5  $\mu$ g/ml polybrene. Cultures were transduced with *Hb9::RFP* lentivirus after 48 hours transduction with seven iMN factors. On day 5, primary mouse cortical glial cells from P1 ICR pups were added to the transduced cultures in glia medium containing MEM (Life Technologies), 10% donor equine serum (HyClone), 20% glucose (Sigma-Aldrich), and 1% penicillin/streptomycin. On day 6, cultures were switched to N3 medium containing DMEM/F12 (Life Technologies), 2% FBS, 1% penicillin/streptomycin, N2 and B27 supplements (Life Technologies), 7.5  $\mu$ M RepSox (Selleck), and 10 ng/ml each of FGF, GDNF, BDNF, and CNTF (R&D). The cultures were maintained in N3 with neurotrophic factors (RepSox, FGF, GDNF, BDNF, and CNTF) and changed every other day.

#### Induced motor neuron survival assay

*Hb9::RFP*<sup>+</sup> iMNs appear between day 13–16 after transduction of iMN factors. The iMN survival assay was initiated on day 17. Starting at Day 17, longitudinal tracking of iMNs was performed using Molecular Devices ImageExpress once every other day for 14–20 days. Tracking of neuronal survival was performed using SVcell 3.0 (DRVision Technologies) or ImageJ. For each neuronal survival assay, iMNs were quantified from three biologically independent iMN conversions per line per condition. Neurons were scored as dead when their soma was no longer detectable by RFP fluorescence. For neurotrophic factor withdrawal conditions, BDNF, GDNF, and CNTF were removed from the culture medium on day 17. For treatment with apilimod, cultures were treated with DMSO or 3  $\mu$ M apilimod (Achemblock, O33822) after neurotrophic factor withdrawal (starting at day 17). For treatment with GW4869 to block exocytosis, cultures were treated with 10  $\mu$ M GW4869 (Cayman Chemical, 13127) along with DMSO or 3  $\mu$ M apilimod after neurotrophic factor withdrawal (starting at day 17). For *PIKFYVE* ASO treatment, the iMN cultures were pretreated one time with 10  $\mu$ M ASOs for 48 hours before neurotrophic factor withdrawal on day 17. For *Rab27a*, *VAMP7*, *ATG7*, *RAB8A*, *HSPA8*, *MCOLN1*, *GORASP1*, and *NSMAF* ASO treatments, the iMN cultures were pretreated one time with 9  $\mu$ M ASOs for 24 hours before neurotrophic factor withdrawal (on day 17) and treated with DMSO or 3  $\mu$ M apilimod starting at day 17. ASO gapmers were designed and produced by IDT. All treatments were maintained for the whole survival assay and the medium was changed every other day. For hazard ratio plots, the hazard rate of the first condition plotted on the far left was used as the control and its hazard rate was set as 1. The red dotted line represents this hazard rate. The hazard rates of all other conditions was divided by this hazard rate to generate the hazard ratios.

#### CRISPR/Cas9 genome editing of iPSCs

CRISPR/Cas9-mediated genome editing was performed in human iPSCs as previously described, using Cas9 nuclease.<sup>8</sup> Single guide RNAs (sgRNAs) targeting both sides of the *C9ORF72* intronic hexanucleotide repeat expansion were designed ([key resources table](#)). To generate isogenic control iPSCs by removing the repeat expansion, *C9ORF72* ALS/FTD patient iPSCs were transfected with human codon-optimized Cas9 (Addgene ID: 31825), the appropriate gRNA constructs by nucleofection (Lonza) according to



the manufacturer's protocol and the homologous recombination donor vector. The surviving colonies were picked on day 7 after transfection and genotyped by PCR amplification and sequencing the targeted genomic site. Colonies showing removal of the repeat expansion were clonally purified on MEF feeders and the resulting colonies were verified by southern blotting.

### **C9ORF72 Southern blotting**

A 241-bp digoxigenin (DIG)-labeled probe was generated from 100 ng control genomic DNA (gDNA) by PCR reaction using Q5 High-Fidelity DNA Polymerase (NEB) with primers shown in the [key resources table](#). Genomic DNA was harvested from iPSCs using cell lysis buffer (100 mM Tris-HCl pH 8.0, 50 mM EDTA, 1% w/v sodium dodecyl sulfate (SDS)) at 55 °C overnight and performing phenol:chloroform extraction. A total of 25 µg of gDNA was digested with AflIII at 37 °C overnight, run on a 0.8% agarose gel, then transferred to a positive charged nylon membrane (Roche) using suction by vacuum and UV-crosslinked at 120 mJ. The membrane was pre-hybridized in 25 ml DIG EasyHyb solution (Roche) for 3 h at 47 °C then hybridized at 47 °C overnight in a shaking incubator, followed by two 5-min washes each in 2X Standard Sodium Citrate (SSC) and in 0.1% SDS at room temperature, and two 15-min washes in 0.1x SSC and in 0.1% SDS at 68 °C. Detection of the hybridized probe DNA was carried out as described in DIG System User's Guide. CDP-Star® Chemiluminescent Substrate (Sigma-Aldrich) was used for detection and the signal was developed on X-ray film (Genesee Scientific) after 20 to 40 min.

### **Repeat primed PCR (RP-PCR)**

Genomic DNA was isolated with Qiagen DNeasy Blood & Tissue Kit (cat 69504). 100 ng of genomic DNA was amplified by PCR using primers listed in the [key resources table](#) in a 20 µl PCR reaction consisting of 0.25 mM each of 7-deaza-2-deoxyguanine triphosphate (deaza-dGTP) (NEB), dATP, dCTP and dTTP, 5% DMSO, 1x Qiagen buffer, 1x Taq polymerase (Roche), 1M betaine and 1 µM each of the three primers. During the PCR, the annealing temperature was gradually decreased from 70 °C and 56 °C in 2 °C increments with a 3-min extension time for each cycle. The PCR products were purified using the QiaQuick PCR purification kit (Qiagen) and analyzed using fragment analysis by Genewiz.

### **Retinoic acid/purmorphamine protocol for iPSC-motor neuron differentiation**

Directed differentiation of iPSC motor neurons were generated for large-scale exosome preparations as previously described with slight modifications.<sup>55</sup> On day 0, iPSCs were dissociated with Accutase (Life Technologies) into single cell suspension and 300,000 iPSCs were seeded into one well of six-well plate (pre-coated with Matrigel (Corning)) in mTeSR medium (Stem Cell Technologies) with 10 µM Rock Inhibitor (Selleck). On day 1, cultures were switched to Neural Differentiation Medium (NDM) consisting of a 1 to 1 ratio of DMEM/F12 (Corning) and Neurobasal medium (Life Technologies), 0.5x N2 (Life Technologies), 0.5x B27 (Life Technologies), 0.1 mM ascorbic acid (Sigma), 1x Glutamax (Life Technologies), 3 µM CHIR99021 (Cayman), 2 µM DMH1 (Selleck) and 2 µM SB431542 (Cayman) were also added. The medium was changed every other day. On day 7, cells were dissociated with Accutase and 4-6 million cells were seeded in a 10-cm dish (pre-coated with Matrigel) in NDM plus 1 µM CHIR99021, 2 µM DMH1, 2 µM SB431542, 0.1 µM Retinoic acid (Sigma), 0.5 µM Purmorphamine (Cayman) and 10 µM Rock Inhibitor. Rock inhibitor was removed on day 9 and the medium was changed every other day. On day 13, cells were dissociated with Accutase and 20 million cells were seeded per well in non-adhesive 6-well plates (Corning) in NDM plus 1 µM Retinoic acid, 1 µM Purmorphamine, 0.1 µM Compound E (Cayman), and 5 ng/ml each of BDNF, GDNF and CNTF (R&D Systems). Cells were used for experiments between days 25–35 of differentiation. The medium was changed every other day.

### **Isolation of extracellular vesicles with polyethylene glycol**

iPSC-derived motor neurons were cultured in non-adhesive 6-well plates. The conditioned medium was harvested 24 hours after DMSO, 3 µM apilimod, 3 µM apilimod with 10 µM GW4869, or 10 µM GW4869 treatment. The harvested medium was centrifuged at 200g for 5 minutes at room temperature to remove cell pellets. The supernatants were collected and centrifuged at 2,000g for 20 minutes at 4 °C to remove remaining cell debris.<sup>23</sup> The supernatants were collected and centrifuged at 10,000g for 30 minutes at 4 °C to remove apoptotic bodies. The obtained supernatants were supplemented with 50% w/v stock solutions of PEG 6000 (Sigma-Aldrich, 81253) and with 1M NaCl to final concentration of 8% PEG and 0.5M NaCl. Samples were mixed gently and incubated at 4 °C overnight with rotating. Extracellular vesicles were concentrated by centrifuge at 3,000g at 4 °C for an hour. Supernatants were removed and pellets were suspended with PBS. Extracellular vesicle samples were kept in -80 °C for long-term storage.

### **Conversion of iPSCs into microglia**

Microglia were generated from human iPSCs as previously described with slight modifications.<sup>56</sup> In brief, 4.5 x 10<sup>6</sup> iPSCs were seeded into a well of an Aggrewell 800 (STEMCELL Technologies) to form embryoid bodies (EBs) in EB medium consisting of mTeSR1 plus BMP4 (Gibco), 50 ng/mL VEGF (Gibco), 20 ng/mL SCF (Miltenyi Biotec), and 10 uM Rock inhibitor (Selleck). A 75% EB medium change was performed daily for 3 days without Rock inhibitor. On Day 4, EBs were transferred to a low-attachment 6-well plate (Corning) and left undisturbed for 3 days. On Day 7, EBs were transferred to a T75 flask pre-coated with growth factor reduced Matrigel (Corning) in X-VIVO15 (Lonza), supplemented with 100 ng/mL M-CSF (Biolegend), 25 ng/mL IL-3 (Biolegend), 1x Glutamax (Life Technologies), 100 U/mL penicillin and 100 µg/mL streptomycin (Gibco), and 55 uM 2-mercaptoethanol (Gibco) and left undisturbed for one week, after which fresh medium added weekly (without replacement). Macrophage precursors began



to emerge about 4 weeks, after which cells were collected weekly and flasks replaced with fresh medium. Harvested cells were plated onto plastic coverslips (Thermo Scientific) at 100,000 per cm<sup>2</sup> and differentiated into microglia-like cells for 10 days or more in differentiation medium consisting of DMEM/F12 (Corning), 1x N2 (Life Technologies), 100 U/mL penicillin and 100 µg/mL streptomycin (Gibco), 50 µM 2-mercaptoethanol (Gibco), 1x Glutamax (Life Technologies), and 100 ng/mL IL-34 (Biolegend).

#### Isolation of nuclear and cytoplasmic cell fractions for immunoblotting

Pellet samples treated with DMSO or 3 µM apilimod from control and C9ORF72 ALS/FTD patient MNs were collected. The nuclear and cytoplasmic cell fractions were prepared using the NE-PER<sup>TM</sup> Nuclear and Cytoplasmic Extraction Reagents (Thermo Scientific<sup>TM</sup>, cat. 78833) according to the manufacturer's instructions. Briefly, the cell pellets were washed once with PBS and centrifuged at 500g for 3 minutes. The cell pellet was suspended with 200 µl of cytoplasmic extraction reagent I (CER-I) by vortexing for 15 seconds. The suspension was incubated for 10 minutes on ice. 11 µl of cytoplasmic extraction reagent II (CER-II) was added to the suspension and the samples were vortexed for 5 seconds, incubated for 1 minute on ice, vortexed for another 5 seconds, and centrifuged at 16,000g for 5 minutes. The supernatant fraction (cytoplasmic extract) was transferred to a new tube and stored. The insoluble (pellet) fraction was re-suspended with 100 µl of nuclear extraction reagent (NER) and subjected to 4 cycles of vortexing for 15 seconds and incubation on ice for 10 minutes. Then samples were centrifuged at 16,000g for 10 minutes. The supernatant fraction (nuclear extract) was transferred to a new tube and stored.

#### Isolation of soluble and insoluble cell fractions for immunoblotting

Pellet samples treated with DMSO or 3 µM apilimod from control and C9ORF72 ALS/FTD patient MNs for 24 hours were collected. The soluble (detergent-soluble) and insoluble (urea-soluble) cell fractions were prepared as previously described.<sup>57</sup> The cell pellets were washed once with PBS and centrifuged at 300g for 5 minutes. The cell pellets were suspended with 200 µl of cold RIPA buffer (Santa Cruz) with a protease inhibitor cocktail (Roche) and a phosphatase inhibitor cocktail (Roche). The suspension was incubated for 10 minutes on ice and then centrifuged at 15,700g for 30 minutes at 4°C. After centrifuging, the supernatants were collected as the soluble (or detergent-soluble) fractions. The detergent-insoluble pellets were washed once with cold RIPA buffer and suspended with 200 µl of urea buffer, containing 7M urea (Sigma), 2M thiourea (Sigma), 4% (wt/vol) CHAPS (Sigma), 30 mM Tris pH8.5 (Teknova), protease inhibitor cocktail (Roche) and phosphatase inhibitor cocktail (Roche). The samples were sonicated 10 seconds for three times in a sonicator water bath, and centrifuged at 100,000g for 30 minutes at 22°C using TLA-45 rotor. The supernatants were collected to a new tube and stored as the insoluble (or urea-soluble) fractions.

#### Negative staining transmission electron microscopy

Exosomes were fixed with 4% formaldehyde in PBS. 200-mesh, formvar-filmed copper grids were plasma-treated (Harrick PDC-001; Harrick Plasma, Ithaca, NY) and immediately incubated on a drop of exosomal vesicles for ten minutes. The grid was blotted and immediately placed on a drop of 1% phosphotungstic acid at pH7 for one minute, blotted again and air dried. The samples were imaged in a Zeiss EM10 transmission electron microscope (Zeiss, White Plains, NY) at 80 kV using a Gatan Erlangshen CCD camera (Gatan, Pleasanton, CA).

#### Correlative Light Electron Microscopy (CL-EM)

C9-ALS1 iMNs were converted on 15 mm Nunc<sup>TM</sup> Thermanox<sup>TM</sup> Coverslips (Thermo Scientific<sup>TM</sup>, 174969) pre-coated with 0.1% gelatin (1 hour, room temperature) and laminin (4°C, overnight). iMNs were treated with 3 µM apilimod for 24 hours and fixed with 4% paraformaldehyde (PFA) in 0.25 M HEPES for 1 hour at room temperature, washed with 0.1 M HEPES for three times, permeabilized with 0.1% Triton-X in 0.1 M HEPES for 10 mins at room temperature, blocked with 10% donkey serum in 1% BSA in 0.1 M HEPES for two hours at room temperature, and incubated with primary antibodies (diluted in 1% BSA in 0.1M HEPES) at 4°C overnight. The following antibodies were used: rabbit anti-TDP-43 Polyclonal antibody (Proteintech, 10782-2-AP, 1:200) and mouse anti-CD63 Monoclonal antibody (BD Transduction Laboratories, 556019, 1:100). Cells were then washed with 0.1M HEPES with 0.1% Triton-X for three times and incubated with Alexa Fluor-conjugated secondary antibodies (Life Technologies) (diluted in 1% BSA in 0.1M HEPES) for 2 hours at room temperature. iMNs were stained with DAPI (Life Technologies) for 10 minutes at room temperature and then mounted on slides with Vectashield (Vector Labs). Neuronal area was determined on the basis of the *Hb9::RFP* signals. High magnification images and z-stack images (fluorescence images) were acquired on an LSM 800 confocal microscope with oil immersion at 63x (Zeiss). Low magnification images (10–20X) were acquired with light microscopy to locate the cell of interest. Cells were fixed with glutaraldehyde and osmium tetroxide, stained with uranyl acetate, dehydrated through graded ethanol, infiltrated with epoxy resin. After polymerizing the block for 18 hours, the cells were separated from the Thermanox coverslip to the block and sectioned into 500 nm sections. The sections were mounted on TEM grid and examined with a transmission electron microscope.

#### Mass spectrometry analysis of exosomes

10 µg of each sample was reduced for 10 minutes at 50°C with 10 mM DTT and alkylated for 30 minutes at room temperature in the dark with 15 mM iodoacetamide. Proteins were precipitated with 8 volumes of ice-cold acetone and 1 volume of ice-cold methanol 2h at -80°C. The pellets were washed 3 times with 250 µl of cold methanol. Protein pellets were resolubilized in 100 µl of 50 mM Tris pH 8.0 and pre-digested with 0.3 µg of Trypsin/LysC for 3 hours at 37°C with agitation. Another 0.3 µg of Trypsin/LysC was added



to the proteins and the digestion was continued overnight at 37°C with agitation. Samples were then acidified with 2% formic acid and the peptides were purified by Strata-X reversed phase SPE (Phenomenex).

Acquisition was performed with a ABSciex TripleTOF 6600 (ABSciex, Foster City, CA, USA) equipped with an electrospray interface with a 25  $\mu$ m iD capillary and coupled to an Eksigent  $\mu$ UHPLC (Eksigent, Redwood City, CA, USA). Analyst TF 1.8 software was used to control the instrument and for data processing and acquisition. The samples were analyzed in SWATH acquisition mode (data independent acquisition). The source voltage was set to 5.5 kV and maintained at 325°C, curtain gas was set at 45 psi, gas one at 25 psi and gas two at 25 psi. Separation was performed on a reversed phase Kinetex XB column 0.3  $\mu$ m i.d., 2.6  $\mu$ m particles, 150mm long (Phenomenex) which was maintained at 60°C. Samples were injected by loop overfilling into a 5  $\mu$ L loop. For the 45 minutes LC gradient, the mobile phase consisted of the following solvent A (0.2% v/v formic acid and 3% DMSO v/v in water) and solvent B (0.2% v/v formic acid and 3% DMSO in EtOH) at a flow rate of 3  $\mu$ L/min.

SWATH samples were processed using a publicly available ion library (SWATH Atlas). Retention times in the library were recalibrated using known abundant proteins and keratins. Proteins were quantified using a maximum of 10 peptides per protein and 4 MS/MS transitions per peptide. The quantification for a protein represents the sum of the area under the curve (AUC) of all the integrated peptides (max 10) for this protein. The intensity of each sample was normalized against every other sample using the total signal. GO analyses have been done using the Reactome FI plugin of Cytoscape, with only the significant genes.

### Western blotting

iPSC-derived motor neurons were collected in RIPA buffer (Santa Cruz) with a protease inhibitor cocktail (Roche). Protein quantity from cell lysates and exosomal samples was measured by the BCA assay (Pierce). Samples were run on a 10% SDS gel, and the proteins were transferred onto Immobilon-FL PVDF Membrane (Millipore). The total protein for each sample was stained by Revert™ 700 Total Protein Stain Kits (LI-COR Biosciences) and quantified. The membrane was blocked with Intercept (TBS) Blocking buffer (LI-COR Biosciences), incubated with primary antibodies overnight at 4°C, washed three times with 0.1% TBS-T, and then incubated with IRDye® 680RD Donkey anti-Rabbit IgG Secondary Antibody or IRDye® 800RD Donkey anti-Mouse IgG Secondary Antibody (LI-COR Biosciences). After washing with 0.1% TBS-T for three times and TBS once, blots were scanned using LI-COR Odyssey CLx imaging system. The following primary antibodies were used: mouse anti-TSG101 (Biosciences, 612697, 1:500), mouse anti-TUJ1 (Biolegend, MMS-435P, 1:2000), rabbit anti-LC3 (MBL, PM036, 1:500), mouse anti-p62 (Biosciences, 610832, 1:1000), rabbit anti-OPTN Polyclonal antibody (Proteintech, 10837-1-AP, 1:500), rabbit anti-phospho-TDP43 (Ser409/410) Polyclonal antibody (Proteintech, 22309-1-AP, 1:500), Rabbit anti-PIP5K3 Polyclonal antibody (Proteintech, 13361-1-AP, 1:300), rabbit anti-TDP-43 Polyclonal antibody (Proteintech, 10782-2-AP, 1:500), mouse anti-poly(GA) Monoclonal antibody (MABN889, Millipore, 1:1000), rabbit anti-TFEB Polyclonal antibody (Bethyl Laboratories, A303-673A, 1:500), rabbit anti-phospho-TFEB (Ser142) Polyclonal antibody (Sigma-Aldrich, ABE1971-I-25UL, 1:500), rabbit anti-Huntingtin Monoclonal antibody (abcam, ab109115, 1:500), rabbit anti-alpha-Synuclein Polyclonal Antibody (Invitrogen, PA5-85791, 1:500), rabbit anti-Fibrillarin Monoclonal antibody (abcam, ab166630, 1:1000) and rabbit anti-HSP90 Polyclonal antibody (Proteintech, 13171-1-AP, 1:1000). The list of antibodies was provided in the [key resources table](#). The full blots for all experiments are shown in Data S1: The regions shown in the main and supplementary figures were highlighted in red for each blot image. Some membranes were cut before immunoblotting in order to enable blotting with multiple antibodies on the same membranes. Cutting sites are labeled with blue lines. Mr = Odyssey® One-Color Protein Molecular Weight Marker (Licor); Mr Duo = Chameleon® Duo Pre-stained Protein Ladder (Licor).

The quantified values were calculated as the relative intensity of indicated proteins normalized to total protein per sample unless otherwise mentioned. For [Figures 3C, 3G, 3I, 3K, S3K, and S3O](#), n=9 independent conversions/condition from three CTRL and three C9-ALS/FTD patients (n=3 per line). For [Figures 3L and S3N](#), n=6 independent conversions/condition from two CTRL and two C9-ALS/FTD patients (n=3 per line). For [Figures 3N, 3O, S3Q, and S3R](#), n=11 independent conversions/condition from three CTRL lines and one corrected isogenic line (n=3 for CTRL2, CTRL3 and C9-ALS/FTD1 corrected isogenic line and n=2 for CTRL5) and n=13 from three C9-ALS/FTD patients (n=6 for C9-ALS/FTD1, n=4 for C9-ALS/FTD2 and n=3 for C9-ALS/FTD3). For [Figures 3Q and S3T](#), n=6 independent conversions/condition from two CTRL (n=3 per line) and n=8 from independent conversions/condition from three C9-ALS/FTD patients (n=2 for C9-ALS/FTD1 and n=3 for C9-ALS/FTD2 and C9-ALS/FTD3). For [Figures 3U and 3V](#), n=9 independent conversions/condition from three CTRL lines and one corrected isogenic line (n=3 for CTRL2 and n=2 for CTRL3, CTRL5 and C9-ALS/FTD1 corrected isogenic line) and n=8 from independent conversions/condition from three C9-ALS/FTD patients (n=2 for C9-ALS/FTD1 and n=3 for C9-ALS/FTD2 and C9-ALS/FTD3). For [Figure S3D](#), n=8 biological replicates (independent conversions) per condition from three CTRL lines (n=3 for CTRL1 and CTRL2, n=2 for CTRL3) and n=8 biological replicates (independent conversions) per condition from three C9-ALS/FTD patients (n=3 for C9-ALS/FTD1 and C9-ALS/FTD3, n=2 for C9-ALS/FTD2). For [Figure S3F](#), n=6 biological replicates (independent conversions) per condition from two CTRL lines and n=9 biological replicates (independent conversions) per condition from three C9-ALS/FTD patients (n=3 per line). For [Figure S3V](#), n=7 independent conversions/condition from two CTRL and one C9-ALS/FTD1 corrected isogenic line (n=3 for CTRL2 and C9-ALS/FTD1 corrected isogenic line and n=2 for CTRL5) and n=9 from independent conversions/condition from three C9-ALS/FTD patients (n=5 for C9-ALS/FTD2 and n=2 for C9-ALS/FTD1 and C9-ALS/FTD3). For [Figure 4D](#), n=4 independent iMN conversions/condition from two CTRL (n=2 for CTRL2 and CTRL3) and n=7 independent conversions/condition from three C9-ALS/FTD lines (n=3 for C9-ALS/FTD1 and C9-ALS/FTD2, and n=1 for C9-ALS/FTD3). For [Figure 4F](#), n=7 independent conversions/condition from three C9-ALS/FTD patients (n=3 for C9-ALS/FTD1 and C9-ALS/FTD2, and n=1 for C9-ALS/FTD3).



### Dipeptide repeat protein expression in iMNs and FACS analysis of GFP+/CFSE+ exosomes

iMNs were generated as described.<sup>55</sup> At day 25 of differentiation, we performed lentiviral transduction of GP(50)-GFP or GFP using previously-published constructs.<sup>58</sup> 3 days after lentiviral transduction of 5–6 embryoid bodies per sample, iMNs were treated with DMSO or 3  $\mu$ M apilimod for 24-hours and supernatant was harvested. 1  $\mu$ l of CFSE dye (Thermo, catalog number: C34554) was added to the supernatant and incubated for 20 minutes at room temperature with gentle agitation. GFP+/CFSE+ exosomes were quantified using flow-cytometry gated for particles smaller than cells. n=3 independent differentiations/condition.

### Poly(GR) Immunoassay

Poly(GR) levels in concentrated exosome and pellet samples from control and C9ORF72 ALS/FTD patient MNs were measured in a blinded manner using Meso Scale Discovery (MSD) platform based Poly(GR) sandwich immunoassay with custom made affinity purified rabbit polyclonal GR antibodies as previously described with minor modifications.<sup>27,28</sup> Biotinylated poly(GR) antibodies at 0.5  $\mu$ g/ml were coated on MSD Gold 96-well single spot streptavidin plates and incubated overnight at 4°C. After washing and blocking, concentrated exosome samples (40  $\mu$ l) were loaded on plates in duplicate wells and incubated for 1.5 hrs at room temperature on a shaking platform. After 3-time washes, plates were loaded with 0.5  $\mu$ g/ml MSD-Gold-Sulfo-tagged poly(GR) detection antibody and incubated for 1 hour at room temperature on a shaking platform followed by three final washes. After adding 1X MSD-Read buffer, plates were immediately read using MSD-QuickPlex SQ 120 reader and data presented as raw electrochemiluminescence (ECL) signals detected from the samples.

For Figure 3R, n=11 biological replicates (independent conversions)/condition from three CTRL lines (n=3 for CTRL2 and n=4 for CTRL1 and CTRL3) and n=7 independent conversions/condition from two C9-ALS/FTD lines (n=3 for C9-ALS/FTD1 and n=4 for C9-ALS/FTD2). For Figure 3S, n=6 biological replicates (independent conversions)/condition from two CTRL lines (n=3 for CTRL2 and CTRL3) and n=9 independent conversions/condition from three C9-ALS/FTD lines (n=3 for C9-ALS/FTD1, C9-ALS/FTD2 and C9-ALS/FTD3). For 3T, n=7 independent conversions/condition from two C9-ALS/FTD patients, n=3 for C9-ALS/FTD1 and n=4 for C9-ALS/FTD2).

### mRFP-GFP-LC3 assay

Retrovirus encoded by pMXs-mRFP-GFP-LC3 was transduced into iMN cultures 24 hours before transduction with iMN reprogramming factors. On day 17, iMNs were treated with DMSO or 3  $\mu$ M apilimod for 24 hours and then fixed with 4% paraformaldehyde at 4°C for 1 hour. The cultures were immunostained with anti-MAP2 antibody to detect motor neurons. Coverslips were imaged on a Zeiss LSM 800 confocal microscope. Quantification was performed using ImageJ.

### RNA-seq and pathway analysis

iPSC-derived motor neurons from C9-ALS/FTD patients (C9-ALS/FTD1) were cultured with DMSO or 3  $\mu$ M apilimod for 24 hours and collected. 3'-Digital gene expression RNA-Seq of all samples was performed by Amaryllis Nucleics. Briefly, mRNA was extracted using the NEBNext Poly(A) mRNA Magnetic Isolation Module according to the manufacturer's instructions. 3' RNA-Seq libraries were generated using the 3'-Digital Gene Expression RNAseq Library Kit (Amaryllis Nucleics). Libraries were sequenced on an Illumina NextSeq 500 sequencer. A total of 10 million to 25 million 80-bp, single-end reads were obtained for each sample. Reads were aligned to the Hg38 transcriptome using HISAT2.<sup>59</sup> A count table was obtained using FeatureCounts with strand specificity enabled. Differential expression analysis was performed using DESeq2.<sup>60,61</sup> The list of differentially expressed genes can be found in the Table S2.

KEGG pathway enrichment analysis, transcription factor enrichment analysis (TFEA) and Bioplanet analysis were performed by providing Enrichr with a list of all genes identified by DESeq2 as having a greater than 95% chance of being differentially expressed.<sup>54</sup>

iPSC-derived motor neurons from C9-ALS/FTD patients (C9-ALS/FTD1 and C9-ALS/FTD3) were cultured with 10  $\mu$ M NC ASO or PIKFYVE ASO for 96 hours and collected. mRNA sequencing was performed by Novogene Co., Ltd. (US). Briefly, total RNA was extracted using RNeasy Plus Mini Kit (Qiagen) according to manufacturer's instructions. Total RNA was then shipped to Novogene Co., Ltd for poly(A) mRNA enrichment and library preparation. Libraries were sequenced on Illumina NovaSeq 6000. A total of 20 million to 40 million 150-bp, paired-end reads were obtained for samples sent. Reads were aligned to the Hg38 transcriptome using STAR.<sup>62</sup> Aligned reads were then quantified using Partek E/M with Partek® Flow® software, v10.0 (Partek Inc.). Gene counts were normalized using median ratio with offset by one before performing differential expression analysis using DESeq2.<sup>61,63</sup>

### Immunocytochemistry

iMNs were fixed in 4% paraformaldehyde (PFA) for 1 hour at 4°C, permeabilized with 0.1% Triton-X for 10 mins at room temperature, blocked with 10% donkey serum in 1% BSA in PBS for two hours at room temperature, incubated with primary antibodies at 4°C overnight. Cells were then washed with 0.1% PBS-T for three times and incubated with Alexa Fluor-conjugated secondary antibodies (Life Technologies) for 2 hours at room temperature. iMNs were stained with DAPI (Life Technologies) for 10 minutes at room temperature and then mounted on slides with Vectashield (Vector Labs). Neuronal area was determined by manual outlining in ImageJ on the basis of the *Hb9::RFP* signals. Images were acquired on an LSM 800 confocal microscope with oil immersion at 63x (Zeiss). iMNs were quantified from two independent conversion/group for each experiment. The following primary antibodies were used: rabbit anti-GR repeat Polyclonal antibody (Proteintech, 23978-1-AP, 1:50), rabbit anti-PR repeat Polyclonal antibody



(Proteintech, 23979-1-AP, 1:50), rabbit anti-TDP-43 Polyclonal antibody (Proteintech, 10782-2-AP, 1:200) and chicken anti-MAP2 antibody (Abcam, ab5392, 1:5000), rabbit anti-GA repeat Polyclonal antibody (Proteintech, 24492-1-AP, 1:50), rabbit anti-TFEB Polyclonal antibody (Bethyl Laboratories, A303-673A, 1:100), rabbit anti-LC3B Monoclonal antibody (Abcam, ab192890, 1:100), mouse anti-LAMP1 Monoclonal antibody (Abcam, ab25630, 1:20), mouse anti-CD63 Monoclonal antibody (BD Transduction Laboratories, 556019, 1:100), rabbit anti-RAB7 Monoclonal antibody (Abcam, ab137029, 1:100), rabbit anti-phospho TDP-43 (pS409/410) Polyclonal antibody (Cosmo Bio, CAC-TIP-PTD-P07, 1:200), goat anti-FUS Polyclonal antibody (Bethyl Laboratories, A303-839A, 1:200) and goat anti-IBA1 Polyclonal antibody (Abcam, ab5076, 1:100). The following dyes and conjugated antibodies were used: LysoSensor™ Green DND-189 (Invitrogen™, L7535, 5  $\mu$ M), LysoTracker™ Deep Red (Invitrogen™, L12492, 100 nM), Alexa Fluor® 647 Anti-LC3B antibody (Abcam, ab225383, 1:100). The list of antibodies was provided in the [key resources table](#).

To quantify the number of punctae in iMNs and MNs, the number of punctae per area ( $\mu\text{m}^2$ ) was measured with Image J using the find maxima detection tool with noise tolerance=50–100 (depending on the antibodies). To quantify the TDP43, TFEB and FUS signals in iMNs, the fluorescence intensity of the TDP43, TFEB or FUS staining in the cell nucleus and cytoplasm were measured using Image J and recorded as the nuclear to cytoplasmic ratio. To quantify the number of colocalized punctae in MNs, the number of colocalized punctae per area ( $\mu\text{m}^2$ ) was measured with Image J using the spots colocalization (ComDet) tool. To quantify the endogenous LAMP1+ punctae in iMNs, the size of LAMP1+ vesicles < 0.81  $\mu\text{m}^2$  were defined as small LAMP1+ vesicles; while vesicles > 0.81  $\mu\text{m}^2$  were defined as large LAMP1+ vesicles. 75% of LAMP1+ vesicles were < 0.81  $\mu\text{m}^2$  in C9-ALS /FTD iMNs treated with apilimod.

### Quantitative real time PCR

Total RNA was extracted from HeLa cells, iPSC-derived motor neurons or mice brain tissue with RNA Extraction Kit (Life Technologies) and reverse transcribed with an Oligo dT primer using a ProtoScript II First Strand Synthesis Kit (NEB). RNA integrity was checked using the NanoDrop 1000 (Thermo). Real-time PCR was performed with iTaq Universal SYBR Green Supermix (Bio-Rad) using primers shown in the [key resources table](#).

### C. elegans paralysis assay on solid media

Standard methods for culturing and handling the worms were used.<sup>45,64</sup> Briefly, 40 age-synchronized L4 worms were transferred to NGM plates and scored daily for paralysis, from day 1 to day 12 of adulthood. Animals were counted as paralyzed if they failed to move upon prodding with a worm pick. Worms were scored as dead if they failed to move their head after being prodded on the nose and showed no pharyngeal pumping. All experiments were conducted at 20 °C and in triplicate, three times. Some experiments were conducted by dissolving apilimod (1 M) into the NGM plates.

### C. elegans neurodegeneration assay

For scoring of neuronal processes for gaps or breakage, worms were selected at day 9 of adulthood for visualization of motor neurons in vivo. Animals were immobilized in 5 mM levamisole dissolved in M9 and mounted on slides with 2% agarose pads. Visualization was done using a Zeiss Axio Imager M2 microscope, using a 10X objective and a 1.0 Optovar. The software used was Zen 2.3 Lite. At least one hundred worms were scored per condition, over 8 distinct experiments.

### Drosophila larval turning assay

All stocks and crosses were maintained on standard yeast/cornmeal/ molasses food (25.0 °C using) a 12 hours light and dark cycle unless otherwise noted. The following stocks were used: (i) GAL4 motor neuron driver D42-GAL4 was used to drive the expression of UAS transgenes using the GAL4-UAS bipartite expression system, (ii)  $w^{1118}$  as used as a genetic background control for TDP-43 transgenes, (iii)  $w^{1118}$ ; UAS-TDP-43<sup>G298S</sup>-YFP used to drive TDP-43 expression in motor neurons was previously described.<sup>65</sup> (iv)  $y[1]v[1]$ ; P{Y[+(7.7) = GSYP]attP2 (Bloomington Drosophila Stock Center (BDSC # 8622)) was used as genetic background control for PIKFYVE RNAi.  $y[1]sc[+]v[1]$ ; P{y[+t7.7]v[+t1.8]=TRiP.GL00246}attP2, used to knock-down PIKFYVE by RNAi was obtained from the Bloomington Drosophila Stock Center (BDSC # 35793). To model C9ORF72 we used  $w^{1118}$ ; P{UAS-poly-GR.PO-100}attP40 (BDSC#58696) with  $y1w^{1118}$ ; attP40 as genetic background control. RNAi experiments were performed with chronic RNAi expression and *Pikfyve* suppression. For apilimod experiments, larvae were fed yeast mixed with either solvent (DMSO) or apilimod (10  $\mu$ M) for 5 days throughout larval development in a chronic treatment paradigm.

Larval turning assays were performed using third instar larvae as previously described.<sup>66</sup> Briefly, third instar larvae were placed on a Petri dish filled with a solidified grape juice/agar medium and allowed to accommodate for a few seconds. Larvae were gently turned ventral side up with a clean paintbrush and the time it took for each larva to turn ventral side down and make the first forward movement was recorded. 33 larvae per genotype, per condition were screened for YFP and evaluated for locomotor function. Larval turning data was analyzed with Graphpad Prism9 for statistical significance using Kruskal-Wallis with multiple comparisons.

### Drosophila C9ORF72 GR.100 NMJ assay

*Drosophila* larvae were reared and immunostained as described.<sup>16</sup> Wild type ( $w^{1118}$ ) or GR.100 larvae ( $w$ ;OK6-Gal4/UAS-100xGR) were reared on individual apple plates at 25 °C and fed yeast mixed with either solvent (DMSO) or apilimod (30  $\mu$ M) for 5 days throughout larval development. Briefly, third-instar larvae were dissected in saline and immunostained with anti-BRP (nc82;



Developmental Studies Hybridoma Bank; 1:100) and anti-HRP (Cyanine 3 (Cy3) conjugated; Jackson ImmunoResearch; 1:400). Alexa Fluor 488-conjugated mouse secondary antibody was used at 1:400 (Jackson ImmunoResearch). NMJs were imaged using a Nikon A1R Resonant Scanning Confocal microscope with a 100x APO 1.4 NA oil immersion objective. BRP puncta were quantified using Nikon Elements Software and data was analyzed using Graphpad Prism.

### TDP-43 mice

TDP-43 transgenic mice were purchased from the Jackson Laboratory (stock no: 012836). The mouse Thy1 promoter drives the expression of the human TDP-43 gene in neurons. TDP-43<sup>Tg/Tg</sup> homozygous mutant mice were generated by crossing TDP-43<sup>Tg/+</sup> heterozygous mice with TDP-43<sup>Tg/+</sup> heterozygous mice. Mice were genotyped using primers listed in the [key resources table](#). Homozygous mutants were identified by a single band at 500 bp, heterozygotes were identified by bands at 303 bp and 500 bp, and wild types were identified by a single band at 303 bp. Homozygous mutants and their corresponding WT littermate controls were used in the experiments.

### ASO Treatment in TDP-43 mice

25  $\mu$ g of negative control or *Pikfyve* ASO was administered by intracerebroventricular injection at P1. The ASOs were synthesized by Integrated DNA Technologies (IDT) and contained chemical modifications designed to bolster their efficacy for inhibition of gene expression. Phosphorothioate bonds were added to the sequence to provide protection from degradation by external nucleases and the modified base 2'-O-methoxy-ethyl (MOE) was also added to facilitate increased nuclease stability and binding affinity of the ASO to the mRNA target of interest. To enhance the purity of the ASOs for the *in vivo* studies, the ASOs were HPLC purified followed by a sodium/salt exchange. Finally, ethanol precipitation was performed to remove any residual chemical impurities. The sequences of the ASOs were listed in the [key resources table](#) (m=2'-O-Methyl, \*-phosphorothioate linkage, MOE = 2'-Methoxyethyl).

P1 neonatal mice were anesthetized by placing them on a paper towel lined bucket of ice to induce hypothermia. The injection site targeting the lateral ventricles was defined as the distance of 40% between the lambda (anatomical landmark) and the eye. To perform the intracerebroventricular (ICV) injection, a 900 series Hamilton syringe (10  $\mu$ L) was adapted with the bottom portion of Neuros model 1701 RN syringe. A 33-gauge needle was inserted 2mm deep perpendicular to the skull surface into the left lateral ventricle. Mice received 3  $\mu$ L of negative control ASO or *Pikfyve* ASO in PBS (a total of 25  $\mu$ g). Within each litter half of the pups received negative control ASO and the other half received *Pikfyve* ASO. Mice were placed under a heat lamp in a new cage containing soiled bedding from the parental cage and were closely monitored until they had fully recovered. Beginning at P5, mice were administered vehicle (3.75% DMSO in 0.9% normal saline) or 6.7  $\mu$ g/kg GW4869 by intraperitoneal (IP) injection every other day throughout the remainder of the experiment.

### Genetic deletion of *Pikfyve* in TDP-43 mice

For experiments using genetic deletion of *Pikfyve*, B6.Cg-PIKFYVE<sup>tm1.1Ashi</sup>/J were purchased from the Jackson Laboratory (stock no: 029331). The *Pikfyve*-flox mice were established by inserting loxP sites flanking *Pikfyve* exon 6 (encoding the FYVE finger domain). Removal of the floxed sequence creates a null allele. Mice heterozygous for the floxed allele (*Pikfyve*<sup>+/<sup>flox</sup></sup>) were bred to germline expressing B6.C-Tg(CMV-Cre)1Cgn/J mice (Jackson lab stock no: 006054) to generate *Pikfyve* mice with a single allele knockout (*Pikfyve*<sup>+/-</sup>). The *Pikfyve*<sup>+/-</sup> mice were crossed with TDP-43<sup>Tg/+</sup> to generate *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/+</sup>. The resulting offspring were then crossed with TDP-43<sup>Tg/+</sup> mice to generate *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/Tg</sup> mice. Mice were genotyped using several primer pairs according to a previous study (Ikonomov et al., Journal of Biological Chemistry, 2011) and listed in the [key resources table](#): S0 and A1 generating products of 1232 bp, 1476 bp or 574 bp, for the wild type allele, the floxed allele or the knockout allele, respectively. S1 and A1, generating products of 1063 bp, 1308 bp or 405 bp, for the wild type, the floxed allele or the knockout alleles, respectively. S2 and A1, generating 663 bp product, 852 bp product or no product, for the wild type, the floxed allele or the knockout alleles, respectively; and S3 and A1, generating 208 bp product, 378 bp product or no product, for the wild type allele, the floxed allele or the knockout allele, respectively.

### Gait impairment, kyphosis, and tremor analysis

Gait impairment, kyphosis, and tremor were examined starting at 14 days of age (first signs of abnormal movement) and continued until 22 days of age. Phenotype scoring was assessed according to a composite rubric previously reported.<sup>67</sup> A score of 0 indicates no phenotype and 4 indicates the most severe phenotype. Statistical significance was calculated by comparing the score at each time point. All investigators were blinded to the genotype and treatment of each experimental group. The humane euthanasia endpoint for the study occurred when the mouse was no longer able to right itself. Hydrogel, powdered pellets, and high caloric diet gel were placed on the bottom of each cage containing TDP-43 mutant mice. The animals were housed in cages under a temperature and humidity-controlled environment and subjected to a standard 12 h light/dark cycle with food and water available *ad libitum*. All procedures were approved by the Institutional Animal Care and Use Committee of the University of Southern California in accordance with guidelines of the National Institutes of Health.

### Cerebrospinal fluid (CSF) Collection

P1 wild type C57BL/6J mice (Jackson laboratory stock number: 000664) were injected by ICV injection with negative control ASO (25  $\mu$ g) or PIKFYVE ASO (25  $\mu$ g). Beginning at P3, mice were administered vehicle (3.75% DMSO in 0.9% normal saline) or



6.7  $\mu\text{g/kg}$  GW4869 by intraperitoneal (IP) injection every other day until P7. CSF was collected on P7, 3 hours after the final IP injection. The neonatal mice were placed on a bucket of ice lined with paper towels and were anesthetized by hypothermia. The skull was exposed by a large incision through the flap of skin on top of the scalp. The site of the lateral ventricles was defined as the distance of 40% between the lambda (anatomical landmark) and the eye. The 33-gauge needle was inserted 2 mm deep perpendicular to the skull surface and clear CSF fluid was aspirated.

For the analysis of CSF by mass spectrometry after vehicle (5% Tween 80, 10% N-methyl-2 pyrrolidone, 85% 30 mM citrate buffer in water, pH 3.1) or apilimod treatment, 6-8 week-old female mice were anesthetized and fur was removed from the dorsal lumbar area, extending from L1 to the sacrum. Ophthalmic ointment was placed on the eyes to prevent drying of the cornea. The injection site was disinfected with 70% ethanol using a sterile swab. Using a fingernail, the groove between L5-L6 was located and the needle was inserted at 90° to the spine on the midline and into the groove. The substance (vehicle or apilimod) was slowly injected and the needle left in place for 5-10 sec prior to withdrawal. 10  $\mu\text{L}$  of vehicle or 50  $\mu\text{M}$  apilimod using a 25  $\mu\text{L}$  Hamilton syringe with a 30-32 gauge needle. For CSF collection, mice were anesthetized with tribromoethanol 4 hours after vehicle or apilimod administration. When the plane of anesthesia was reached, CSF was collected through the cisterna magna ( $\sim 1\sim 3$   $\mu\text{L}$ / mouse). All samples were flash frozen on dry ice and then stored in  $-80^\circ\text{C}$  until shipped to the study sponsor.

### Mouse tissue collection

Wildtype or TDP43 Tg/Tg mice (postnatal day1) were treated with 25  $\mu\text{g}$  negative control ASO or *Pikfyve* ASO via intracerebroventricular injection at postnatal day 1. For tissue collection, P19-P21 mice were anesthetized and intracardiac perfusion was performed with cold PBS. Brains were separated into two hemispheres sagittally. One hemisphere was frozen at  $-80^\circ\text{C}$  for immunoblotting or RNA analysis. The other brain hemisphere and spinal cords were fixed in cold 4% PFA overnight prior to being cryoprotected in 30% sucrose in PBS+0.01% sodium azide in preparation for immunofluorescence analysis. For CSF collection, ASO injected mice were euthanized at P7.

### Histology

Cryoprotected tissues were embedded in O.C.T compound (Fisher Scientific, catalog number 23-730-571) and cryosectioned to 16  $\mu\text{m}$  slices using Leica CM3050S cryostat. Sections were stored at  $-80^\circ\text{C}$ . For immunofluorescence staining with primary antibodies, slides were treated with citrate buffer (pH 6)/0.05% tween-20 for 20 minutes at  $80^\circ\text{C}$  prior to antibody staining. All primary antibody stainings were done with 3 overnight incubations at  $4^\circ\text{C}$ . Secondary antibody incubation was done at room temperature in the dark for one hour. The list of antibodies was provided in the [key resources table](#). For NeuroTrace (Fluorescent Nissl Stain, Thermo Fisher Scientific, N21480) staining, the slides were rehydrated in 10x PBS for at least 40 minutes and washed in 1x PBS/0.1% Triton X-100 for 10 minutes. The slides were washed in 1x PBS 2 times for 5 minutes each and then incubated with diluted NeuroTrace (1:50 in PBS) in the dark for 20 minutes. Slides were washed for 10 minutes in PBS/0.1% Triton X-100, 2 x 5 minutes in PBS and 2 hours at room temperature or overnight at  $4^\circ\text{C}$  in PBS prior to mounting. To quantify the ventral motor neurons in the lumbar spinal cord, 4-5 sections per animal were stained with NeuroTrace and imaged using a Zeiss LSM 880 at 10x. Both lateral motor column and medial motor column motor neurons in each half spinal cord were quantified and the average total motor neuron counts per hemicord was recorded for individual animals.

To quantify the mislocalization of TDP43 into the cytoplasm, the nuclear:cytoplasmic ratio was measured using a total TDP43 antibody (10782-2-AP, Proteintech, 1:400). The fluorescence intensity of the TDP43 staining in the cell nucleus and cytoplasm were measured using Image J and recorded as the nuclear to cytoplasmic ratio. Motor neurons in the ventral spinal cord were imaged using Zeiss LSM 880 with oil immersion at 63x. 15-25 neurons were quantified per animal. To quantify the phosphorylated and aggregated TDP43+ punctae, brain and spinal cord sections were stained with phospho(Ser403/404)-TDP43 antibody (Proteintech, 66079-1-Ig, 1:100) or phospho(Ser409/410)-TDP43 antibody (Cosmo Bio, CAC-TIP-PTD-P07, 1:200) and imaged at 40x with either a Zeiss LSM800 (spinal cord) or LSM880 (brain). The number of phosphorylated-TDP43+ punctae per cytosolic area was measured with Image J using the find maxima detection tool with noise tolerance=70. Neurons and its cell area were identified using Tuj1 antibody staining (Neuromics, CH23005, 1:100) and cell nucleus was identified using DAPI. Only the cytosolic area in neurons were quantified. 15-25 neurons were quantified per animal. To quantify the number of colocalized punctae, spinal cord sections were stained with rabbit anti-LAMP1 (Abcam, ab24170, 1:100), mouse anti-RAB7 (Santa Cruz, sc-376362, 1:100) and mouse anti-LC3B (GeneTex, GT3612, 1:100) and images at 40x or 64x with a Zeiss LSM800. The number of colocalized punctae per area ( $\mu\text{m}^2$ ) was measured with Image J using the spots colocalization (ComDet) tool.

For glial analysis in the spinal cord, numbers of GFAP+ astrocytes (Abcam, ab7260, 1:100) or IBA1+ microglia (GeneTex, GT101495, 1:100) in a  $150\mu\text{m} \times 150\mu\text{m}$  area of the ventral horn were quantified manually in Image J. The cell density in each ventral horn section was recorded and 4-6 ventral horn sections were quantified for each animal. The average cell density from each animal was used for statistical analysis.

### AAV-(GR)<sub>100</sub> mouse model

The AAV9-CMV-eGFP-rBG and AAV9-CMV-eGFP-hGR<sub>100</sub>-rBG were prepared for generating viruses as previously described.<sup>49</sup> Intracerebroventricular injection of 2  $\mu\text{L}$  ( $1 \times 10^{10}$  genomes  $\mu\text{L}^{-1}$ ) of AAV-eGFP or AA-VeGFP-(GR)100 virus were performed in postnatal day1 (P1) C57BL/6J pups. Postnatal day 4 (P4) mouse pups received 25  $\mu\text{g}$  of NC or *Pikfyve* ASO by intracerebroventricular



injection. Mice were placed under a heat lamp in a new cage containing soiled bedding from the parental cage and were closely monitored until they had fully recovered. The humane euthanasia endpoint for the study occurred when the mouse was no longer able to right itself. The animals were housed in cages under a temperature and humidity-controlled environment and subjected to a standard 12 h light/dark cycle with food and water available *ad libitum*. All procedures were approved by the Institutional Animal Care and Use Committee of the University of Southern California in accordance with guidelines of the National Institutes of Health. The brain sections were stained with anti-GFP Polyclonal antibody (Aves Lab, GFP-1010, 1:100) and rabbit anti-GR repeat Polyclonal antibody (Proteintech, 23978-1-AP, 1:50). The spinal cord sections were stained with NeuroTrace (Fluorescent Nissl Stain, Thermo Fisher Scientific, N21480).

### Hindlimb clasping test

Hindlimb clasping test was examined starting 14 days of age (first signs of abnormal movement) and continued until 26 days of age. Mice were suspended by the tail and hindlimb clasping was observed for 30 seconds. Phenotypic scoring was assessed according to a composite rubric previously reported.<sup>68</sup> All investigators were blinded to the genotype and treatment of each experimental group.

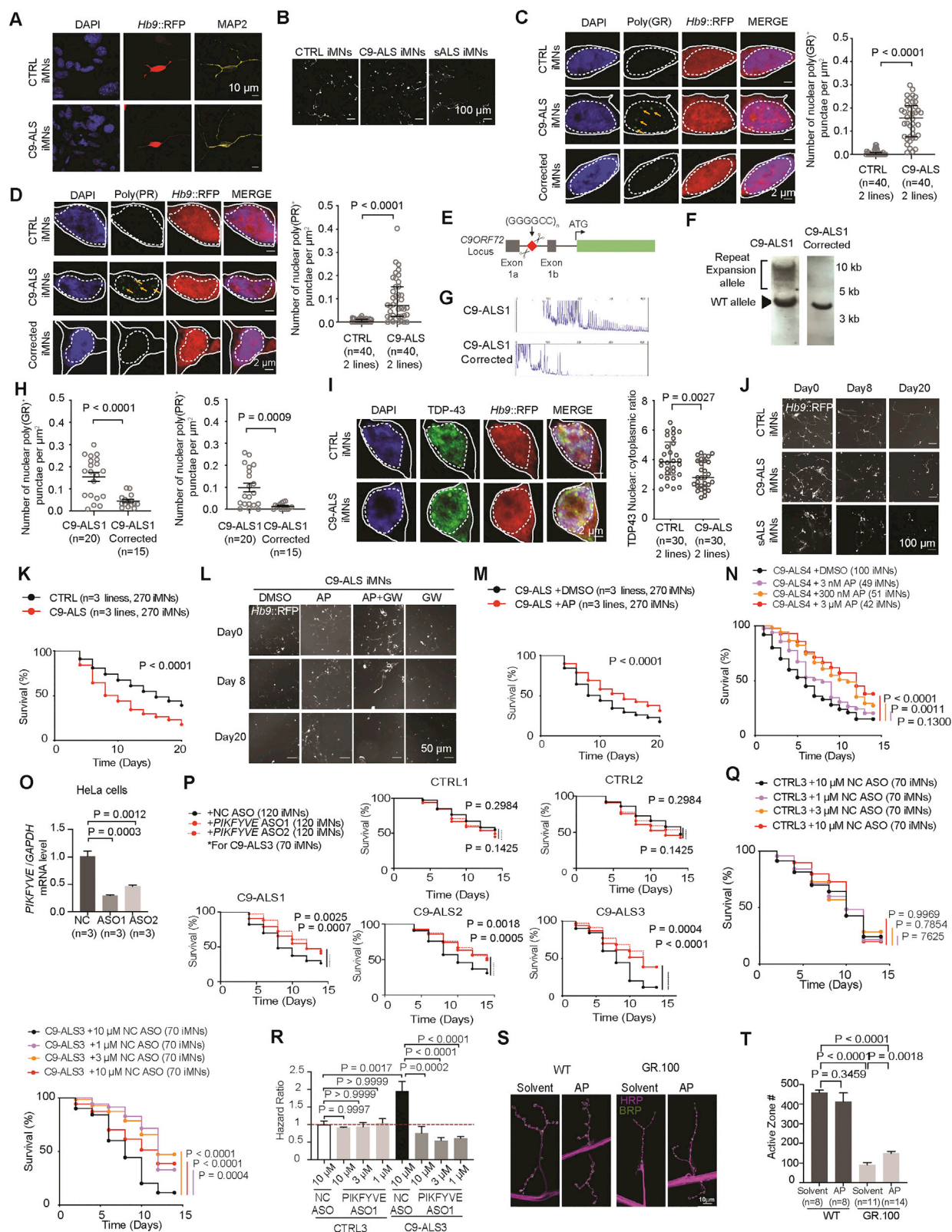
### QUANTIFICATION AND STATISTICAL ANALYSIS

Statistical analysis and data visualization were performed with the statistical software package GraphPad Prism version 9.1.0. For all experiments, the normal distribution of data sets was tested by the D'Agostino-Pearson omnibus normality test. Differences between two groups were analyzed using a two-tailed Student's *t* test, unless the data were non-normally distributed, in which case two-sided Mann-Whitney testing was used. Differences between multiple groups were analyzed using Ordinary one-way ANOVA. *P*-values were corrected for multiple comparisons using statistical hypothesis testing (Šídák method) unless the data were non-normally distributed for which nonparametric Kruskal-Wallis testing and Dunn's test (for correcting multiple comparison) were used. Mean and standard error of the mean was used for normally distributed data sets, and the median and interquartile range were used for non-normally distributed data sets. Significance was assumed at  $p < 0.05$ . Sample size estimation was performed by power analysis using preliminary data (<https://www.stat.ubc.ca/~rollin/stats/ssize/n2.html>). Any outliers that were excluded were determined by an outlier ROUT test in GraphPad Prism.

For iMN survival studies, statistical analysis was performed using two-sided log-rank test to account for the events that did not occur (meaning that iMNs that did not degenerate at the end of the survival assay). For each line, iMNs were randomly selected for tracking and used to generate the Kaplan-Meier survival curves. The number of iMNs counted per group was specified in the Kaplan-Meier plot or method section for each survival experiment. The hazard rates were calculated with the Mantel-Haenszel method and statistical analysis was performed with one-way ANOVA. The number of biological replicates (independent conversions) per group was specified in each figure. For immunocytochemistry, iMNs were quantified from two independent conversions per group. The number of iMNs counted per group was specified in each figure. For qRT-PCR, poly(GR) immunoassay and western blotting, the number of biological replicates (independent conversions) was specified in each figure or method section. For animal studies, the number of animals per group was specified in each figure.



# Supplemental figures



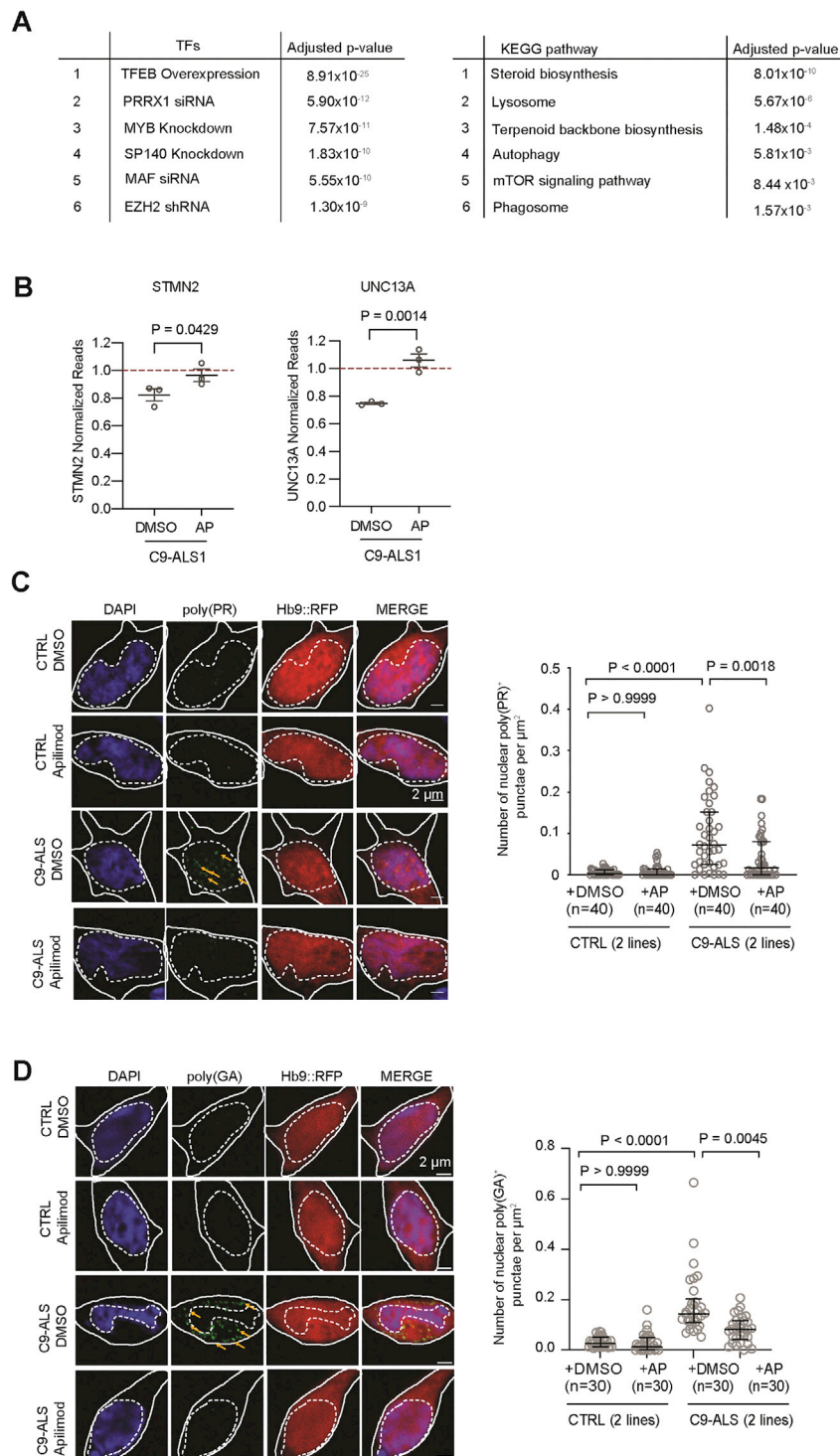
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**Figure S1. *PIKFYVE* inhibition ameliorates C9ORF72 ALS/FTD neurodegeneration, related to Figure 1**

- (A) MAP2 immunostaining of iMNs expressing *Hb9::RFP*.
- (B) Images of *Hb9::RFP*+ iMNs generated from CTRL/C9-ALS/FTD/sporadic ALS lines at the start of the survival assay.
- (C and D) Immunostaining/quantification of endogenous poly(GR)+ (C) or poly(PR)+ (D) punctae in CTRL/C9-ALS/FTD iMNs. Gray circles = the number of nuclear poly(GR)+ or poly(PR)+ punctae/ $\mu\text{m}^2$  in one iMN. Mann-Whitney test. Mean  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.
- (E) Removal of the *C9ORF72* repeat expansion by CRISPR-Cas9 editing. Guide RNAs shown as scissors.
- (F) Southern blot examining the *C9ORF72* repeat length in C9-ALS/FTD1 and its corrected isogenic line. Normal *C9ORF72* allele = 3.8-kb band.
- (G) Repeat-primed PCR analysis of the repeat expansion in C9-ALS/FTD1 and its isogenic control line.
- (H) Quantification of poly(GR)+ and poly(PR)+ punctae in C9-ALS/FTD1 and isogenic control iMNs. Gray circles = the number of nuclear poly(GR)+ or poly(PR)+ punctae/ $\mu\text{m}^2$  in one iMN. Unpaired t test. Mean  $\pm$  SEM.
- (I) Immunostaining/quantification of total TDP-43 in CTRL/C9-ALS/FTD iMNs. Gray circles = the ratio of nuclear:cytoplasmic TDP-43 from one iMN. Mann-Whitney test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.
- (J) Images of CTRL, C9-ALS/FTD, and sporadic ALS iMNs during the survival assay.
- (K) Survival of CTRL/C9-ALS/FTD iMNs (all lines/genotype in aggregate). Log rank test.
- (L) Images of C9-ALS/FTD iMNs during the survival assay when treated with DMSO, apilimod, apilimod + GW4869, or GW4869.
- (M) Survival of C9-ALS/FTD iMNs (three lines in aggregate) in DMSO/apilimod. Log rank test.
- (N) Survival curves of C9-ALS/FTD iMNs in DMSO/apilimod. Log rank test.
- (O) *PIKFYVE* mRNA levels (relative to *GAPDH*) in HeLa cells treated with indicated ASOs. One-way ANOVA. Mean of three independent ASO treatments/condition  $\pm$  SEM.
- (P) Survival of control/C9-ALS/FTD iMNs treated with indicated ASOs. Log rank test.
- (Q) Survival of CTRL3 and C9-ALS3 treated with negative control ASO/*PIKFYVE* ASO1. Log rank test.
- (R) Hazard ratio of iMNs from CTRL3 and C9-ALS3 treated with negative control/*PIKFYVE* ASO1. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (S) Images of muscle and NMJs in wild-type/*C9ORF72* GR.100 *Drosophila* larvae treated with DMSO/apilimod and immunostained with anti-BRP (active zone marker) and anti-HRP (neuronal membrane marker).
- (T) Quantification of active zones at NMJs in the indicated genotypes and conditions. Unpaired t test. Mean  $\pm$  SEM.





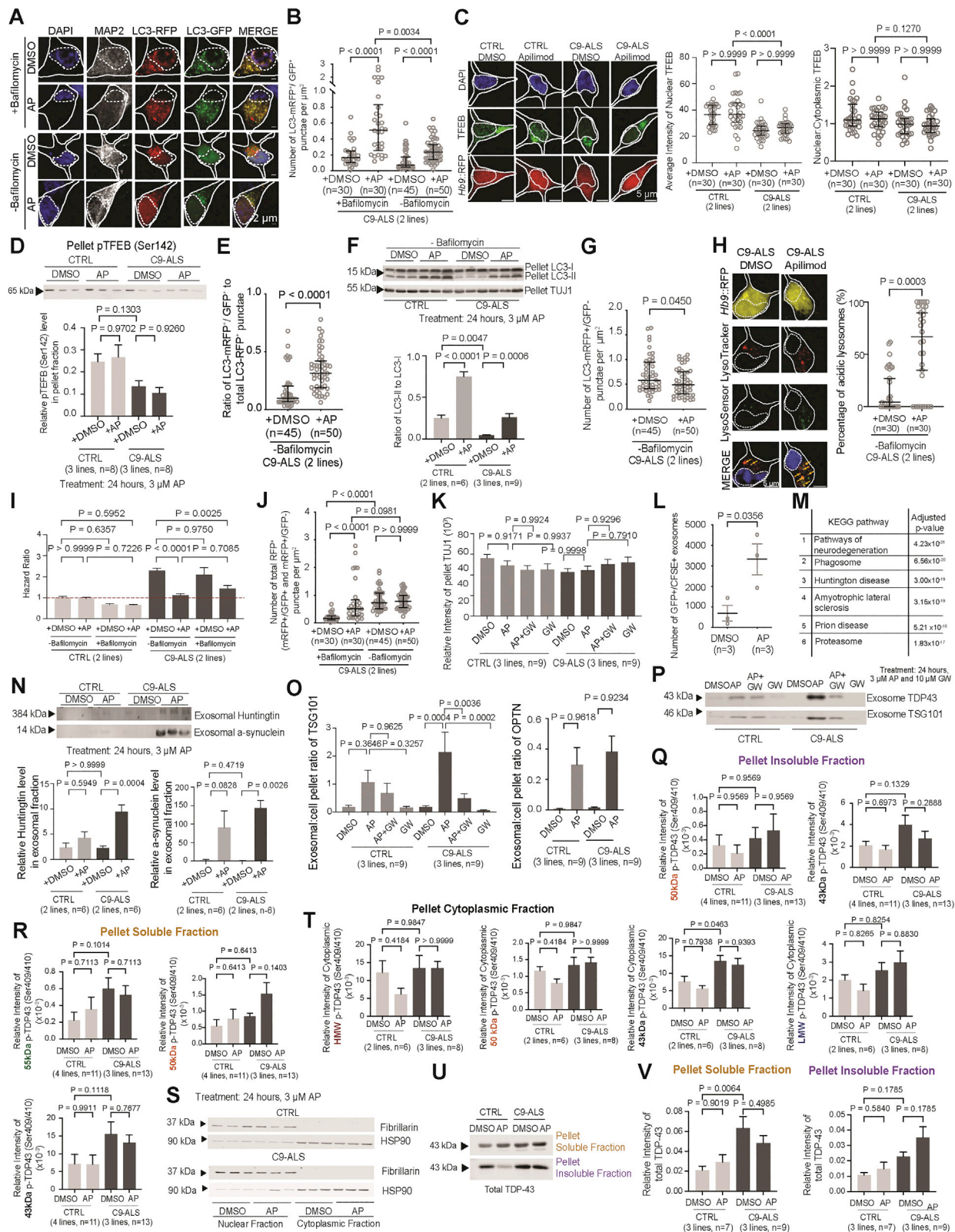
**Figure S2. *PIKFYVE* inhibition ameliorates disease pathology in *C9ORF72* ALS/FTD iMNs, related to Figure 2**

(A) Transcription factor enrichment and KEGG pathway analysis of differentially expressed genes from RNA-seq data of C9-ALS/FTD iMNs treated with DMSO/apilimod.

(B) Normalized expression (from RNA-seq data) of *STMN2* and *UNC13A* from C9-ALS/FTD iMNs treated with DMSO/apilimod. Unpaired t test. Mean  $\pm$  SEM.

(C and D) Immunostaining/quantification of poly(PR)+ (C) and poly(GA)+ (D) punctae in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of nuclear poly(PR)+ or poly(GA) punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.





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**Figure S3. *PIKFYVE* inhibition increases exocytosis of aggregation-prone proteins from iMNs, related to Figure 3**

(A and B) Images/quantification of LC3-mRFP-GFP3 fluorescence in C9-ALS/FTD iMNs in DMSO/apilimod  $\pm$  bafilomycin. Gray circles = the number of LC3-GFP+/mRFP+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.

(C) Immunostaining/quantification of TFEB in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the mean intensity of nuclear TFEB (left panel) and the average nuclear:cytoplasmic TFEB ratio (right panel) from one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.

(D) Immunoblots/quantification of phosphorylated (Ser142) TFEB in pellet fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.

(E) Quantification of the “no bafilomycin” conditions from (A). Gray circles = the ratio of LC3-mRFP+/GFP+:total LC3-mRFP+ (LC3-mRFP+/GFP+ and LC3-mRFP+/GFP–) punctae in one iMN. Mann-Whitney test. Median  $\pm$  interquartile range.

(F) Immunoblots/quantification of LC3 (LC3-I above 15 kDa and LC3-II below 15 kDa) in pellet fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.

(G) Quantification of the no bafilomycin conditions from (A). Gray circles = the number of LC3-mRFP+/GFP– punctae/ $\mu\text{m}^2$  in one iMN. Mann-Whitney test. Median  $\pm$  interquartile range.

(H) Staining/quantification of LysoSensor Green and LysoTracker in C9-ALS/FTD iMNs treated with DMSO or apilimod. Gray circles = the percentage of LysoSensor+ punctae/total LysoTracker+ punctae in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.

(I) Hazard ratio of CTRL/C9-ALS/FTD iMNs treated with DMSO, apilimod, DMSO + bafilomycin, or apilimod + bafilomycin. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.

(J) Quantification of (A), the DMSO/apilimod conditions  $\pm$  bafilomycin. Gray circles = the number of total LC3-mRFP+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range.

(K) Quantification of TUJ1 in pellet fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO, apilimod, apilimod + GW4869, or GW4869. One-way ANOVA. Mean  $\pm$  SEM.

(L) FACS analysis of GFP+/CFSE+ exosomes from CTRL iMNs treated with DMSO/apilimod. Data points = an independently differentiated culture. Unpaired t test. Mean  $\pm$  SEM.

(M) KEGG pathways of differentially expressed proteins in the exosomal fraction of C9-ALS/FTD iMNs treated with DMSO/apilimod.

(N) Immunoblots/quantification of Huntingtin and alpha-synuclein in exosomal fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.

(O) Ratios of TSG101 and OPTN in the exosome:cell pellet fractions of CTRL/C9-ALS/FTD iMNs. One-way ANOVA. Mean  $\pm$  SEM.

(P) Immunoblots of total TDP43 (43 kDa) and TSG101 in exosomal fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO, apilimod, apilimod + GW4869, or GW4869.

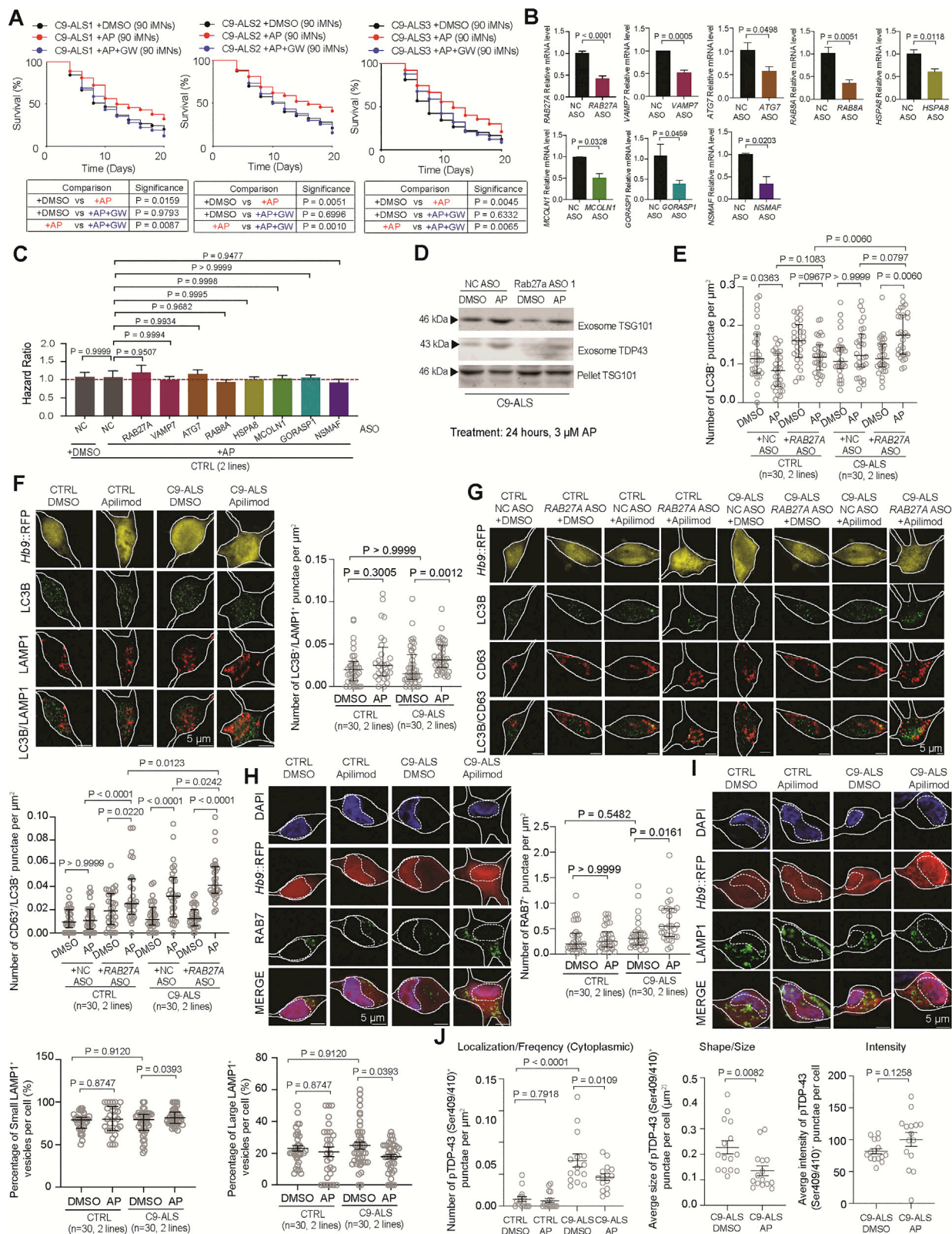
(Q and R) Quantification of 50 and 43 kDa pTDP-43 (Ser409/410) in the pellet insoluble (Q) or soluble (R) fraction from Figure 3M. One-way ANOVA. Mean  $\pm$  SEM.

(S) Immunoblots of fibrillarin (37 kDa) and HSP90 (90 kDa) in nuclear or cytoplasmic fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod.

(T) Quantification of high-molecular weight (HMW, >75 kDa), 50 kDa, 43 kDa, and low-molecular weight (LMW, <43 kDa) pTDP-43 in the cytoplasmic soluble fraction from Figure 3P. One-way ANOVA. Mean  $\pm$  SEM.

(U and V) Immunoblots/quantification of total TDP-43 in soluble/insoluble fractions from CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. One-way ANOVA. Mean  $\pm$  SEM.



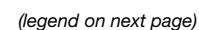




**Figure S4. *PIKFYVE* inhibition clears pTDP-43 through amphisome and multivesicular body exocytosis, related to Figure 4**

- (A) Survival C9-ALS/FTD iMNs treated with DMSO, apilimod, or apilimod + GW4869. Log rank test. p values were not corrected for multiple comparisons.
- (B) mRNA levels of *RAB27A* (relative to *HPRT*) or other indicated genes (relative to *GAPDH*) in iMNs treated with negative control ASO or an ASO targeting each gene. Unpaired t test. Mean  $\pm$  SEM.
- (C) Hazard ratio of CTRL iMNs treated with the ASOs indicated plus DMSO/apilimod. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (D) Immunoblots of TSG101 and pTDP-43 (Ser409/410) (50 kDa) in exosomal fractions from C9-ALS/FTD iMNs treated with DMSO/apilimod plus negative control/*RAB27A* ASO1.
- (E) Quantification of endogenous LC3B+ punctae in CTRL or C9-ALS/FTD iMNs treated with negative control/*RAB27A* ASO and DMSO/apilimod. Gray circles = the number of LC3B+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range.
- (F) Immunostaining/quantification of endogenous LC3B+/LAMP1+ punctae in CTRL or C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of LC3+/LAMP1+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid lines outline the cell body.
- (G) Immunostaining/quantification of endogenous LC3B+/CD63+ punctae in CTRL/C9-ALS/FTD iMNs treated with negative control/*RAB27A* ASO and DMSO/apilimod. Gray circles = the number of LC3B+/CD63+ punctae/ $\mu\text{m}^2$  from one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid lines outline the cell body.
- (H) Immunostaining/quantification of endogenous RAB7+ punctae in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of RAB7+ punctae/ $\mu\text{m}^2$  from one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.
- (I) Immunostaining/quantification of endogenous LAMP1+ punctae in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the percentage of small LAMP1+ vesicles/cell (left) and large LAMP1+ vesicles/cell (right) from one iMN. One-way ANOVA. Mean  $\pm$  SEM. Solid and dotted lines outline the cell body and nucleus, respectively.
- (J) Quantification of pTDP-43+ (Ser409/410) punctae in CTRL/C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of cytoplasmic pTDP43+ punctae/ $\mu\text{m}^2$  (left), the average size (in  $\mu\text{m}^2$ ) of pTDP43+ punctae/neuron (middle), and the average intensity of pTDP43+ punctae/neuron (right) in one iMN. One-way ANOVA (left); unpaired t test (middle and right). Mean  $\pm$  SEM.



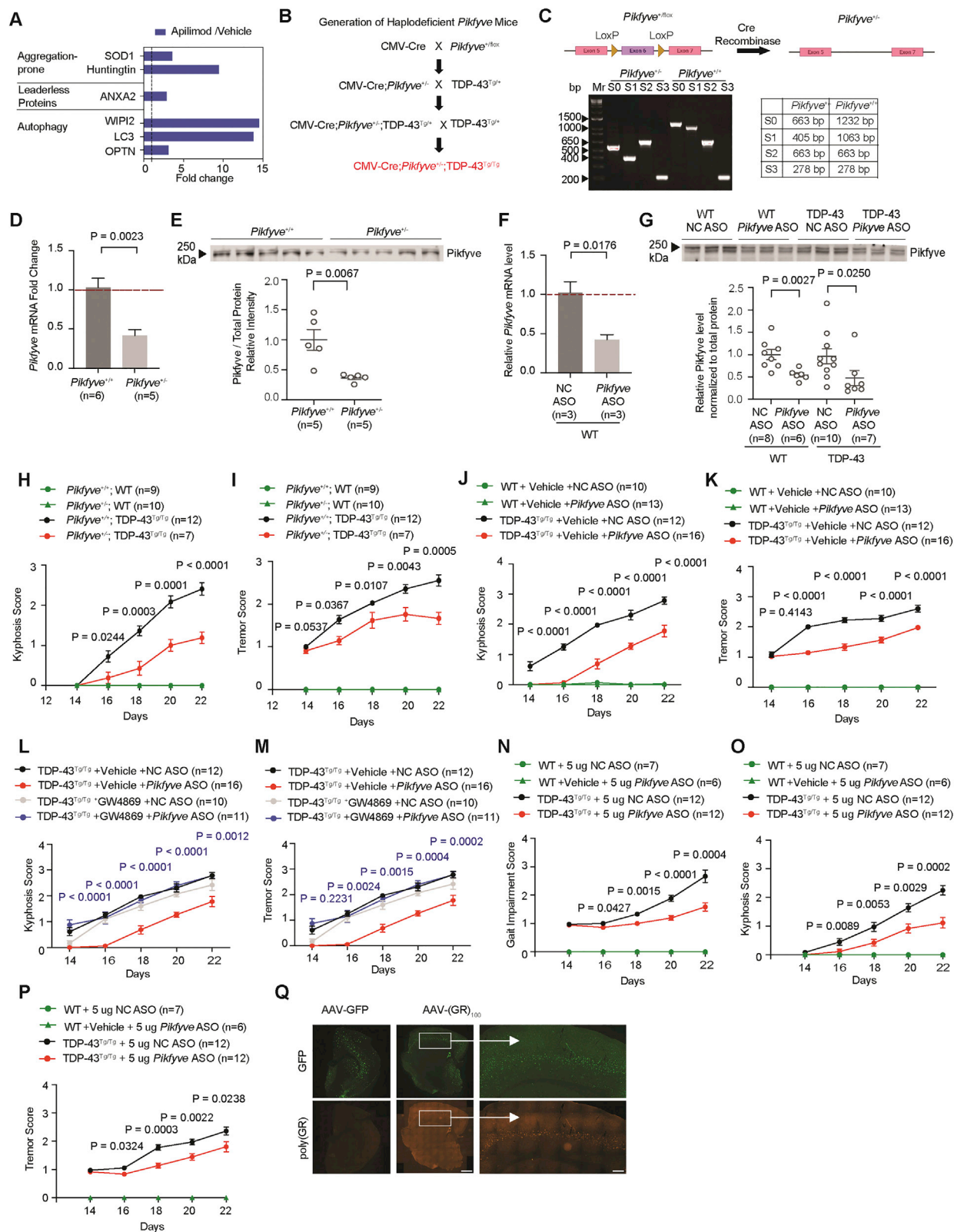




**Figure S5. PIKFYVE inhibition improves iMN proteostasis and survival for diverse forms of ALS, related to Figure 5**

- (A) Images of sporadic ALS iMNs during the survival assay when treated with DMSO, apilimod, apilimod + GW4869, or GW4869.
- (B) Survival of CTRL iMNs treated with DMSO and eight sporadic ALS (sALS) iMNs treated with DMSO/apilimod. Log rank test.
- (C) Survival of iMNs from sALS9 in DMSO/apilimod. Log rank test.
- (D) Hazard ratios of iMNs from sALS9 treated with DMSO/apilimod. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (E) Hazard ratios of iMNs from three CTRL and six sALS lines in aggregate (sALS1–sALS6) treated with negative control ASO, *PIKFYVE* ASO1, or ASO2. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (F) Survival of iMNs from two *FUS* ALS (H517Q, R522R mutation) and *TARDBP* ALS (G298S mutation) lines treated with DMSO/apilimod. Log rank test.
- (G) Immunostaining/quantification of FUS in iMNs from two CTRL and two *FUS* ALS (H517Q, R522R mutation) lines treated with DMSO/apilimod. Gray circles = the nuclear:cytoplasmic FUS ratio from one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range. Solid and dotted lines outline the cell body and nucleus, respectively.
- (H) Hazard ratios of iMNs from CTRL, two sporadic ALS, and one *FUS*-ALS (R522R mutation) lines treated with negative control, *VAMP7*, or *ATG7* ASO plus DMSO/apilimod. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (I) Hazard ratios of iMNs from CTRL and three sporadic ALS lines treated with DMSO, apilimod or apilimod + GW4869. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (J) Survival of iMNs from three sporadic ALS lines treated with DMSO, apilimod, or apilimod + GW4869. Log rank test.
- (K and L) Hazard ratios of iMNs from one CTRL and one *TARDBP* ALS (G298S mutation) (K) or one *FUS* ALS (H517Q) line (L) treated with DMSO, apilimod, or apilimod + GW4869. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (M) Survival of iMNs from a *FUS* ALS (H517Q) and *TARDBP* ALS line treated with DMSO, apilimod, or apilimod + GW4869. Log rank test.
- (N) Hazard ratios of iMNs from two CTRL lines treated with DMSO and one *FUS* ALS (R522R mutation) line treated with DMSO, apilimod, DMSO + bafilomycin, or apilimod + bafilomycin. Mean of independent iMN conversions  $\pm$  SEM. One-way ANOVA.
- (O) Images of worms: control, untreated (top), or apilimod-treated (bottom) TDP-43<sup>A315T</sup> transgenics expressing unc-47::GFP in motor neurons. Arrows indicate gaps or breaks along neuronal processes that are scored as neurodegeneration events.
- (P) Apilimod delayed paralysis in TDP-43<sup>A315T</sup> transgenics. Log rank test.  $n > 200$  worms/experiment.
- (Q) Apilimod suppressed motor neuron degeneration in TDP-43<sup>A315T</sup> transgenics. One-way ANOVA.





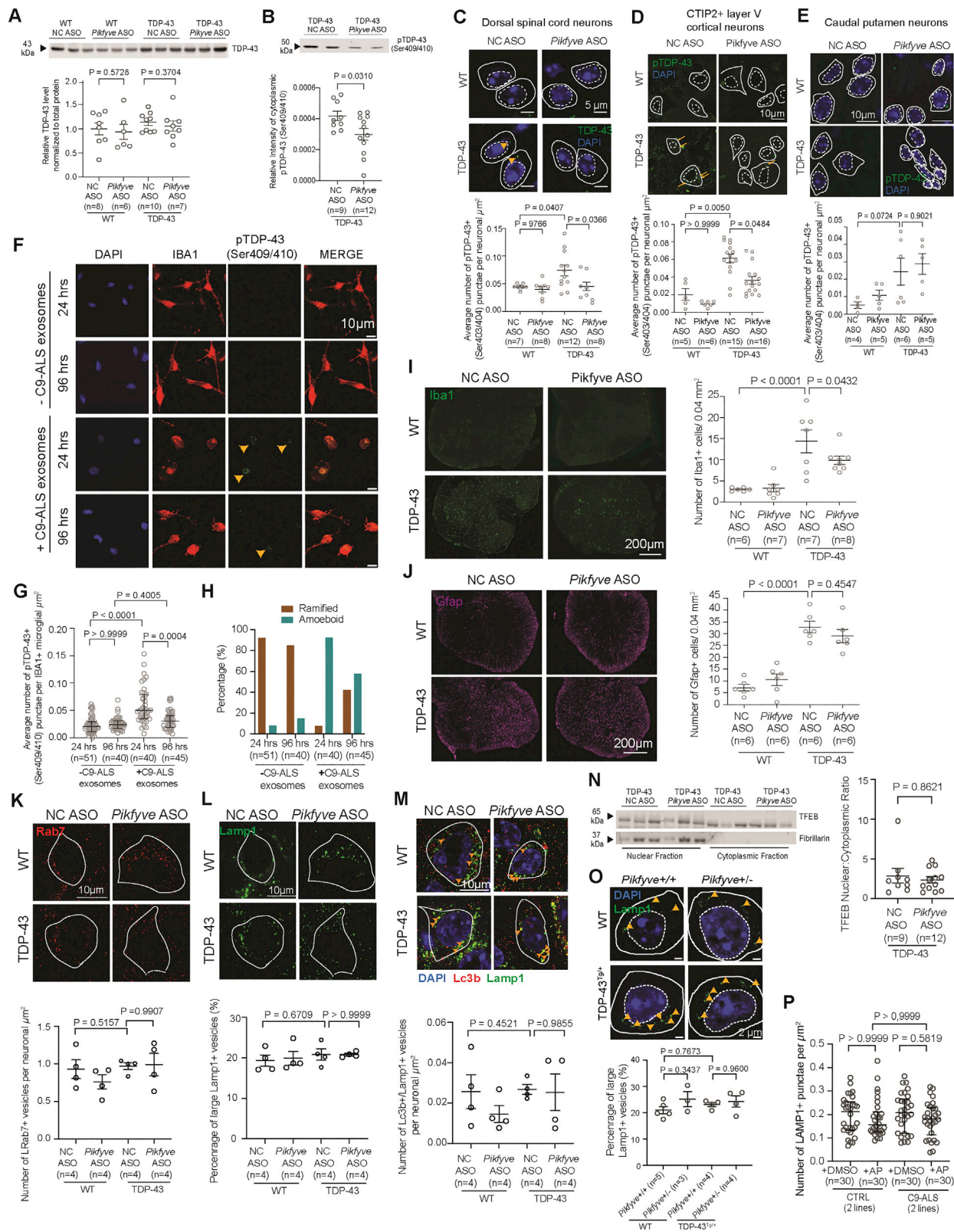
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**Figure S6. *Pikfyve* suppression improves motor function and extends survival of TDP-43 and *C9ORF72* mice, related to Figure 6**

- (A) Fold-change of selected protein levels in adult mouse CSF after intrathecal injection of apilimod compared to vehicle.  $n = 6$  and 4 mice for the vehicle- and apilimod-treated group, respectively. Values = the average of two technical replicates/condition/group.
- (B) Mating scheme for generation of a single-allele deletion of *Pikfyve* in TDP-43<sup>Tg/Tg</sup> mice.
- (C) Targeting strategy to disrupt exon 6 of the *Pikfyve* gene using the Cre-lox system.
- (D) *Pikfyve* mRNA levels in *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/Tg</sup> mouse brain (normalized to *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/Tg</sup> mice) at 15–18 days of age. Unpaired t test. Mean  $\pm$  SEM. Data points = one mouse.
- (E) Immunoblot of PIKFYVE in whole brains of *Pikfyve*<sup>+/-</sup>;WT and *Pikfyve*<sup>+/-</sup>;WT mice at P15–P18. Unpaired t test. Mean  $\pm$  SEM. Data points = one mouse.
- (F) *Pikfyve* mRNA levels (relative to *Ppia*) in brains of P7 WT mice treated with negative control/*Pikfyve* ASO at P0.  $n = 3$  mice/group. Unpaired t test. Mean  $\pm$  SEM.
- (G) Immunoblot of PIKFYVE levels in whole brains of WT and TDP-43<sup>Tg/Tg</sup> mice at P20 after treatment with negative control/*Pikfyve* ASO at P1. Mann-Whitney test. Median  $\pm$  interquartile range. Data points = one mouse.
- (H and I) The kyphosis (H) and tremor scores (I) of *Pikfyve*<sup>+/-</sup>;WT, *Pikfyve*<sup>+/-</sup>;WT, *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/Tg</sup>, and *Pikfyve*<sup>+/-</sup>;TDP-43<sup>Tg/Tg</sup> mice. Unpaired t test. Mean  $\pm$  SEM.
- (J and K) The kyphosis (J) and tremor scores (K) of WT or TDP-43<sup>Tg/Tg</sup> mice treated with vehicle and negative control/*Pikfyve* ASO. Unpaired t test. Mean  $\pm$  SEM.
- (L and M) The kyphosis (L) and tremor scores (M) of TDP-43<sup>Tg/Tg</sup> mice treated with vehicle + NC ASO, vehicle + *Pikfyve* ASO, GW4869 + negative control ASO, and GW4869 + *Pikfyve* ASO. Unpaired t test. Mean  $\pm$  SEM.
- (N–P) The gait impairment (N), kyphosis (O), and tremor (P) scores of WT or TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Unpaired t test. Mean  $\pm$  SEM.
- (Q) Immunostaining of GFP and poly(GR) in the cortex of mice transduced with AAV-eGFP or AAV-eGFP-(GR)<sub>100</sub>. Rectangle = magnification of the area.





(legend on next page)



**Figure S7. *Pikfyve* suppression reduces TDP-43 and C9ORF72 pathology and neurodegeneration *in vivo*, related to Figure 7**

(A) Immunoblot of total TDP-43 levels in whole brains of WT/TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. One-way ANOVA. Mean  $\pm$  SEM. Data points = one mouse.

(B) Immunoblot of pTDP-43 (Ser409/410) levels in the cytoplasmic fraction of whole brain lysates from TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Unpaired t test. Mean  $\pm$  SEM. Data points = one mouse.

(C–E) Immunostaining/quantification of pTDP-43+ (Ser403/404) punctae in large TUJ1+ neurons in the dorsal spinal cord (C), in layer V neurons in the motor cortex layer (D), and in neurons in the caudal putamen (E) from WT/TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Solid and dotted lines outline the cell body and nucleus, respectively. Arrows mark pTDP-43+ punctae. Data points = the average number of pTDP-43+ punctae in neurons/ $\mu\text{m}^2$  for one mouse. (C) One-way ANOVA. Mean  $\pm$  SEM. (D and E) Kruskal-Wallis test. Mean  $\pm$  SEM.

(F) Immunostaining of pTDP-43+ (Ser409/410) punctae in CTRL iPSC-derived IBA1+ microglia treated with/without exosomes harvested from C9-ALS/FTD iMNs for 24 or 96 h. Arrows mark pTDP-43+ punctae in IBA1+ microglia.

(G and H) Quantification of (F). For (G), gray circles = the number of pTDP-43+ punctae in IBA1+ microglia/ $\mu\text{m}^2$  in one IBA1+ microglial cell treated with/without exosomes harvested from C9-ALS/FTD iMNs. Kruskal-Wallis test. Median  $\pm$  interquartile range. For (H), values = the percentage of ramified or amoeboid IBA1+ microglia.

(I) Immunostaining/quantification of IBA1+ microglia in the spinal cord ventral horn of WT or TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the number of IBA1+ microglia/ventral horn section in one mouse. One-way ANOVA. Mean  $\pm$  SEM.

(J) Immunostaining/quantification of GFAP+ astrocytes in the spinal cord ventral horn of WT or TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the number of GFAP+ astrocytes/ventral horn section in one mouse. One-way ANOVA. Mean  $\pm$  SEM.

(K–M) Immunostaining/quantification of RAB7+ punctae (K), percentage of large LAMP1+ vesicles (L), or the number of LC3B+/LAMP1+ punctae (M) in large TUJ1+ neurons in the spinal cord ventral horn from WT/TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the average number of RAB7+ punctae/ $\mu\text{m}^2$  (K), the percentage of large LAMP1+ vesicles (L), or the number of LC3B+/LAMP1+ punctae (M) in TUJ1+ neurons for one mouse. One-way ANOVA. Mean  $\pm$  SEM.

(N) Immunoblot of TFEB levels in the nuclear or cytoplasmic fraction of whole brain lysates from TDP-43<sup>Tg/Tg</sup> mice treated with negative control/*Pikfyve* ASO. Data points = the nuclear:cytoplasmic TFEB ratio in one mouse. Mann-Whitney test. Mean  $\pm$  SEM.

(O) Immunostaining/quantification of the percentage of large LAMP1+ vesicles in large TUJ1+ neurons in the spinal cord ventral horn from 10-month-old *Pikfyve*<sup>+/+</sup>;WT, *Pikfyve*<sup>+/+</sup>;WT, *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/+</sup>, and *Pikfyve*<sup>+/+</sup>;TDP-43<sup>Tg/+</sup> mice. Solid and dotted lines outline the cell body and nucleus, respectively. Arrows mark LAMP1+ punctae. Data points = the percentage of large LAMP1+ vesicles in TUJ1+ neurons for one mouse. One-way ANOVA. Mean  $\pm$  SEM.

(P) Quantification of LAMP1+ punctae in CTRL and C9-ALS/FTD iMNs treated with DMSO/apilimod. Gray circles = the number of LAMP1+ punctae/ $\mu\text{m}^2$  in one iMN. Kruskal-Wallis test. Median  $\pm$  interquartile range.