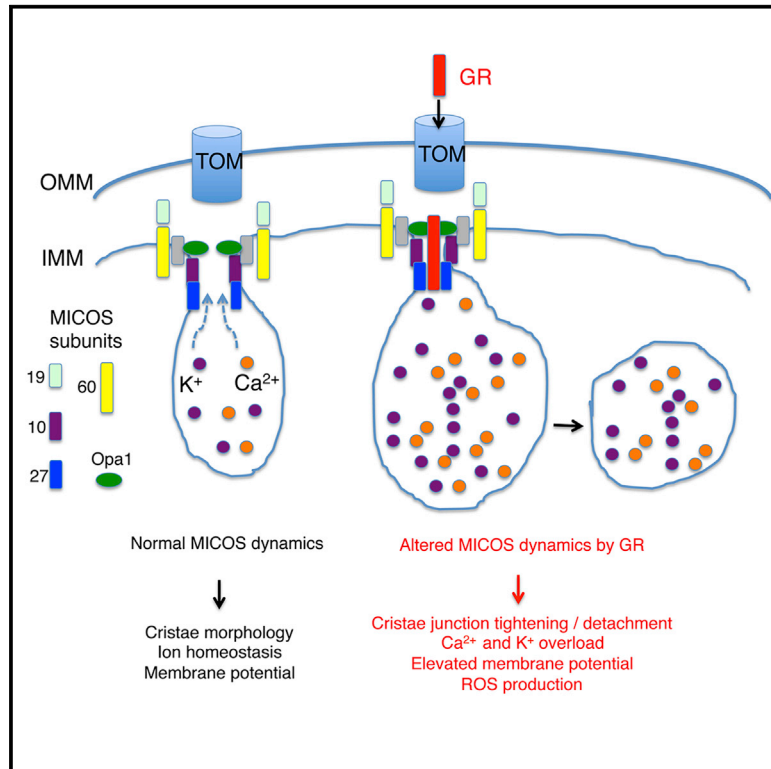


## Altered MICOS Morphology and Mitochondrial Ion Homeostasis Contribute to Poly(GR) Toxicity Associated with C9-ALS/FTD

### Graphical Abstract



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### In Brief

G4C2 repeat expansion in *C9ORF72* is the most common genetic cause of ALS. Repeat-encoded poly(GR) is believed to contribute to disease, albeit with not fully defined mechanisms. Li et al. show that poly(GR) targets the MICOS complex, causing mitochondrial toxicity. Manipulating MICOS is beneficial in animal models and patient cells.

### Highlights

- C9-ALS/FTD-associated poly(GR) causes muscle toxicity in a *Drosophila* model
- Poly(GR) enters mitochondria and binds to MICOS proteins, altering MICOS dynamics
- Poly(GR) alters mitochondrial cristae morphology, ion homeostasis, and metabolism
- Manipulation of MICOS or restoration of ion homeostasis rescues disease phenotypes



## Report

# Altered MICOS Morphology and Mitochondrial Ion Homeostasis Contribute to Poly(GR) Toxicity Associated with C9-ALS/FTD

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## SUMMARY

Amyotrophic lateral sclerosis (ALS) manifests pathological changes in motor neurons and various other cell types. Compared to motor neurons, the contribution of the other cell types to the ALS phenotypes is understudied. G4C2 repeat expansion in *C9ORF72* is the most common genetic cause of ALS along with frontotemporal dementia (C9-ALS/FTD), with increasing evidence supporting repeat-encoded poly(GR) in disease pathogenesis. Here, we show in *Drosophila* muscle that poly(GR) enters mitochondria and interacts with components of the Mitochondrial Contact Site and Cristae Organizing System (MICOS), altering MICOS dynamics and intra-subunit interactions. This impairs mitochondrial inner membrane structure, ion homeostasis, mitochondrial metabolism, and muscle integrity. Similar mitochondrial defects are observed in patient fibroblasts. Genetic manipulation of MICOS components or pharmacological restoration of ion homeostasis with nigericin effectively rescue the mitochondrial pathology and disease phenotypes in both systems. These results implicate MICOS-regulated ion homeostasis in C9-ALS pathogenesis and suggest potential new therapeutic strategies.

## INTRODUCTION

Amyotrophic lateral sclerosis (ALS) is characterized primarily by the progressive degeneration of upper motor neurons (UMNs) in the motor cortex and lower motor neurons (LMNs) in the brainstem and spinal cord (Brown and Al-Chalabi, 2017), with possible LMN to UMN spread (Chou and Norris, 1993). The earliest pathophysiological events initiating the disease remain to be elucidated, although accumulating evidence suggests that pathological changes can occur distally at the neuromuscular junction (NMJ) in advance of MN loss and onset of clinical symptoms (Moloney et al., 2014). Traditionally considered a MN disease, ALS is emerging as a “multisystem” disease with pathological changes in various cell types, including peripheral immune cells, neurons, microglia, astrocytes, and muscle cells (Pansarasa et al., 2014; Loeffler et al., 2016). Skeletal muscle defects are common and occur early in ALS (Pansarasa et al., 2014; Loeffler et al., 2016). In the copper-zinc superoxide dismutase (SOD1)-G93A mouse model of ALS, for example, muscle dysfunction represents one of the earliest pathological events, preceding MN death (Dobrowolny et al., 2008), and the mus-

cle-specific expression of SOD1-G93A can lead to MN death (Wong and Martin, 2010). One possible mechanism of muscle contribution to MN health in ALS is NMJ stability; defects in the NMJ can cause MN denervation and death in a dying-back manner (Moloney et al., 2014). In C9-ALS patients (Jokela et al., 2016), BAC transgenic mice (Liu et al., 2016), and patient induced pluripotent stem cell (iPSC)-derived muscle fibers (Lynch et al., 2019), muscle cell pathology is observed, underlining the relevance of muscle to the pathogenesis of this most common form of ALS. However, the nature of muscular defects in ALS, its pathogenic mechanism, and its contribution to the broad spectrum of disease phenotypes are not well understood.

Expansion of G4C2 repeats in *C9ORF72* accounts for ~40% of familial and 5%–10% of sporadic ALS cases, with repeat numbers ranging from a few dozen to thousands (DeJesus-Hernandez et al., 2011; Renton et al., 2011). A number of mechanisms of disease pathogenesis by G4C2 repeat expansion have been proposed, including haplo-insufficiency of *C9ORF72*, toxicity associated with RNA foci formed by sense and anti-sense RNAs, or proteotoxicity induced by dipeptide repeat (DPR) proteins translated from G4C2 repeat-carrying



transcripts (Taylor et al., 2016; Gendron and Petrucelli, 2018). Increasing evidence emphasizes the contribution of DPR toxicity in C9-ALS/frontotemporal dementia (FTD), especially arginine-containing DPR proteins (GR and PR), with diverse processes ranging from nucleolar function and nucleocytoplasmic transport to RNA granule dynamics, mRNA splicing and translation, and the DNA repair pathway having been implicated (Taylor et al., 2016; Gendron and Petrucelli, 2018; Yuva-Aydemir et al., 2018). The relative contribution of the various mechanisms to the onset and progression of this multisystem disease, however, remains uncertain. *Drosophila* has been an excellent model organism for investigating pathogenic mechanisms of human neurological disorders (Jaiswal et al., 2012), and a large body of work has been done on C9-ALS/FTD-related models in *Drosophila* (Xu et al., 2013; Mizielinska et al., 2014; Freibaum et al., 2015; Yang et al., 2015; Zhang et al., 2015; Lee et al., 2016; Simone et al., 2018; Xu and Xu, 2018; Berson et al., 2019; He et al., 2019; Lopez-Gonzalez et al., 2019; Moens et al., 2019). Almost all of these studies are focused on neuronal settings. Thus, whether disease gene products associated with C9-ALS/FTD affect muscle cells to cause the ALS aspects of the C9-ALS/FTD spectrum of phenotypes, and if so, the cellular mechanisms involved, remains unclear.

## RESULTS

### Muscle Mitochondrial Defects Caused by Poly(GR) Expression in Flies

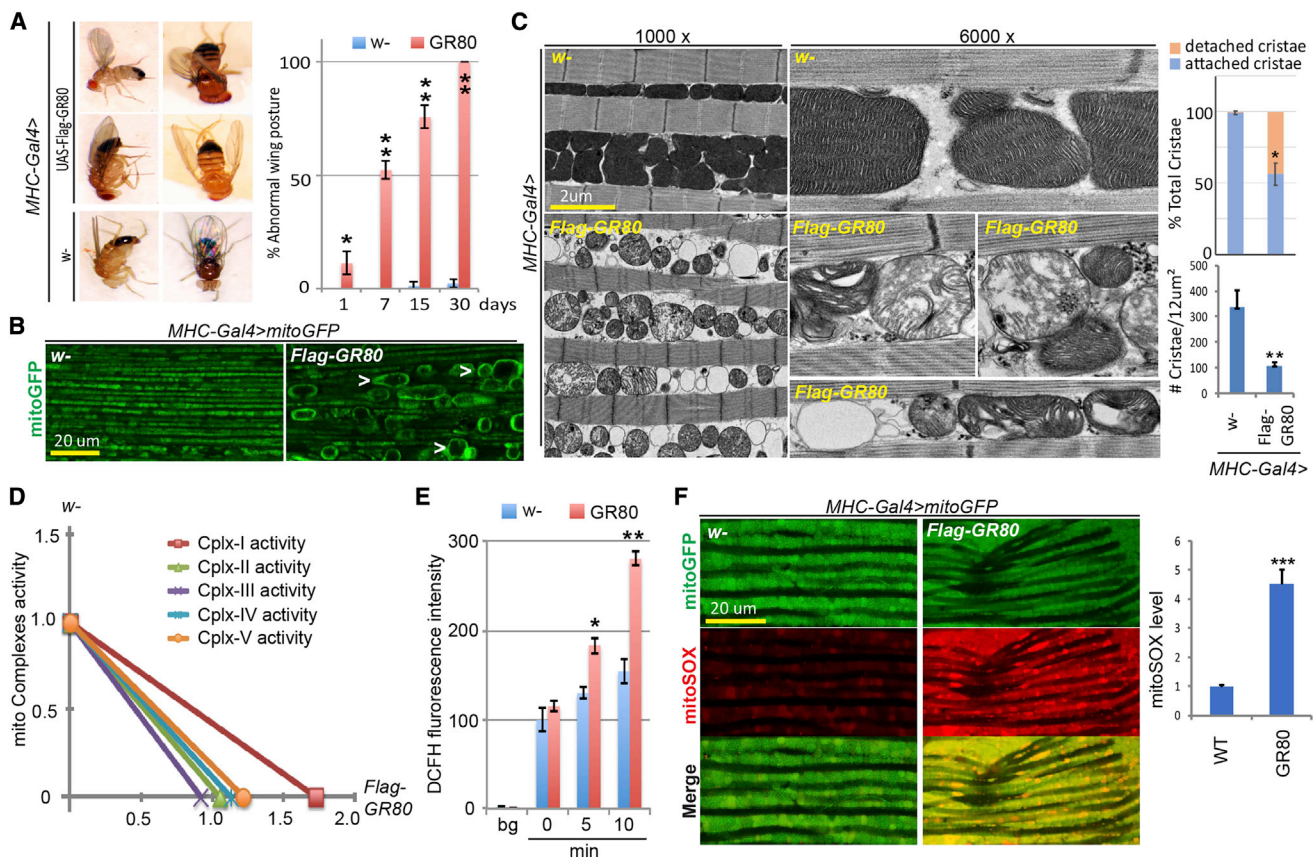
We used *Drosophila* as a model to assess the effect of GR in muscle. Available poly(GR) transgenic lines (Mizielinska et al., 2014; Yang et al., 2015) were specifically expressed in fly muscle using the *UAS-Gal4* system. To discern the effect of repeat length on toxicity, transgenes expressing GR36, GR80, and GR100 were used. The GR80 protein has a FLAG tag at the N terminus to aid protein detection (Yang et al., 2015). A *G4C2-36* transgene potentially expressing all types of DPRs and a *36RO* transgene expressing *G4C2-36* repeat RNA only (Mizielinska et al., 2014) was also included. The development, morphology, behavior, or survival appeared normal in newly enclosed flies of all genotypes. Around 1 week after eclosion, however, ~50% of the *GR100* and *GR80* transgenic flies started to exhibit held-up or droopy wing postures. After 2–3 weeks, virtually 100% of the flies showed abnormal wing posture and were flightless (Figures 1A, S1A, and S1B). *GR36* and *G4C2-36* transgenic flies exhibited a later onset and less penetrant wing posture phenotype than *GR100* and *GR80* flies, whereas *36RO* flies were phenotypically normal (Figures S1A and S1B). Quantification of GR expression in the various lines (Figure S1C) indicated that *GR100* was expressed at a slightly lower level than *GR80*. However, *GR36* expression level was higher than *GR100*, although its phenotype was weaker. Thus, GR repeat length correlates with toxicity. In subsequent studies, we focused on analyzing *GR80* flies, because this is the only GR line with a FLAG tag at the N terminus that facilitates later biochemical and cell biological studies of GR protein, and we have length-matched and FLAG-tagged *GA80* and *PR80* as controls, which did not show obvious wing phenotype (Figure S1A).

The wing posture phenotype in the various fly lines correlated with indirect flight muscle (IFM) integrity (Figure S1D), which is often associated with mitochondrial health. We examined mitochondrial morphology in *GR80* fly IFM using a matrix-targeted mito-GFP reporter. Compared to the controls, *GR80* fly muscle contained swollen mitochondria when analyzed at 2–3 weeks of age (Figure 1B). Many mitochondria were round and had areas of matrix devoid of mito-GFP. Transmission electron microscopy (TEM) revealed swollen mitochondria, as well as small and round mitochondria devoid of electron-dense matrix material in *GR80* flies (Figure 1C). Closer examination further revealed aberrant cristae structure. Compared to control muscle mitochondria abundant in lamellar cristae, each connected to the inner mitochondrial membrane (IMM) via cristae junctions (CJs), many mitochondria in *GR80* flies were devoid of CJs, forming concentric cristae stacks inside the matrix or short tubular cristae detached from IMM, indicating that *GR80* damages CJ structures. Similar mitochondrial defects were seen in *GR100* flies (Figures S1E and S1F). No obvious mitochondria defect was observed in length-matched *GA100*, *PA100*, or *PR100* fly muscle (Figures S1E and S1F) or *GA80* and *PR80* fly muscle (Figure S1G), despite similar levels of expression of FLAG-tagged *GA80*, *PR80*, and *GR80* (Figure S1H).

We next analyzed mitochondrial function in *GR80* fly muscle. There was increased ATP production in *GR80* and *GR100* flies (Figures S1I and S1J). This suggests that ATP production by oxidative phosphorylation (OxPhos), a key function of mitochondria, was not compromised in *GR80* flies, or that it was compensated for by other means. We also found that complex I activity was elevated, whereas complex II–V were largely unaltered in *GR80* flies (Figure 1D). Elevated complex I activity may account for the increased ATP level. As complex I is the major site of reactive oxygen species (ROS) production, we tested whether the redox state is altered in *GR80* flies. Using the ROS dyes DCFH (Figure 1E) and MitoSOX (Figure 1F), we detected elevated ROS in *GR80* fly IFM. *GR80* thus causes structural and functional changes in muscle mitochondria.

### Mitochondrial Localization of Poly(GR)

Our observation of mitochondrial changes in *GR80* flies raised the possibility that *GR80* may associate with mitochondria. Using confocal microscopy, we could detect the cytosolic localization of *GR80*, with some colocalizing with mitochondria in fly muscle (Figure 2A) and mammalian cells (Figure 2B). Using immunoelectron microscopy (immuno-EM), we confirmed *GR80* mitochondrial localization (Figure 2C). To further confirm this, and to determine the specific compartment of *GR80* localization, we purified mitochondria from *GR80* fly muscle and prepared mitochondrial subfractions. Western blot analysis indicated the robust expression of *GR80* in fly muscle (Figure S2A). The majority of mitochondrial *GR80* was localized to the IMM in a pattern similar to the intermembrane space (IMS)/IMM marker *Opa1* (Figure 2D). *GR80* was sensitive to proteinase K treatment of mitoplasts but not intact mitochondria preparations, suggesting that it is facing the IMS (Figure S2B). The enriched mitochondrial localization was rather specific for *GR80*, as neither *GA80* nor *PR80* exhibited obvious mitochondrial localization (Figures S2C and S2D).



**Figure 1. Mitochondrial Defects in GR80 Transgenic Fly Muscle**

(A) Images showing abnormal wing posture phenotypes in *Mhc-Gal4>GR80* flies. The bar graph shows age-dependent progression of the wing posture phenotype (n = 90 flies, 3 biological repeats). The error bars signify standard deviation (SD) in this and all of the other figures.

(B) Monitoring of mitochondrial morphology in thoracic flight muscle of *Mhc-Gal4>GR80* flies expressing a mito-GFP reporter. The images are representative of 3 independent samples.

(C) TEM images of IFM mitochondria of control (*Mhc-Gal4/w-*) and *Mhc-Gal4>GR80* flies at lower (1,000 $\times$ ) and higher (6,000 $\times$ ) magnifications. The graphs show quantification (n = 5 sections from 3 independent samples).

(D) Mitochondrial respiratory chain activities in *GR80* compared to control (*Mhc-Gal4/w-*) flies. The data represent 1-time experiment.

(E) Quantification of ROS levels measured with DCFH fluorescence in *GR80* fly muscle (n = 3).

(F) Mito-SOX staining of mitochondrial ROS levels in control and *GR80* flies expressing mito-GFP. The graph shows quantification (n = 3).

\*p < 0.05 and \*\*p < 0.01, 1-way ANOVA test followed by Student-Newman-Keuls (SNK) test plus Bonferroni correction. See also Figure S1.

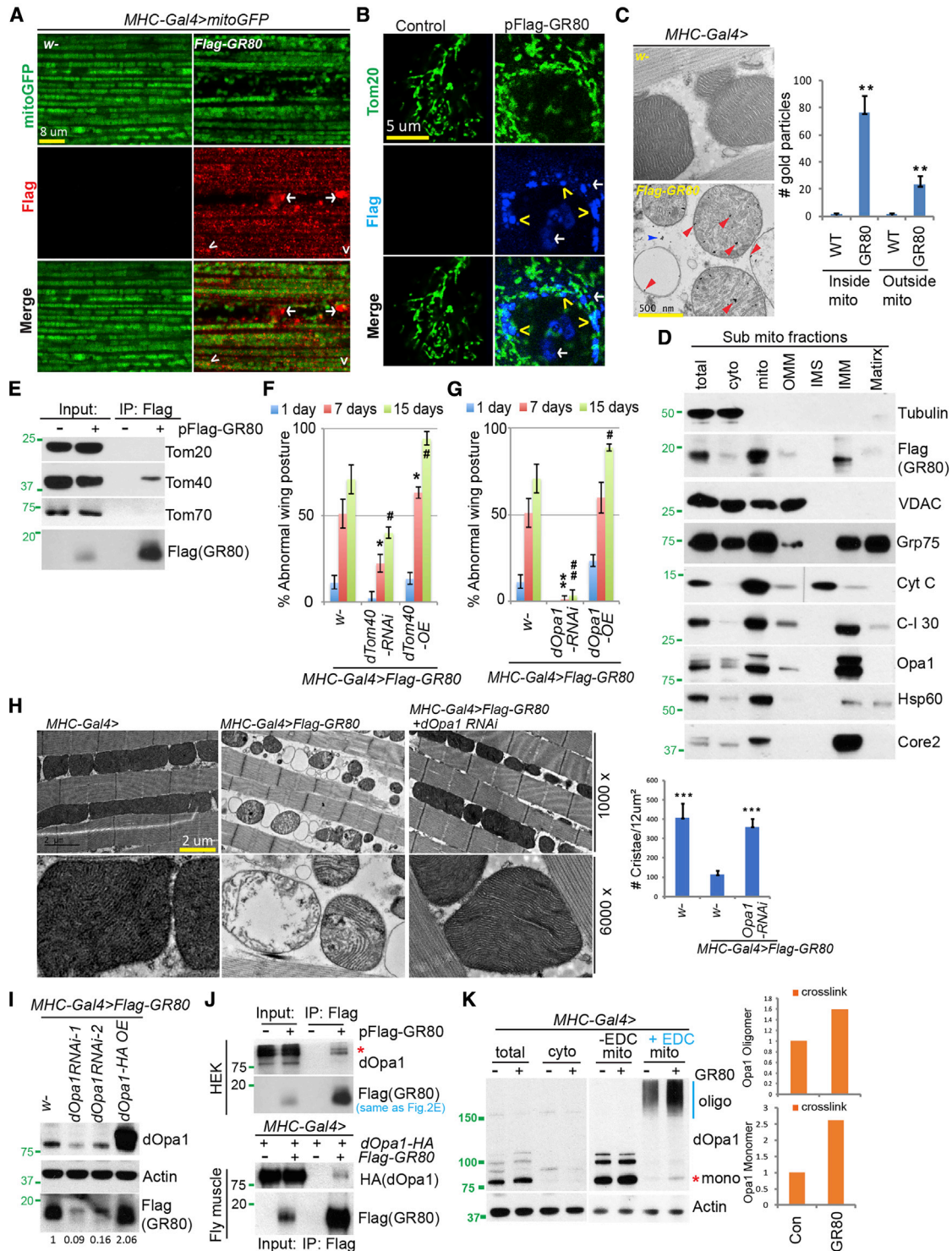
For GR80 to reach IMS/IMM, it most likely has to go through the translocase of the outer membrane/inner membrane (TOM/TIM) import complex, and interact with Tom40, a key component of the TOM complex forming the import channel. GR80 interacted with Tom40, but not Tom20 or Tom70, in co-immunoprecipitation (coIP) assays (Figure 2E). Functionally, Tom40 overexpression (OE) increased the GR80 protein level and enhanced the GR80-induced wing posture phenotype, whereas partial Tom40 knockdown by RNAi had opposite effects (Figures 2F, S2E, and S2F).

### Opa1 Mediates the Effects of Poly(GR) on CJ

We next investigated how GR80 instigates mitochondrial toxicity. By binding to Tom40, GR80 may clog the TOM/TIM channel, thereby impairing mitochondrial import globally. Inconsistent with this idea was our observation that the import of mito-GFP was not significantly affected by GR80. Given the IMS/IMM

localization of GR80 and its profound effects on CJ structure, we tested whether GR80 may alter IMS/IMM structures related to CJ formation. Using immuno-EM, we observed GR80 localization to CJ (Figure S2G). Tuning down the expression of Opa1, a key determinant of CJ remodeling (Frezza et al., 2006), completely suppressed GR80 effect on wing posture and cristae morphology, which was accompanied by a reduction in GR80 level (Figures 2G–2I). In contrast, Opa1 OE significantly enhanced GR80 toxicity and protein level (Figures 2G and 2I). Although Opa1 is known to act in mitochondrial fusion, its modulation of GR80 toxicity is most likely unrelated to that, as altering fission-fusion balance by the loss- or gain-of-function of Drp1, Fis1, or Marf had no obvious effect on GR80 toxicity (Figures S2H and S2I). These results implicate CJ as a key site of GR80 toxicity.

We next assessed the molecular basis underlying the genetic interaction between GR80 and Opa1. In coIP assays we



**Figure 2. Mitochondrial Localization and Opa1 Interaction of GR80**

(A and B) Immunostaining showing mitochondrial localization of FLAG-GR80 in transgenic fly muscle (A) or transfected HeLa cells (B). Mito-GFP (A) or Tom20 staining (B) marks the mitochondria. Arrowhead: mitochondrial localization; arrow: non-mitochondrial localization. The images are representative of 3 independent samples.

(C) Immuno-EM analysis showing mitochondrial localization of FLAG-GR80 in fly muscle. Red and blue arrows mark the signals inside and outside the mitochondria, respectively. The graph shows quantification (n = 4, 3 biological repeats).

(legend continued on next page)

detected an association between GR80 and the long-form of Opa1 (Figure 2J), suggesting that they may interact inside mitochondria. More supporting evidence came from immuno-EM, which showed a juxtaposition of GR80 and Opa1 near CJs (Figure S2J). Next, we tested the effect of GR80 on the formation of Opa1 oligomers, the presumed molecular staples that constrict and hold the inner membranes at CJs (Frezza et al., 2006). GR80 promoted Opa1 oligomer formation and also had a mild effect on the total Opa1 (monomer) level (Figures 2K and S2K). It is possible that by promoting Opa1 oligomer formation, GR80 causes moderate overall stabilization of steady-state Opa1 monomers. Consistent with the biochemical data, GR80 fly muscle mitochondria exhibited narrower and presumably tightened CJs, which could be loosened by reducing Opa1 oligomer formation through Opa1 RNAi (Figures S2L–S2N).

### Alteration of Mitochondrial Contact Site and Cristae Organizing System (MICOS) Structure and Function by Poly(GR)

The Opa1 interactome in *Drosophila* includes components of the Mic60-Mic19 subcomplex of MICOS (Banerjee and Chinthapalli, 2014), which is centrally involved in CJ formation and function (Koob and Reichert, 2014; Schorr and van der Laan, 2018). We therefore tested whether GR80 may affect MICOS. In coIP assays, GR80 exhibited a physical association with MICOS components in mammalian cells (Figure 3A) and fly muscle (Figure 3B). Functionally, as with Opa1, the partial reduction of components of the Mic60-Mic19 subcomplex effectively rescued the GR80-induced wing posture phenotype (Figure S3A). This was correlated with the restoration of cristae number and the reduction in the GR80 protein level (Figures 3C and 3D). Note that the knockdown of Mic60 and Mic19 resulted in characteristic concentric cristae, a characteristic phenotype seen when these factors are inhibited (John et al., 2005). Conversely, Mic19 (CHCHD3) OE led to an increased GR80 level and an enhanced GR80 toxicity on wing posture (Figures 3E and S3B). It thus appears that at least part of the GR80 toxicity comes from its effect on the Mic60-Mic19 subcomplex. Moreover, MICOS may affect GR80 metabolism, as the GR80 level is altered by genetic manipulations of MICOS components.

MICOS is organized into the Mic60-Mic19 and Mic10 (Minos1)-Mic27 (Apool) subcomplexes, which are presumably connected by Mic12 (QIL1) (Guarani et al., 2015). We next tested the relationship between GR80 and the Mic10-Mic27 complex. In coIP assays, we found robust GR80 interaction with Mic27 and weaker interaction with Mic10 (Figures 3F and S3C). Interac-

tion was confirmed by coIP in the reverse direction (IP with Mic27/Apool and detection of GR80) and under non-crosslinking conditions (Figure 3G). Mic27 is the only MICOS factor that exhibited strong coIP with GR80 under such conditions, suggesting that it is the more proximal subunit of MICOS directly interacting with GR80. In the presence of GR80, the interaction between Mic10 and Mic27 was weakened, whereas the interactions of Mic10 with Mic60, Mic19, Opa1, and LETM1 were strengthened (Figure 3H), suggesting that GR80 alters MICOS dynamics and intersubunit interactions. Functionally, RNAi of Mic10 or Mic27, the efficiency of which was verified by qRT-PCR (Figure S3D), reduced the GR80 protein level (Figure 3I) and rescued the GR80-induced wing posture defect (Figure S3E).

Based on the biochemical and genetic data, we hypothesized that GR80 may enter MICOS and cause the malformation and tightening of CJ, impairing the exchange of metabolites and osmolytes. This may affect ion homeostasis and, in turn, the mitochondrial membrane potential (MMP). GR80 fly mitochondria maintained a high MMP (Figures 3J and S3F) and contained a higher Ca<sup>2+</sup> level as monitored with the genetically encoded mito-GCaMP indicator and the Rhod2-AM dye (Figure S3G), in keeping with MMP being a driving force of calcium entry into mitochondria. Tuning down levels of MICOS components lowered the MMP (Figure 3K), suggesting that GR80 acts through MICOS to alter MMP and mitochondrial ion homeostasis. This is presumably mediated by GR80 interference with the activity of LETM1, a protein with K<sup>+</sup> and Ca<sup>2+</sup> transporter activity (Nowikovsky et al., 2004; Jiang et al., 2009) and an interacting partner of the Mic60/Mic19/Opa1 subcomplex (Banerjee and Chinthapalli, 2014). There was an increased mito-K<sup>+</sup> level as detected with MitoPOP (Austin et al., 2017) in GR80 fly muscle (Figure S3H). Moreover, LETM1 OE or knockdown of the MICOS component decreased the mito-K<sup>+</sup> level (Figures S3H and S3L). This was also accompanied by a reduction in GR80 level (Figure S3I), suggesting that the restoration of mitochondrial ion homeostasis may affect the MICOS structure and, in turn, GR80 stability.

To further test the roles of MMP and altered ion homeostasis in GR80 toxicity, we treated GR80 flies with nigericin, a K<sup>+</sup>/H<sup>+</sup> antiporter that can re-balance mitochondrial matrix ion levels (Nowikovsky et al., 2007). Treatment with low concentrations of nigericin effectively restored mitochondrial morphology (Figure 3L), accompanied by the rescue of the fly wing posture defect (Figure 3M). The GR80 level was also significantly reduced by nigericin treatment (Figure 3N). The nigericin effect on the GR80

(D) Immunoblots of sub-mitochondrial fractions, with various mitochondrial marker proteins showing localization pattern of FLAG-GR80.

(E) CoIP assays using FLAG-GR80-transfected HEK293T cells to show specific GR80-Tom40 interaction.

(F) Bar graph showing quantification of the effect of Tom40 RNAi or overexpression (OE) on the wing posture in GR80 flies (n = 90 flies, 3 biological repeats).

(G) Bar graph showing quantification of the effect of Opa1 RNAi or OE on GR80 induced wing posture defect (n = 90 flies, 3 biological repeats).

(H) TEM images showing effect of Opa1 RNAi on GR80-induced mitochondrial morphology defect in fly thoracic muscle. The graph shows quantification (n = 5 sections, 3 biological repeats).

(I) Immunoblots showing effect of Opa1 RNAi or OE on GR80 protein level in GR80 fly muscle.

(J) CoIP assay showing GR80-Opa1 interaction in HEK293T cells (top) and fly muscle (bottom).

(K) Immunoblots showing effect of GR80 on Opa1 oligomer formation in fly thoracic muscle. 1-Ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC) was used to cross-link Opa1. The positions of Opa1 monomer (mono) and oligomers (oligo) are indicated. The graph shows quantification (n = 2).

\*p < 0.05, #p < 0.05, \*\*p < 0.01, and ###p < 0.01, 1-way ANOVA test followed by SNK test plus Bonferroni correction. The immunoblots are representative of at least 2 independent repeats. See also Figure S2.



protein level was specific, as it did not affect the GA80 level (Figure S3J). A similar effect on GR80 was observed in mammalian cells treated with nigericin or monensin, another ionophore with a similar mechanism of action (Figure 3O). Moreover, the aberrant interactions between MICOS subunits caused by GR80 (Figure S3K) and the elevated mito- $K^+$  in *GR80* flies were rescued after nigericin treatment (Figures S3H and S3L). Thus, by restoring mitochondrial ion homeostasis, nigericin may affect the MICOS structure and, in turn, GR80 stability, similar to the effects of the genetic manipulation of MICOS, although other mechanisms of nigericin action cannot be excluded.

### Rescue of Mitochondrial Defects in C9-ALS/FTD Patient Fibroblasts by Genetic Manipulation of MICOS Components or by Nigericin Treatment

Finally, we tested whether the pathogenic mechanisms uncovered in flies are relevant to GR toxicity in patient cells. In fibroblasts from C9-ALS/FTD patients carrying *C9ORF72* G4C2 repeat expansion (Kramer et al., 2016), GR was present in the mitochondrial fraction, as shown by dot blot (Figure S4A) and immunostaining (Figure S4B), although nuclear GR was also detected (Figure S4B). There was increased mitochondrial complex I activity (Figure 4A) and ATP production (Figure S4C) in patient cells and detachment and loss of cristae in patient mitochondria, as revealed by EM (Figure 4B). There was also increased MMP, mito- $Ca^{2+}$ , mitochondrial ROS, and mito- $K^+$  in patient fibroblasts (Figure 4C). Thus, altered mitochondrial inner membrane structure and ion homeostasis are conserved features of C9-ALS. We also observed GR interactions with Mic27/Apool in patient fibroblasts (Figure S4D). Nigericin treatment (Figures 4D–4F) or knockdown of key MICOS components (Figures S4E–S4G) rescued the mitochondrial defects in patient fibroblasts. Nigericin treatment also reduced the GR-induced mito- $K^+$  increase in HEK293 cells transfected with GR80 (Figure S4H). Nigericin treatment (Figure 4G) or knockdown of key MICOS components (Figure S4I) reduced the GR level in patient cells, but nigericin had no effect on the GA level (Figure S4J). The coexistence of GR and other DPRs (GA and PR), the mitochondrial accumulation of GR, and the specific effect of nigericin on the GR level were observed in independent patient fibroblast cell lines (Figures S4K–S4M). These data support the notion that poly(GR)-induced alterations of MICOS and ion homeostasis contribute to mitochondrial toxicity in C9-ALS.

### DISCUSSION

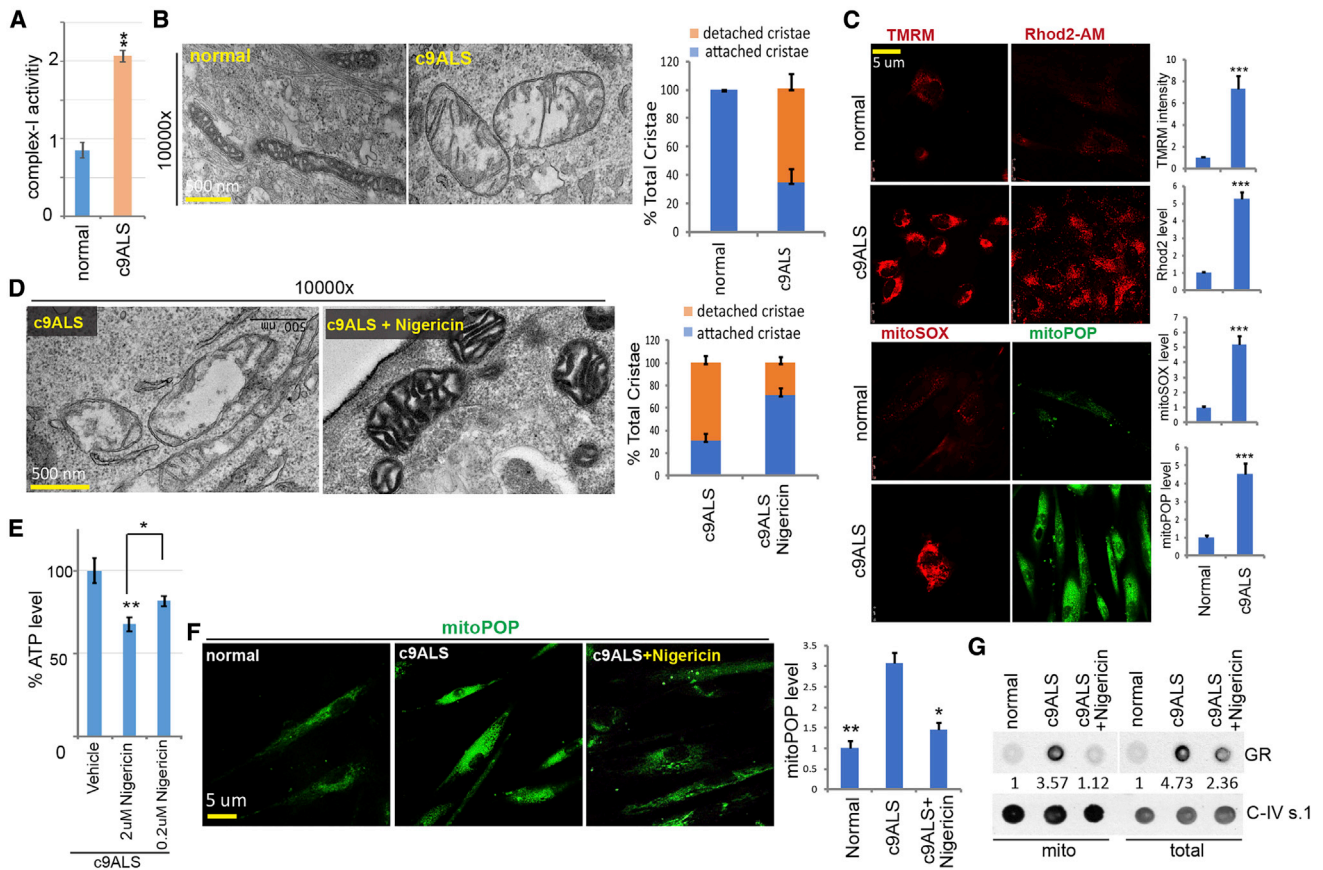
Mitochondrial dysfunction is a common and early feature observed in neurodegenerative diseases. The pathogenic role of mitochondrial dysfunction has been extensively studied in SOD1-linkd ALS (Palomo and Manfredi, 2015) and is also linked to C9-ALS/FTD (Yang et al., 2015; Konrad et al., 2017; Choi et al., 2019), but the underlying mechanism is not well defined. We show that poly(GR) can enter mitochondria and associate with MICOS, altering the balance and stoichiometry of the MICOS subcomplexes and impairing mitochondrial function. Our findings of poly(GR) entering mitochondria and causing mitochondrial toxicity resonate with previous studies (Choi et al., 2019) that identify mitochondrial protein ATP5A1 as a binding partner of poly(GR). These results support the notion that G4C2 repeat-derived DPRs can target diverse essential organelles and cellular structures to cause cellular toxicity and contribute to C9-ALS/FTD pathogenesis. The composition and length of DPRs produced in disease have not been systematically determined. It is possible that they vary in a cell-type-specific manner and may interact with different partners or interfere with different cellular structures. This may confer cell-type selectivity of vulnerability. Supporting this notion, a recent study found a specific increase in poly-GR among the DPRs tested, and evidence of mitochondrial defect, in iPSC-derived C9-ALS/FTD muscle cells (Lynch et al., 2019). Future studies will test whether MICOS defects may underlie the mitochondrial defects in patient iPSC-derived C9-ALS/FTD muscle cells.

Our results implicate a causal and direct link between MICOS dysfunction and C9-ALS/FTD. MICOSs are dynamic structures that organize the IMM into domains of a broader respiratory network. In addition to forming membrane boundaries that limit the dynamic distribution of membrane proteins, the MICOSs at CJ help partition metabolites between the intracristal spaces and IMSs and thus influence OxPhos. How these structures change their number, composition, position, morphology, and interactions with other mitochondrial complexes in response to metabolic demand or stress is poorly understood. Our results indicate that by interacting with MICOS component Mic27, poly(GR) weakens the interaction between Mic27 and Mic10 while strengthening the interaction between Mic10 and Mic60/Mic19/Opa1/LETM1 proteins. Mic10 and Opa1 have both been shown to form oligomers, which are required for membrane

### Figure 3. Effect of GR80 on MICOS and Mitochondrial Ion Homeostasis

(A and B) CoIP assay showing FLAG-GR80 interaction with MICOS components Mic60 (Mitofilin) and Mic19 (CHCHD3) in HEK293T cells (A) or between FLAG-GR80 and Myc-tagged mitofilin in fly muscle tissue (B).  
(C) TEM images showing effect of RNAi of Mic19 or Mic60 on cristae morphology in *GR80* flies.  
(D and E) Immunoblots showing effects of Mic19 or Mic60 RNAi (D) or Mic19 OE (E) on GR80 level in fly muscle.  
(F) CoIP assay showing stronger GR80-Mic27 interaction compared with GR80-Mic10 (Minos1) interaction in HEK293T cells.  
(G) Reverse coIP under non-crosslinking condition showing Mic27/GR80 interaction.  
(H) CoIP assay showing the effect of GR80 on Mic10 interaction with other MICOS components in HEK293 cells.  
(I) Immunoblots showing the effect of Mic27 or Mic10 RNAi on GR80 protein level in fly muscle.  
(J and K) Increased MMP in *GR80* fly muscle as detected with the tetraethylbenzimidazolylcarbocyanine iodide (JC-1) (J) or tetramethylrhodamine methyl ester (TMRM) (K) dye, and its rescue by the genetic manipulation of MICOS components (K). The graphs show quantification (n = 5, 3 biological repeats). \*\*\*p < 0.001 in Student's t test.  
(L and M) Effect of nigericin treatment on mitochondrial morphology (L) and wing posture (M) of *GR80* flies. \*p < 0.05,  $\chi^2$  test (n = 1).  
(N and O) Immunoblots showing the effect of nigericin on GR80 level in fly muscle (N) and the effect of nigericin or monensin on GR80 level in HEK293T cells (O). Immunoblots are representative of at least 2 independent repeats. See also Figure S3.





**Figure 4. Mitochondrial Defects in C9-ALS Patient Fibroblasts and Effect of Nigericin and Genetic Manipulation of MICOS Components**

(A) Measurement of complex I activity in control and patient fibroblasts (n = 3).  
 (B) TEM images of mitochondria in control and patient fibroblasts. The graph shows quantification (n = 3).  
 (C) Altered MMP, mito-Ca<sup>2+</sup>, mito-ROS, and mito-K<sup>+</sup> in patient fibroblasts. The graphs show quantification (n = 3).  
 (D–F) Effects of nigericin treatment on mitochondrial morphology (D), ATP level (E), or mito-K<sup>+</sup> level (F) in patient fibroblasts. The graphs show quantification (n = 4).  
 (G) Dot blot assay showing effect of nigericin on GR protein levels in total or mitochondrial fraction (top) in patient fibroblasts. Complex IV subunit 1 (upper) or actin (lower) serves as a loading control.

\*p < 0.05, \*\*p < 0.01, and \*\*\*p < 0.001, 1-way ANOVA test followed by SNK test plus Bonferroni correction (E and F), or 2-tailed Student's t test (A and C). Immunoblots are representative of at least 2 independent repeats. See also Figure S4.

bending at the CJ in the case of Mic10 (Barbot et al., 2015), and CJ remodeling in response to metabolic demand and apoptotic stimuli in the case of Opa1 (Frezza et al., 2006). We show that GR80 promotes Opa1 oligomer formation and tightening of CJ and that by interacting with MICOS components, poly(GR) itself becomes stabilized. This vicious loop may perpetuate GR toxicity in the disease state.

We find that genetic manipulation of MICOS has a profound effect on poly(GR) level, suggesting that a dynamic equilibrium between the mitochondrial and extra-mitochondrial pools of poly(GR) exists and that mitochondria play an important role in poly(GR) metabolism. How poly(GR) interacts with and impairs these multi-protein assemblies and the mechanism of MICOS and mitochondria in regulating poly(GR) metabolism are interesting questions for future investigation. Since aberrant cristae morphology is frequently seen in disease (Wollweber et al., 2017) and during aging (Takahashi et al., 1970), other disease-causing genes or the aging process may directly or indirectly

affect the MICOS to alter mitochondrial structure and function, and further studies of the mechanisms involved will offer novel insights into the pathophysiology.

Our results also implicate altered mitochondrial ion homeostasis in C9-ALS/FTD pathogenesis. The K<sup>+</sup>/H<sup>+</sup> antiporter nigericin can effectively rescue mitochondrial cristae morphology and various physiological parameters in fly models and independent patient fibroblast cell lines. Nigericin was previously shown to rescue LETM1 deficiency-induced mitochondrial swelling and ion dyshomeostasis in yeast and mammalian cells (Nowikovsky et al., 2007; Dimmer et al., 2008), suggesting that LETM1 dysfunction may be relevant to poly(GR) toxicity. Nigericin can also affect cancer-related signaling and has been explored as an anticancer agent in preclinical studies (Hegazy et al., 2016). However, given its prominent pro-inflammatory activity, its use in ALS patients may require the careful determination of an effective therapeutic window, as inflammation is implicated in ALS pathogenesis (McCauley and Baloh, 2019). Should future

studies validate the beneficial effects of MICOS modification and nigericin treatment in animal models that do not rely on GR OE and in C9-ALS/FTD patient iPSC-derived muscle cells or motor neurons, further modification of nigericin or related K<sup>+</sup>/H<sup>+</sup> antiporters is warranted to develop potential therapeutic agents.

## 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.celrep.2020.107989>.

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## AUTHOR CONTRIBUTIONS

S.L. and Z.W. designed the study, performed the experiments, analyzed the data, co-wrote the manuscript, and contributed equally to the study; Y.L. and I.T. performed the experiments and analyzed the data; D.D.S., A.M., and F.-B.G. provided the key reagents; H.V. supervised the EM and muscle histology study and analyzed the data; and B.L. conceived and supervised the study, performed the experiments, wrote the manuscript, and provided funding.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Mouse anti-FLAG	Sigma-Aldrich	F1804
Rabbit anti-FLAG	Sigma-Aldrich	F7425
Chicken anti-GFP	Abcam	ab13970
Mouse anti-TOM20	Santa Cruz Biotech	sc17764
Mouse anti-Opa1	BD Biosciences	612806
Mouse anti-Tom40	Santa Cruz Biotech	sc365467
Mouse anti-Tom70	Santa Cruz Biotech	sc390545
Mouse anti-VDAC1	Santa Cruz Biotech	sc390996
Rabbit anti-cytochrome C	Abcam	ab90529
Rabbit anti-C-I30	Abcam	ab14711
Mouse anti-Core2	Santa Cruz Biotech	sc390378
Mouse anti-alpha-tubulin	Proteintech	66031
Rabbit anti-Grp75	Cell Signaling	2816
Mouse anti-Hsp60	Santa Cruz	sc-59567
Mouse anti-C-IV s.1	Abcam	ab14705
Rat anti-HA	Roche	3F10
Mouse anti-actin	Sigma-Aldrich	A2228
Rabbit anti-mitofilin	Abcam	ab48139
Rabbit anti-CHCHD3	Abcam	ab98975
Mouse anti-ApoL	Santa Cruz Biotech	sc-390958
Mouse anti-LETM1	Abcam	ab55434
Mouse anti-Myc	Santa Cruz Biotech	9E10
Rabbit anti-Minos1	Abcam	ab84969
Rabbit anti-fly-Opa1	Sigma-Aldrich	M6319
Rat anti-poly (GR)	Millipore	MABN778
Rabbit anti-poly (GR)	Millipore	ABN1361
Rabbit anti-poly(GA)	Proteintech	24492-1-AP
Rabbit anti-poly(PR)	Proteintech	23979-1-AP
Rabbit anti-ATP6	Abcam	AB102573
Goat anti-Mouse IgG-HRP	Santa Cruz	sc-2005
Goat anti-Rabbit IgG HRP	Santa Cruz	sc-2004
Goat anti-Chicken IgY (H+L) Secondary Antibody, Alexa Fluor 488	Invitrogen	A11039
Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor 488	Invitrogen	A11034
Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor 568	Invitrogen	A11036
Goat anti-Mouse IgG (H+L) Cross-Adsorbed Secondary Antibody, Alexa Fluor 568	Invitrogen	A11004
Goat anti-Mouse IgG (H+L) Cross-Adsorbed Secondary Antibody, Alexa Fluor 633	Invitrogen	A21050

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**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
DAPI	Sigma-Aldrich	D9542
<b>Chemicals, Peptides, and Recombinant Proteins</b>		
Nigericin	Cayman Chemical	11437
Monensin sodium salt	Sigma-Aldrich	M5273
EDC (1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride)	Thermo Scientific	22980
Lipofectamine 3000	Invitrogen	L3000015
Lipofectamine RNAi-MAX	Invitrogen	13778150
Tris base	Sigma-Aldrich	11814273001
Glycine	Sigma-Aldrich	G8898
SDS	Sigma-Aldrich	L3771
Anti-FLAG M2 affinity gel	Sigma-Aldrich	A2220
Normal goat serum	Jackson ImmunoResearch	005-000-121
Pierce 16% Formaldehyde (w/v), Methanol-free	Thermo Fisher	28908
Triton™ X-100	Sigma Aldrich	T9284
DMEM, high glucose, GlutaMAX Supplement	GIBCO	10566016
Schneider's Medium	GIBCO	21720-024
Percoll	GE Healthcare	17089101
EDTA	Sigma-Aldrich	E9884
EGTA	Sigma-Aldrich	E3889
Dimethyl sulfoxide	Sigma-Aldrich	D8418
2XLaemmli sample buffer	Biorad	161-0737
Protease inhibitor cocktail	Bimake	B14012
<b>Critical Commercial Assays</b>		
ATP Bioluminescence Assay Kit HS II	Roche	11699709001
Complex I Enzyme Activity Microplate Assay Kit	Abcam	ab109721
Protease K digestion kit	Thermo Fisher	AM2548
RNeasy Mini kit	QIAGEN	74104
iScript cDNA synthesis ki	Biorad	1708890
PowerUp SYBR Green Master Mix	Applied Biosystems	A25741
Western Lightning Plus-ECL	PerkinElmer Inc.	NEL105001EA
HyBlot CL Autoradiography Film	Denville Scientific Inc.	1159M38
Tetramethylrhodamine (TMRM)	Invitrogen	T668
MitoSox	Invitrogen	M36008
Rhod-2, AM	Invitrogen	R1244
MitoPOP	Dr. Karin Nowikovsky	N/A
JC-1	Invitrogen	T3168
Q5 Site-Directed Mutagenesis Kit	New England Biolabs	E0554S
NuPAGE® MOPS SDS running buffer	Invitrogen	NP0001
NuPAGE 4-12% Bis-Tris Protein Gels	Invitrogen	NP0321
<b>Oligonucleotides</b>		
qRT-PCR primer: actin42A Forward: TCTTACTGAGCGCGGTTACAG	This paper	N/A
qRT-PCR primer: actin42A Reverse: ATGTCGCGCACAATTTAC	This paper	N/A

(Continued on next page)

<b>Continued</b>		
REAGENT or RESOURCE	SOURCE	IDENTIFIER
qRT-PCR primer: mic10 Forward: TCACCCTGCTCTTCTTCCG	This paper	N/A
qRT-PCR primer: mic10 Reverse: ACGATATGCCACGCCAC	This paper	N/A
qRT-PCR primer: Apool Forward: AATCATGGCGGCGGTTGC	This paper	N/A
qRT-PCR primer: Apool Reverse: TGTGCCTTAGCGGTTCC	This paper	N/A
Stealth RNAi siRNA Negative Control Hi GC	Invitrogen	12935-400
Stealth RNAi <sup>TM</sup> siRNA of Opa1	Invitrogen	HSS107432
Stealth RNAi <sup>TM</sup> siRNA of Letm1	Invitrogen	HSS106021
Stealth RNAi <sup>TM</sup> siRNA of Tom40	Invitrogen	HSS145636
Stealth RNAi <sup>TM</sup> siRNA of Mitofilin-IMMT	Invitrogen	HSS116992
Stealth RNAi <sup>TM</sup> siRNA of CHCHD3	Invitrogen	HSS147816
Stealth RNAi <sup>TM</sup> siRNA of Apool	Invitrogen	HSS175195
Stealth RNAi <sup>TM</sup> siRNA of Minos1	Invitrogen	AS029RSI
Recombinant DNA		
pcDNA3.1-Flag-GR80	Dr Fen-Biao Gao	N/A
pCMV6-C1Orf151 (Minos1)-DDK-Myc	Origene	RC212930
Software and Algorithms		
SPSS	IBM	<a href="https://www.ibm.com/analytics/spss-statistics-software">https://www.ibm.com/analytics/spss-statistics-software</a>
Excel	Microsoft	<a href="https://www.microsoft.com/en-us/microsoft-365/excel">https://www.microsoft.com/en-us/microsoft-365/excel</a>
Fiji ImageJ	National Institute of Health	<a href="https://imagej.nih.gov/ij/">https://imagej.nih.gov/ij/</a>
Deposited Data		
Mendeley Dataset	This Paper	<a href="https://data.mendeley.com/datasets/sd3rtdy7sn/1">https://data.mendeley.com/datasets/sd3rtdy7sn/1</a>
Experimental Models: Cell Lines		
HEK293 cell line	ATCC	CRC-1573
HeLa cell line	Gift from Dr. Richard Youle	N/A
Normal fibroblast cell line #1	Coriell Institute	ND29510
Normal fibroblast cell line #2	Coriell Institute	ND29971
C9ALS fibroblast cell line #4	Gift from Dr. Aaron Gitler	<a href="#">Kramer et al., 2016</a>
C9ALS fibroblast cell line #6	Gift from Dr. Aaron Gitler	<a href="#">Kramer et al., 2016</a>
C9ALS fibroblast cell line #7	Gift from Dr. Aaron Gitler	<a href="#">Kramer et al., 2016</a>
Experimental Models: Organisms/Strains		
<i>UAS-Flag-GR80</i>	Gift from Dr. Fen-Biao Gao	<a href="#">Yang et al., 2015</a>
<i>UAS-Flag-GA80</i>	Gift from Dr. Fen-Biao Gao	<a href="#">Yang et al., 2015</a>
<i>UAS-Flag-PR80</i>	Gift from Dr. Fen-Biao Gao	<a href="#">Yang et al., 2015</a>
<i>UAS-mito-GFP</i>	Gift from Dr. William Saxton	N/A
<i>UAS-mito-GCaMP</i>	Gift from Dr. Fumiko Kawasaki	N/A
<i>UAS-Myc-Mitofilin</i>	Gift from Dr. Xinnan Wang	N/A
<i>UAS-dOpa1-Flag</i> , from Dr. Leo Pallanck	Gift from Dr. Leo Pallanck	N/A
<i>UAS-dOpa1-RNAi</i>	Gift from Dr. Leo Pallanck	N/A
<i>UAS-GR100</i>	Bloomington Drosophila Stock Center (BDSC)	#58696
<i>UAS-PA100</i>	BDSC	#58699
<i>UAS-G4C2-36</i>	BDSC	#58688

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**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<i>UAS-PR100</i>	BDSC	#58698
<i>UAS-G4C2-36-RNA Only</i>	BDSC	#58689
<i>UAS-PA36</i>	BDSC	#58695
<i>UAS-PR36</i>	BDSC	#58694
<i>UAS-GA100</i>	BDSC	#58697
<i>UAS-GA36</i>	BDSC	#58693
<i>UAS-GR36</i>	BDSC	#58692
<i>UAS-Drp1</i>	BDSC	#51647
<i>UAS-Marf-RNAi</i>	BDSC	#55189
<i>Chchd3 RNAi</i>	BDSC	#51157
<i>Chchd3 RNAi</i>	BDSC	#38984
<i>Chchd3 EP</i>	BDSC	#11599
<i>Mitofilin RNAi</i>	BDSC	#63994
<i>Letm1 RNAi</i>	BDSC	#37502
<i>Tom40 RNAi</i>	BDSC	#26005
<i>Mic10 RNAi</i>	BDSC	#63669
<i>ApoL RNAi</i>	BDSC	#66933
<i>Minos1-HA</i>	FlyORF	F002914
<i>Letm1-HA</i>	FlyORF	F001238
<i>Tim17b RNAi</i>	VDRC	v103529

**RESOURCE AVAILABILITY**

**Lead Contact**

Further information and requests for resources and reagents may be obtained from the Lead Contact, Bingwei Lu ([bingwei@stanford.edu](mailto:bingwei@stanford.edu))

**Materials Availability**

All unique/stable reagents generated in this study are available from the Lead Contact with a completed Materials Transfer Agreement.

**Data and Code Availability**

Original/source data for all figures in the paper is available at Mendeley Data: <https://data.mendeley.com/datasets/sd3rtdy7sn/1>

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**

**Drosophila Stocks**

The indicated *UAS* RNAi and OE fly lines were crossed to *Mhc-Gal4* driver line for muscle expression. See [Key Resources Table](#) for genotypes. Fly culture and crosses were performed according to standard procedures and raised at indicated temperatures. Flies were generally raised at 25°C and with 12/12 hr dark/light cycles. Fly food was prepared with a standard receipt (Water, 17 L; Agar, 93 g; Cornmeal, 1,716 g; Brewer's yeast extract, 310 g; Sucrose, 517 g; Dextrose, 1033 g). Unless otherwise indicated in the figure legend, male flies at 2-3 weeks of age were used for the experimental procedures described.

**Cell Lines**

HeLa cell was a gift from Dr. Richard Youle (NINDS/NIH). HEK293T cells were purchased from ATCC. Control fibroblast line #1 and control fibroblast line #2 were obtained from the Coriell Institute. Control fibroblast line #1 was from a 55-year old female subject (Coriell ID ND29510). Control fibroblast line #2 was from a 61-year female subject (Coriell ID ND29971). The three C9orf72 expansion ALS patient fibroblast lines were generously shared by Dr. Aaron Gitler and were described before ([Kramer et al., 2016](#)). C9ALS fibroblast line #4 was from a 55-year old male patient, C9ALS fibroblast line #6 was from a 61-year old male patient, and C9ALS fibroblast line #7 was from a 62-year old male patient. Unless otherwise indicated, C9ALS fibroblast line#6 was used for most experiments. HEK293T cells, control fibroblasts and C9ALS patient fibroblasts were cultured under standard tissue culture conditions (1x DMEM medium - GIBCO, 10% FBS, 5% CO<sub>2</sub>, 37°C).

## METHOD DETAILS

### ***Drosophila* behavioral tests and ATP measurement**

For all wing posture assays, male flies at 1 day, 7-day and 14~15-day old were scored and all the experimental groups were aged at 25°C. 25~30 male flies were collected/raised in one vial and 3~4 independent vials were counted per genotype.

ATP measurements in fly thoracic indirect flight muscle were performed according to published protocols, using a luciferase-based bioluminescence assay (ATP Bioluminescence Assay Kit HS II, Roche Applied Science). For each test, three thoraces were carefully dissected by removing heads, wings, legs and abdomen from whole flies, and the remaining part quickly homogenized in 100  $\mu$ l lysis buffer (provided by kit). The tissue lysates were then boiled for 5 min at 100°C and briefly cleared by centrifugation at 20,000 g for 2 mins. The supernatant was transferred to a new tube and kept on ice. 2.5  $\mu$ l of cleared tissue lysate was mixed with 187.5  $\mu$ l dilution buffer and 10  $\mu$ l luciferase reagent. The luminescence signal was briefly mixed and immediately measured by a Lumat LB 9507 tube luminometer (Berthold Technologies). For each genotype or drug treatment group, at least 3~4 independent tests were assayed.

### **Immunoblots and Immunostaining**

Antibodies used for western blot and immunostaining were mouse anti-Flag (1:1,000, Sigma-Aldrich, F1804), rabbit anti-Flag (1:1,000, Sigma-Aldrich, F7425), mouse anti-TOM20 (1:1,000, Santa Cruz, sc17764), mouse anti-Tom40 (1:1000, Santa Cruz, sc365467), mouse anti-Tom70 (1:1000, Santa Cruz, sc390545), mouse anti-VDAC1 (1:1000, Santa Cruz, sc390996), rabbit anti-cytochrome C (1:1000, Abcam, ab90529), rabbit anti-C-130 (1:1000, Abcam, ab14711), mouse anti-Core2 (1:1000, Santa Cruz, sc390378), mouse anti-OPA1 (1:1000, BD Biosciences 612806), rat anti-HA (1:1000, Roche, 3F10), mouse anti-actin (1:5000, Sigma, A2228), rabbit anti-mitofilin (1:1000, Abcam, ab48139), rabbit anti-CHCHD3 (1:1000, Abcam, ab98975), mouse anti-Apool (1:1000, Santa Cruz, sc-390958), mouse anti-LETM1 (1:1000, Abcam, ab55434), mouse anti-Myc (1:1000, Santa Cruz, 9E10), rabbit anti-Minos1 (1:1000, Abcam, ab84969), rabbit anti-fly-Opa1 (1:1000, Sigma, M6319). Antibodies for dot blot: rat anti-poly (GR) (1:500, Millipore, MABN778), rabbit anti-poly(GA) (1:500, Proteintech, 24492-1-AP), rabbit anti-ATP6 (1:1000, abcam, AB102573), mouse anti-Flag (1:1000, Sigma, F1804), mouse anti-actin (1:3000, Sigma, A2228).

### **Nigericin treatment of *Drosophila***

Newly hatched flies were collected and placed into vials with instant fly food (Carolina Biological Supply Company, USA) mixed with different doses (0, 2, 10 and 50  $\mu$ M) of Nigericin (Cayman Chemical, 11437) from the stock solution (50 mM dissolved in DMSO). The flies were fed for 7 days and changed with fresh vial every 24 hours. After treatments, fly thoracic samples were harvested and used for sample preparation.

### **Fly muscle staining**

For immunohistochemical analysis of mitochondrial morphology of adult fly indirect flight muscles, 5-day old male flies from 29°C were assayed. In muscle staining, at least 5 individuals were examined for each genotype and the representative images were presented. Dissected tissue samples were briefly washed with 1x PBS and fixed with 4% formaldehyde in 1x PBS containing 0.3% Triton X-100 for 45 minutes at room temperature. Fixatives were subsequently blocked with 1x PBS containing 5% normal goat serum and incubated for 60 minutes at room temperature followed by incubation with primary antibodies at 4°C overnight. The primary antibodies used were: chicken anti-GFP (1:5,000, Abcam) and mouse anti-FLAG (1:1,000; Sigma). After three washing steps with 1x PBS/0.25% Triton X-100 each for 15 minutes at room temperature, the samples were incubated with Alexa Fluor® 594-conjugated and Alexa Fluor® 488-conjugated (1:500, Molecular Probes) for 3 hours at room temperature and subsequently mounted in Slow-Fade Gold (Invitrogen).

### **MitoSox, TMRM, Rhod2, JC-1, and mitoPOP staining and mito-GCaMP imaging**

Fly thoraxes were isolated by removing heads, wings, legs and abdomen from whole flies and dissected in the Schneider's Medium (cat#: 21720-024, GIBCO™). Samples were briefly washed once with fresh medium and stained in Schneider's medium at room temperature for 30 minutes in the dark box. The dilutions of the dyes are: MitoSox 5  $\mu$ M, TMRM 200 nM, JC-1 5  $\mu$ M, Rhod-2 5  $\mu$ M and mitoPOP 5  $\mu$ M. After staining, the samples were washed 3 times with Schneider's Medium and directly applied to observation with Leica SP8 confocal microscope. MitoSox, TMRM, Rhod2, and JC-1 were purchased from Molecular Probes. Mito-POP was described before (Austin et al., 2017). For mito-GCaMP live imaging, flies thoraxes were isolated as indicated before. The thoracic muscles were dissected in the Schneider's Medium. Samples were briefly washed once with fresh medium and then quickly observed under confocal microscope (Leica SP8). For data quantification, signal intensity was measured and calculated using NIH ImageJ.

### **Mitochondrial respiratory complexes activity assay**

The individual activity of the mitochondrial respiratory complexes in Figure 1D was measured at the service of Metabiolab Inc., (France). Mitochondrial complex- I activities shown in Figures 4A and S4G were measured by using a Complex I Enzyme Activity Microplate Assay Kit (Colorimetric) (ab109721, abcam), with the 450 nm signal measured using a plate-reader (Cytation3, BioTek Inc).



### Dot-blot assay

Samples (fly thoraxes and human fibroblasts) were lysed in cell lysis buffer [50 mM Tris-HCl, pH7.4, 150 mM NaCl, 5 mM EDTA, 10% glycerol, 1% Triton X-100, 0.1 mg/ml cycloheximide, 1x RNase inhibitor, and Complete protease inhibitor cocktail (cat#: B14012, Bimake)]. The extracts were briefly pre-cleared by centrifugation and measured concentration by the BCA method (BioRad). The sample concentrations were adjusted to 2  $\mu\text{g}/\mu\text{l}$  and mixed with 2x SDS sample buffer (BioRad). The samples were blotted on Hybond<sup>TM</sup>-C super NC membrane, air-dried, blocked in 5% dry milk (in TBST) and incubated with primary antibody for overnight at 4°C. The secondary antibody incubation and developing were performed following similar procedures as for western blots. For data quantification, signal intensity was measured and calculated using NIH ImageJ.

### Transmission EM, immune-gold EM, and analysis of mitochondrial morphology

For TEM, dissected fly adult thoraces from 2-3 week old flies, or control and C9orf72 patient fibroblasts were fixed 1 hour in modified Trump's fixative (0.1 M sodium cacodylate buffer, 1% glutaraldehyde, and 4% formaldehyde) at room temperature and kept at 4°C overnight. All samples were processed, trimmed and stained by standard methods. For immunogold labeling, samples were fixed in Karnovsky's fixative: 2% Glutaraldehyde (EMS Cat# 16000) and 4% paraformaldehyde (EMS Cat# 15700) in 0.1M Sodium Cacodylate (EMS Cat# 12300) pH 7.4 for 1 hr. The fix was replaced with cold/aqueous 1% Osmium tetroxide (EMS Cat# 19100) and were then allowed to warm to Room Temperature (RT) for 2 hr rotating in a hood, washed 3X with ultrafiltered water, then en bloc stained in 1% Uranyl Acetate at RT 2hrs while rotating. Samples were then dehydrated in a series of ethanol washes for 30 minutes each @ RT beginning at 50%, 70% EtOH then moved to 4°C overnight. They were place in cold 95% EtOH and allowed to warm to RT, changed to 100% 2X, then Propylene Oxide (PO) for 15 min. Samples are infiltrated with EMBED-812 resin (EMS Cat#14120) mixed 1:2, 1:1, and 2:1 with PO for 2 hr each with leaving samples in 2:1 resin to PO overnight rotating at RT in the hood. The samples are then placed into EMBED-812 for 2 to 4 hours then placed into molds w/labels and fresh resin, orientated and placed into 65°C oven overnight. Sections were taken around 80nm, picked up on formvar/Carbon coated slot Cu grids, stained for 40seconds in 3.5% Uranyl Acetate in 50% Acetone followed by staining in Sato's Lead Citrate for 2 minutes. For immunolabelling, sections were subjected to micro etching with 10% periodic acid in water for 15 minutes on parafilm, followed by 3x wash with ddH<sub>2</sub>O/5 min each. This was followed by elution of osmium with 10% sodium metaperiodate in H<sub>2</sub>O for 15 minutes on parafilm. After 3x washes with ddH<sub>2</sub>O/5 min each, samples were blocked in Standard Block (0.5% BSA, 0.5% Ovalbumin in PBST) for 30 min, quenched with 0.5M glycine in PBS for 10 min, washed with PBS 2x/2 min each, and blocked again with Standard Block for 20 min. Primary antibodies were added at 1:1000 for mouse anti-Flag and 1:50 for rabbit anti-Opa1 and incubated at 4°C O/N. Samples were washed 2x in Standard Block/10 min each, 2x in PBS/10 min each, and then incubated with secondary antibodies (EM GAR15 and EM GMHL5) at 1:20 for 1 hour and 20 min at RT. Samples were washed 2x with PBST/5 min each, 2x with ultrapure H<sub>2</sub>O (drop then stream), and contrast stained with 3.5% Uranyl Acetate in 50% Acetone (35 s then rinse in stream) and 0.2% Lead Citrate (35 s then rinse in stream). Samples were observed in the JEOL JEM-1400 120kV. Images were taken using a Gatan Orius 832 4k X 2.6k digital camera with 9 $\mu\text{m}$  pixel.

For mitochondrial morphology analyses, in Figures 1C and 2H, the cristae number of mitochondria in indirect flight muscle were counted in the TEM images with the magnification of 6,000 x. A 2  $\mu\text{m}$  x 6  $\mu\text{m}$  rectangle area was selected in the TEM mitochondrial images and all the mitochondrial cristae in this area were counted. 5 individual samples were counted per genotype. In Figure 2C, the number of gold particles in indirect flight muscle samples were counted in the TEM images with the magnification of 6,000 x. A 2  $\mu\text{m}$  x 4  $\mu\text{m}$  rectangle area was selected in the TEM images and the gold particles in this area were counted and categorized as inside mitochondria or outside mitochondria. 4 individual samples were counted per genotype. In Figure S2G, the gold particle numbers in indirect flight muscle samples were counted in the TEM images with the magnification of 8,000 ~12,000 x. A 2  $\mu\text{m}$  x 4  $\mu\text{m}$  rectangle area was selected in the TEM images and the gold particles in this area which were close to the mitochondrial inner membrane (less than 40 nm) were counted. 4 individual samples were counted per genotype. In Figure S2J, the gold particle numbers of GR80 and OPA1 in fly indirect flight muscle samples were counted in TEM images with the magnification of 8,000 ~12,000 x. A 2  $\mu\text{m}$  x 4  $\mu\text{m}$  rectangle mitochondria area was selected and the gold particles in this area that were close to the mitochondrial inner membrane (less than 40 nm) were counted. 4 individual samples were counted per genotype. In Figures S2L and S2M, the cristae junction diameters in fly indirect flight muscle samples were measured by ImageJ in the TEM images with the magnification of 8,000 ~12,000 x, n = 22 ~25 cristae junctions from mitochondria of 4 different samples were measured per genotype. Methods similar to mitochondrial morphology quantification in fly muscle samples were used for patient fibroblast analysis in Figures 4B, 4D, and S4D. The total cristae number as well as attached and detached cristae number of mitochondria in the normal and C9ALS patient fibroblast were counted at magnification of 5,000 x, A 2  $\mu\text{m}$  x 6  $\mu\text{m}$  rectangle area was selected from the TEM images for analysis. All the mitochondrial cristae in the area were counted. 6 individual samples were counted per genetic background.

### Cell lines, plasmids, cell culture and cell transfection conditions

Regular HeLa cells and HEK293T cells (ATCC) were cultured under standard conditions (1x DMEM medium, 5% FBS, 5% CO<sub>2</sub>, 37°C). HeLa and HEK293T Cell transfections were performed by using Lipofectamine 3000 (cat#: L3000015, Invitrogen), and si-RNA knockdown experiments were performed using Lipofectamine RNAiMAX reagent (cat#: 13778150, Invitrogen), according to manufacturer's instructions.

The *pcDNA3-Flag-GR80* plasmid was reported before (Yang et al., 2015). The *Minos1-Myc* plasmid was modified from the *pCMV6-C1Orf151 (minos1)-DDK-Myc* (Origene-RC212930) plasmid by deleting the DDK sequence using the Q5 Site-Directed Mutagenesis Kit (E0554S, NEB).

### RNAi in mammalian cells and patient fibroblasts

All siRNAs used for the RNAi experiments were purchased from Thermo Fisher. Briefly, HeLa cells and patient fibroblast cells were transfected with lipofectamine RNAiMAX reagent (Invitrogen, 13778) according to standard protocol. After 72 hours transfection, cells were washed with warm PBS, followed by lysis and western blot analysis. siRNA sequences used for individual are as follows: siOPA1(HSS107432), siLetm1(HSS106021), siTom40(HSS145636), siMitofilin-IMMT(HSS116992), siCHCHD3(HSS147816), siApoL(HSS175195), siMinos1(AS029RSI). C9-ALS patient fibroblasts and matched control fibroblasts were described before (Kramer et al., 2016) and kindly provided by Dr. Aaron Gitler.

### RT-PCR

For qRT-PCR analysis of RNAi efficiency, total RNAs were extracted from fly thoraxes by using the RNeasy® Minikit (QIAGEN), and cDNAs were obtained by reverse transcription using the iScript cDNA synthesis kit (Bio-Rad). cDNA templates were then subjected to reaction with PowerUp SYBR Green Master Mix (Applied Biosystems) and analyzed by StepOnePlus real-time PCR system following manufacture's standard PCR thermal cycling procedure. Calculated data were obtained by using StepOne software V2.3. Relative mRNA levels normalized to *actin* for statistical quantification in Excel. PCR primers used are as follows:

Actin42A sense: 5'TCTTACTGAGCGCGGTTACAG3'  
 Actin42A anti-sense: 5'ATGTCGCGCACAAATTCAC3'  
 Mic10 sense: 5' TCACCCTGCTTCTTCCG3'  
 Mic10 anti-sense: 5' ACGATATGCCACGCCAC3'  
 ApoL sense: 5'AATCATGGCGGCGGTTGC3'  
 ApoL anti-sense: 5'TGTGCCTTAGCGGTTCC3'

### Mitochondrial assays

Tetramethylrhodamine (TMRM) (T668) was purchase from Invitrogen. For cell staining, normal and C9orf72 patient fibroblast were cultured in standard condition. One day later, cell growth medium was removed and 100 nM TMRM solution was added into cell to incubate for 30min. Live cell image was taken under 574 nm emission filter by using Leica SP8.

For muscle staining, fly muscles with various genetic manipulations were dissected in cold 1xPBS. Muscles were subsequently incubated with 100 nM TMRM solution in PBS 30min at room temperature, followed by taking images using Leica SP8 confocal microscope. Red mitochondrial superoxide indicator-mitoSOX (M36008) was obtained from Molecular Probes. Briefly, normal and C9orf72 patient fibroblast were cultured in 24 well plates with 12 mm coverslip. Cells were incubated with 1 $\mu$ M mitoSOX for 10min at 37°C. Later on, cells were washed with warm PBS for 3 times. Live cell images were taken using the 580 nm emission filter of Leica SP8 confocal microscope. For data quantification, signal intensity was measured and calculated using NIH ImageJ.

### Mitochondria isolation and cross-linking

Intact mitochondria from *in vitro* human cells and fly tissues were purified and quality controlled for the absence of contamination by other organelles according to established procedures. For analysis of fly samples, male flies at appropriate ages were used for thoracic muscle dissection. Samples were homogenized using a Dounce homogenizer. After two steps of centrifugation (1,500 g for 5 minutes and 13,000 g for 17 minutes), the mitochondria pellet was resuspended and washed twice with HBS buffer (5 mM HEPES, 70 mM sucrose, 210 mM mannitol, 1 mM EGTA, 1x protease inhibitor cocktail), then resuspended and loaded onto Percoll gradients. After centrifugation (16700 rpm, 15 minutes, Beckman SW-40Ti rotor), the fraction between the 22% and 50% Percoll gradients containing intact mitochondria was carefully transferred into a new reaction tube, mixed with 2 volumes of HBS buffer, and pelleted by centrifuging at 20,000 g for 20 minutes at 4°C to collect the mitochondrial samples for further analyses.

After purification, mitochondrial samples were resuspended and incubated with different cross-linking reagents – 1% Formaldehyde in pre-chilled HBS, 10 mM EDC (#22980, Thermo Fisher) in HBS buffer. The samples were incubated under 25°C and stopped at different time point by adding SDS-sample buffer and boiling. The reaction products were subjected to different analyses by following established procedures.

For the mitoplast preparation and proteinase K sensitivity assays, HEK293T cells were transfected with Flag-GR80 plasmid. After 48h transfection, mitochondria fraction was isolated. Purified mitochondria were re-suspended in a hypotonic buffer containing 20 mM HEPES-KOH pH 7.2 and incubated on ice 30 minutes to allow for mitochondrial swelling. The mitoplasts were isolated by centrifugation at 64,000 g for 30 minutes. Crude mitochondria or mitoplasts were treated with 1 $\mu$ g/ml proteinase K and incubated on ice for 30 min. The reactions were terminated by adding 2x SDS loading buffer (Biorad), followed by SDS-PAGE and western blot analyses.

### Immunohistochemical analysis of cultured cells

For immunohistochemical analysis of human cells, HeLa cells, HEK293 cells, or human fibroblasts were cultured on the ethanol-cleaned cover glass. Cells were washed with 1x PBS 3 times and fixed with 4% formaldehyde in 1x PBS for 30 minutes at room temperature, later washed and permeabilized with 1x PBS containing 0.25% Triton X-100 for 15 minutes. The fixed samples were subsequently blocked with 1x PBS containing 5% normal goat serum and incubated for 1 hour at room temperature followed by

incubation with primary antibodies at 4°C overnight. The primary antibodies used were mouse anti-Flag (1:1,000, Sigma-Aldrich), rabbit anti-Flag (1:1,000, Sigma-Aldrich), rabbit anti-TOM20 (1:1,000, Santa Cruz), rabbit anti-GR (ABN1361, Sigma-Aldrich, 1:300). The secondary antibodies used were Alexa Fluor® 488, 594 and 633-conjugated antibodies (1:500, Molecular Probes).

#### Co-immunoprecipitation (co-IP), SDS-PAGE, and western blot analyses

For Flag-GR80 co-IP, HEK293T cells were transiently transfected with *pcDNA3.1-Flag-GR80* plasmid. 72 hours post-transfection, cells were either processed directly for lysis (non-crosslinking condition), or subjected to cross-linking by applying UV cross-linking and 0.5% formaldehyde in 1x PBS to the attached cells on the Petri dish (crosslinking condition). We homogenized the cells in the lysis buffer [50 mM Tris-HCl, pH7.4, 150 mM NaCl, 5 mM EDTA, 10% glycerol, 1% Triton X-100, 0.1mg/ml cycloheximide, 1x RNase inhibitor, and Complete protease inhibitor cocktail (cat#: B14012, Bimake)], additional Phosphatase Inhibitor Cocktail (cat#: B15001, Bimake) will be applied if phosphorylation signal is to be detected. After centrifugation at 10,000 g for 5 min, the supernatant was subjected to immunoprecipitation using M2 affinity agarose (Anti-FLAG M2 affinity gel, cat#: A2220, Sigma-Aldrich) at 4°C overnight with gentle shaking. Subsequently, the beads were washed three times (10 minutes each) at 4°C in lysis buffer, mixed with 2x SDS Sample buffer, and loaded onto SDS-PAGE gels.

For denaturing-IP, after cross-linking with 1% FA, samples were denatured by adding 2x SDS-sample buffer and boiling, then diluted 6 times with pre-chilled cell lysis buffer. IP experiment was performed by providing 35 µl M2-FLAG beads (Sigma) and incubated at 4°C overnight.

NuPAGE 4%–12% Bis-Tris Protein Gels (cat#: NP0321BOX, Invitrogen) and NuPAGE® MOPS SDS running buffer (cat#: NP0001, Invitrogen) were used for SDS-PAGE and immunoblot analyses according to standard procedures. For data quantification of western blots, signal intensity was measured and calculated using NIH ImageJ.

#### QUANTIFICATION AND STATISTICAL ANALYSIS

All analyses were performed with SPSS (IBM, USA). Error bars represent standard deviation (SD). For pairwise comparisons, we used two-tailed Student's *t* test. For comparing multiple groups, we used one-way ANOVA test followed by Student–Newman–Keuls test (SNK test) plus Bonferroni correction (multiple hypotheses correction). Data in [Figure S2M](#) was analyzed and presented by the Box-and-Whisker Plot. In statistical comparisons, \*, # and & indicating  $p < 0.05$  and \*\*, ## and && indicating  $p < 0.01$  were considered as significant differences.

**Cell Reports, Volume 32**

**Supplemental Information**

**Altered MICOS Morphology and Mitochondrial Ion**

**Homeostasis Contribute to Poly(GR) Toxicity**

**Associated with C9-ALS/FTD**

**Shuangxi Li, Zhihao Wu, Yu Li, Ishaq Tantray, Diego De Stefani, Andrea Mattarei, Gopinath Krishnan, Fen-Biao Gao, Hannes Vogel, and Bingwei Lu**

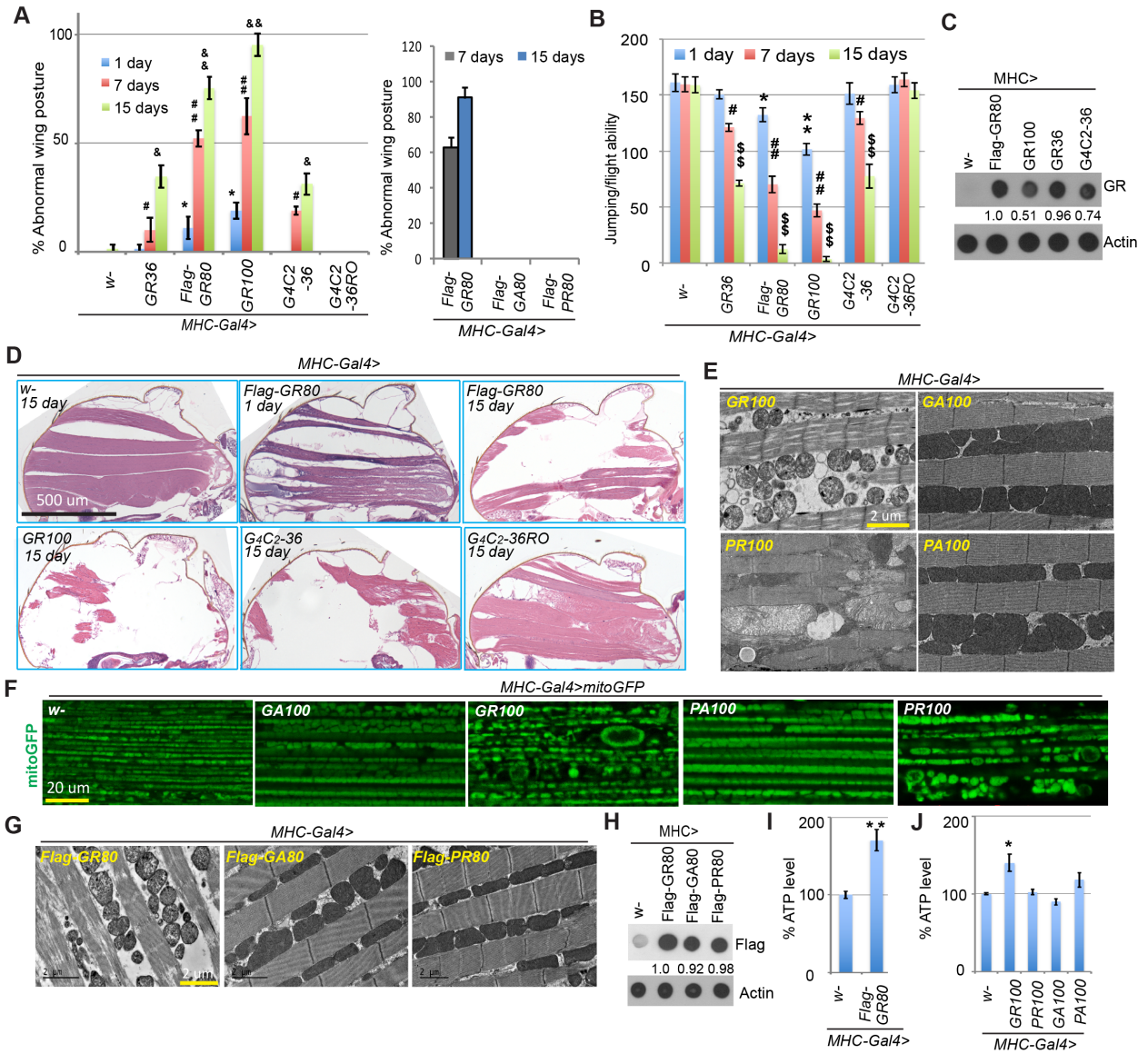
## **Supplemental Materials**

### **This file includes:**

Supplemental Figures S1 to S4

Legends for Supplemental Figures S1 to S4

Figure S1



**Figure S1. Analyses of transgenic flies expressing G4C2 repeats, GR or other DPRs,  
Related to Figure 1.**

(A, B) Analyses of wing posture (A) and jumping/flight ability (B) of the various transgenic flies (n=30 males in A and 10 males in B; 3 biological repeats).

(C) Dot blots quantifying GR expression in the various transgenic fly lines.

(D) H&E staining of paraffin sections of thoracic muscle showing effect of various transgene expression on indirect flight muscle integrity in the corresponding genotypes shown in A.

(E) TEM images of IFM mitochondrial morphology in the various transgenic flies.

(F) Monitoring of mitochondrial morphology in thoracic muscle of various transgenic flies co-expressing a mito-GFP reporter.

(G) TEM images of IFM mitochondrial morphology in transgenic flies expressing Flag-tagged GR80, GA80, and PR80.

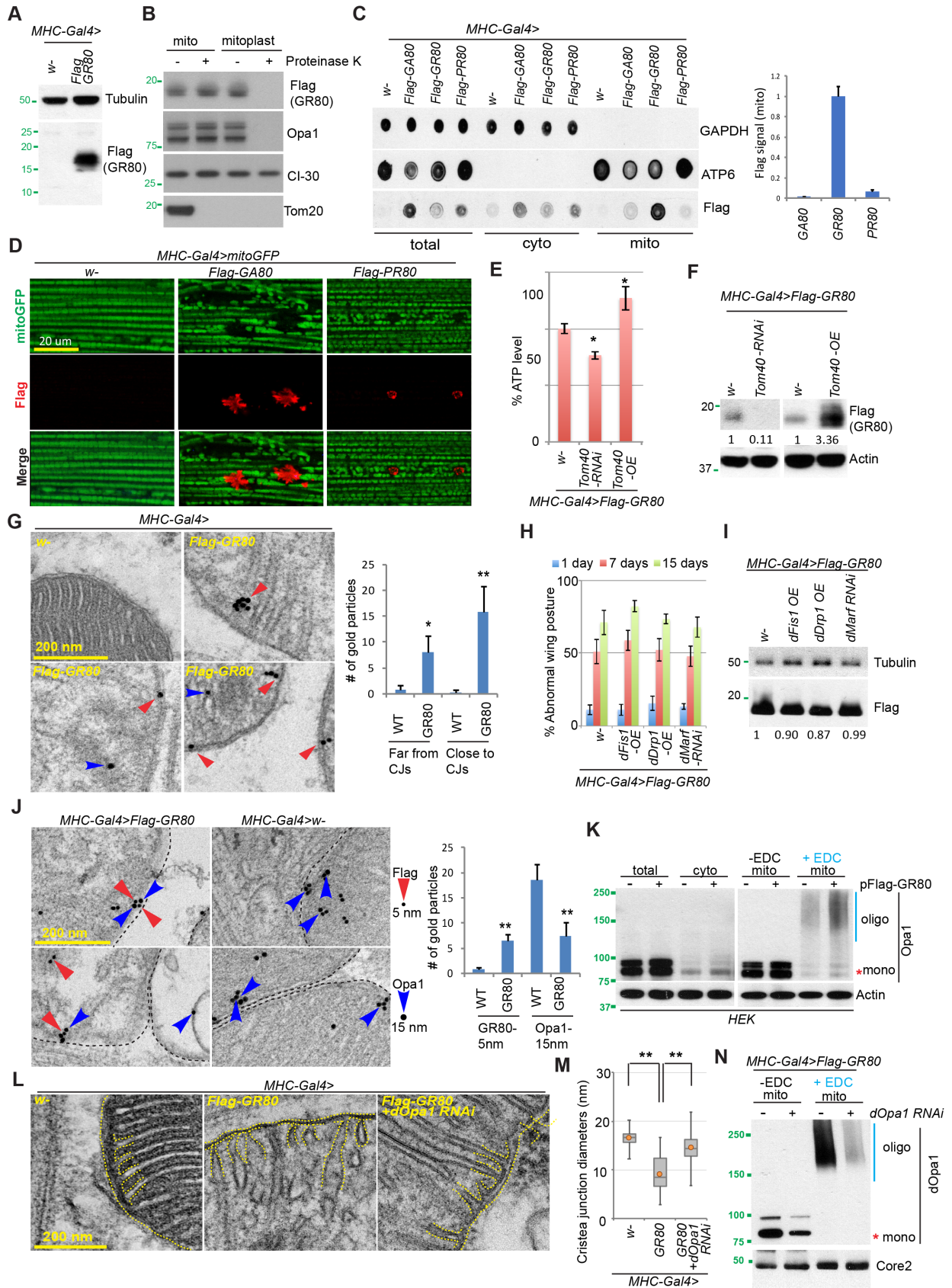
(H) Dot blots quantifying Flag-tagged GR80, GA80, and PR80 expression in the transgenic fly lines.

(I, J) Quantification showing increased ATP level in *GR80* (I) and *GR100* (J) flies.

(n=30 flies, 4 biological repeats).

\*, #, &, and \$ indicate  $p < 0.05$ ; \*\*, ##, &&, and \$\$ indicate  $p < 0.01$  in one-way ANOVA test followed by Student–Newman–Keuls test (SNK test) plus Bonferroni correction. Immunoblots are representative of at least two independent repeats. Images in D-G are representative of three independent samples.

Figure S2





**Figure S2. Additional analysis of the mitochondrial localization and Opa1 interaction of GR80, Related to Figure 2.**

(A) Immunoblots showing robust expression of Flag-GR80 in fly thoracic muscle extract from *GR80* flies. Tubulin serves as loading control.

(B) Proteinase K sensitivity assay of intact mitochondria and mitoplasts showing similar behavior of GR80 and IMS-facing Opa1.

(C) Dot blots showing the presence of GR80, but not GA80 or PR80, in the mitochondrial fraction prepared from thoracic muscle of transgenic flies. GAPDH serves as cytosolic marker; mt:ATP6 serves as mitochondrial marker. The residual Flag signals in Flag-GA80 and Flag-PR80 mitochondrial fractions are largely background signals. Graph shows quantification (n=3).

(D) Immunostaining showing lack of mitochondrial localization of Flag-GA80 or Flag-PR80 in fly thoracic flight muscle expressing a mito-GFP reporter. Images are representative of three independent samples.

(E) Bar graph showing quantification of the effect of Tom40 RNAi or OE on ATP level in *GR80* flies (n=4).

(F) Immunoblots showing the effect of Tom40 RNAi or OE on GR80 protein level in fly thoracic muscle.

(G) Immune-EM images showing mitochondrial localization of Flag-GR80 in fly thoracic muscle. Arrowheads indicate GR80 localized to the matrix (blue) or close to the cristae (red). Graph shows quantification (n=4).

(H, I) Lack of obvious effect on wing posture (H) or GR80 protein expression (I) by genetic manipulation of mitochondrial fission or fusion related proteins (n=3).

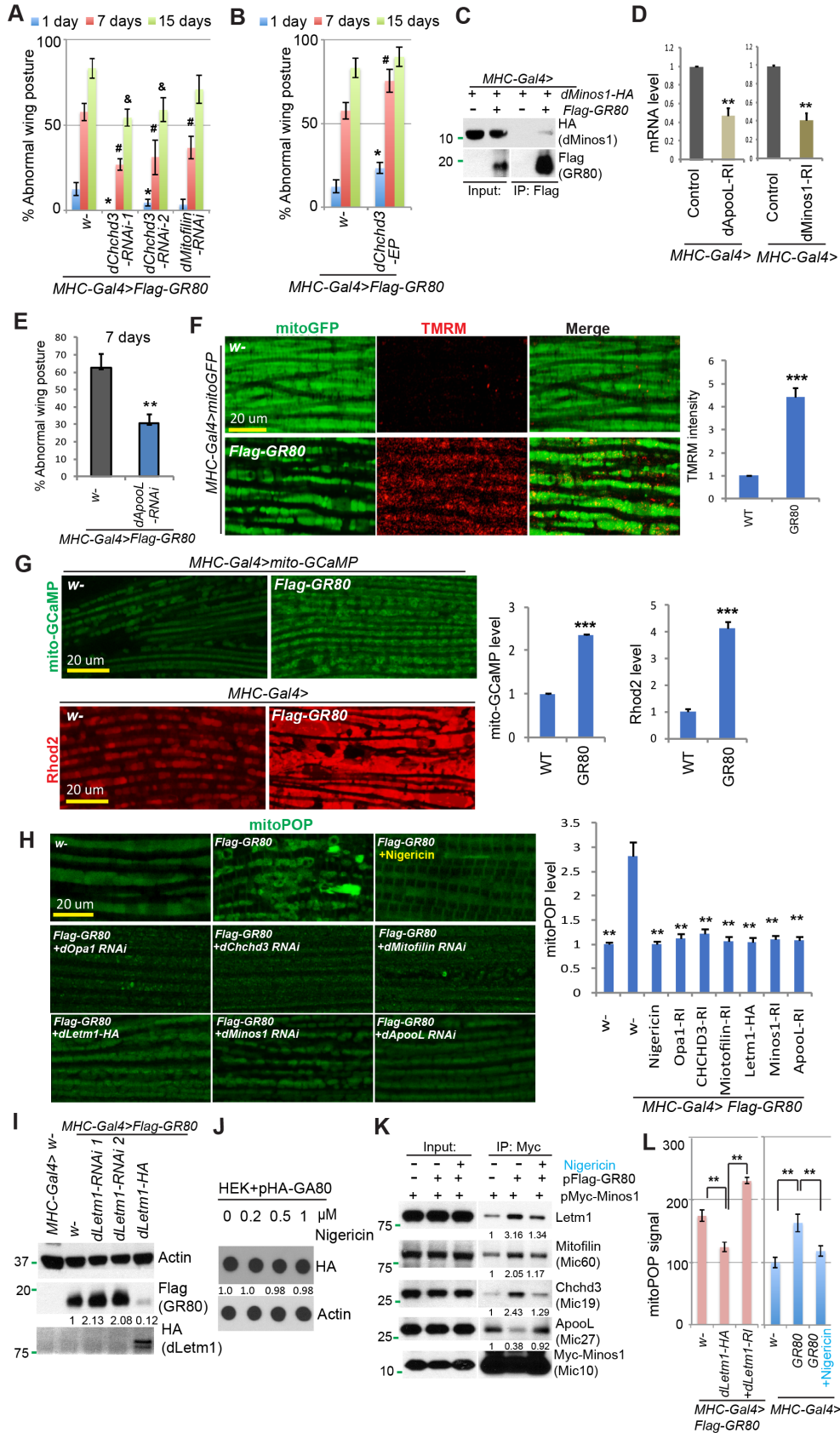
(J) Double-label immune-EM analysis of Flag-GR80 and Opa1 co-localization on mitochondrial membrane in fly thoracic muscle. Red and blue arrows mark Flag-GR80 and Opa1 signals, respectively, labeled with different sized gold particles. Graph shows quantification (n=4).

(K) Immunoblots showing effect of Flag-GR80 expression on Opa1 oligomer formation in HEK293T cells. EDC was used to cross-link Opa1 proteins. The positions of Opa1 monomers and oligomers are indicated.

(L-N) Effect of Opa1 RNAi on CJ morphology (L), CJ diameter (M), or Opa1 oligomer formation (N) in *GR80* thoracic muscle (n=22~25 CJs analyzed, 3 biological repeats for M).

\* and \*\* indicate  $p < 0.05$  and  $P < 0.01$  in one-way ANOVA test followed by Student–Newman–Keuls test (SNK test) plus Bonferroni correction. Data in M was analyzed and presented by the Box-and-Whisker Plot. \*\*,  $p < 0.01$ . Immunoblots are representative of at least two independent repeats.

Figure S3



**Figure S3. Additional data on GR80 effects on MICOS and mitochondrial ion homeostasis, Related to Figure 3.**

(A, B) Bar graphs showing quantification of the effects of RNAi-mediated knockdown of CHCHD3 and mitofilin (A) or OE of CHCHD3 with an EP line (B) on wing posture in *GR80* flies (n=90 flies analyzed, 3 biological repeats).

(C) Immunoblots showing Co-IP between Flag-GR80 and Minos1-HA in fly muscle.

(D) Quantification of mRNA level by qRT-PCR showing the knockdown efficiency of the respective RNAi lines (n=3).

(E) Bar graph showing quantification of the effect of Mic27 (Apool) RNAi on the wing-posture phenotype of *GR80* flies (n=3).

(F) Confocal images showing TMRM staining of MMP in *GR80* fly muscle. Graph shows quantification (n=6).

(G) Images showing mito- $\text{Ca}^{2+}$  levels in *GR80* fly muscle monitored using mito-GCaMP reporter or Rhod2-AM dye. Graphs show quantification (n=3).

(H) Images showing mito- $\text{K}^{+}$  level monitored using mito-POP in *GR80* fly muscle after nigericin treatment or after genetic manipulation of MICOS components. Graph shows quantification (n=3).

(I) Immunoblots showing effects of knockdown or OE of LETM1 on GR80 protein expression in fly thoracic muscle.

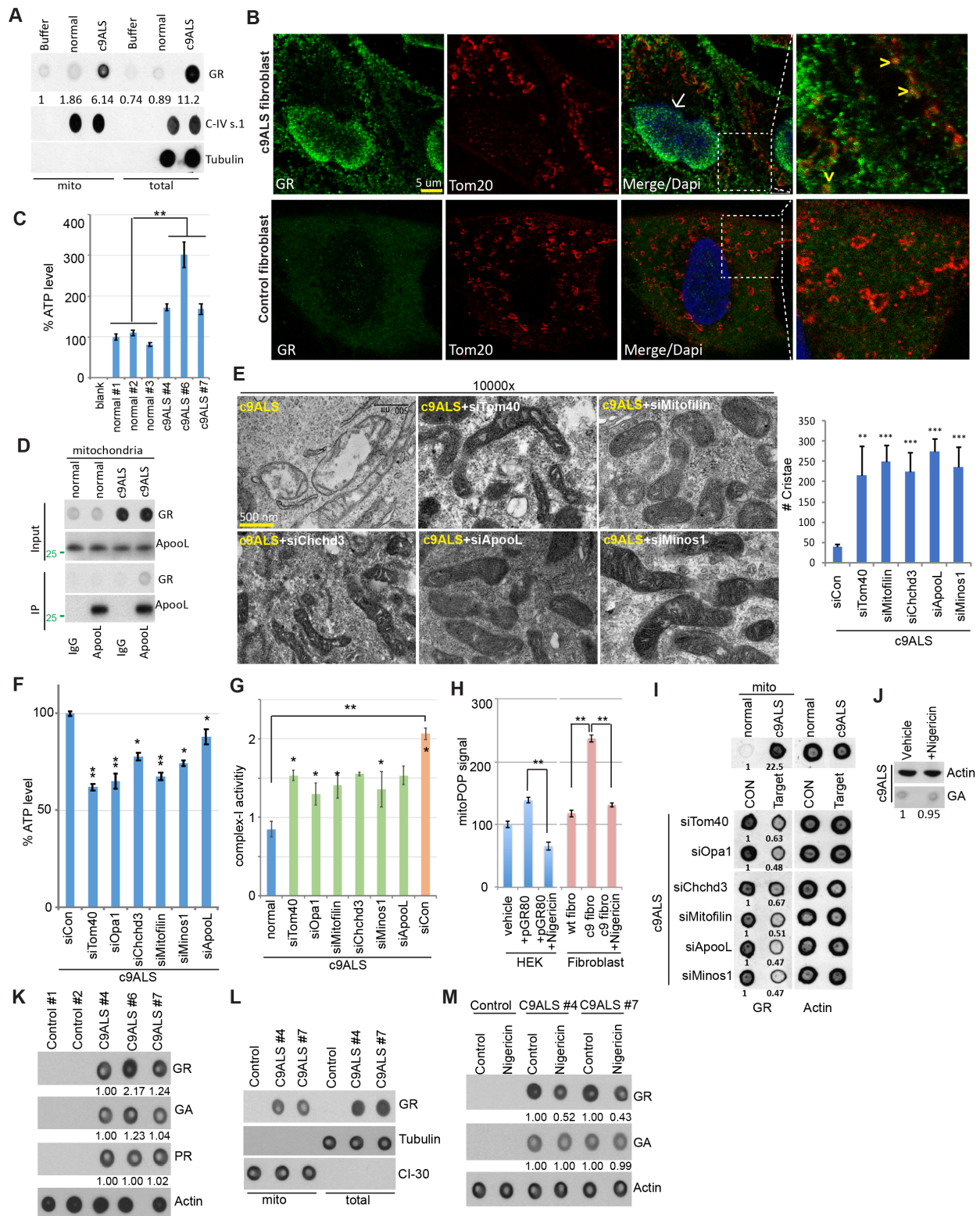
(J) Dot blot showing lack of effect of nigericin on GA80 level in HEK293 cells.

(K) Immunoblots showing effect of nigericin treatment on the interaction between Minos1 and MICOS components in transfected HEK293T cells.

(L) Bar graph showing quantification of mito-K<sup>+</sup> level in purified mitochondria from *GR80* fly muscle with or without genetic manipulation of LETM1 (left) or with or without nigericin treatment (right) (n=3).

\*, #, &, and \$ indicate p<0.05; \*\* and \*\*\* indicate p<0.01 and P<0.001, respectively, in one-way ANOVA test followed by Student–Newman–Keuls test (SNK test) plus Bonferroni correction. Immunoblots are representative of at least two independent repeats.

Figure S4



**Figure S4. Effect of genetic manipulation of MICOS components on mitochondrial defects in C9ALS patient fibroblasts, Related to Figure 4.**

(A) Dot blot assay analyzing GR peptide expression in the mitochondria of C9ALS patient fibroblasts. Complex-IV, subunit 1 (C-IV s.1) serves as mitochondrial marker, tubulin as cytosolic marker.

(B) Immunostaining showing GR expression in patient fibroblasts and its localization to mitochondrial and nuclear compartments. Arrowhead and arrow mark mitochondrial and nuclear signals of GR, respectively. Images are representative of three independent samples.

(C) Quantification of ATP level in normal and C9ALS patient fibroblasts (n=3).

(D) Co-IP assay showing Apool/GR interaction in patient fibroblasts. Apool is detected by standard western blot, GR by dot blot.

(E) TEM images showing aberrant mitochondrial morphology in C9ALS patient fibroblasts and the effect of siRNA-mediated knockdown of various MICOS components in rescuing the cristae loss phenotype of patient cells. Graph shows quantification (n=10, 3 biological repeats).

(F) Quantification of ATP level in C9ALS fibroblasts showing effects of siRNA-mediated knockdown of various MICOS components. Scrambled siRNA (siCON) serves as control (n=3).

(G) Measurements of complex-I activity in C9ALS fibroblasts showing effects of siRNA-mediated knockdown of various MICOS components (n=3).

(H) Measurement of mito-K<sup>+</sup> level in purified mitochondria from HEK293T cells transfected with GR80 and treated with nigericin or LETM1 siRNA, or from control and C9ALS fibroblasts with or without nigericin treatment (n=4).

(I) Dot blot analysis showing effects siRNA-mediated knockdown of various MICOS components on GR peptide expression. Actin serves as loading control.

(J) Dot blot assay showing lack of effect of nigericin on GA level in patient fibroblasts.

(K) Dot blot assays showing the co-existence of GR, GA, and PR in three independent patient fibroblast cell lines. Note that line #6 is used in most experiments. Actin serves as loading control.

(L) Dot blot assays showing the presence of GR in the mitochondrial fraction of additional patient fibroblast cell lines #4 and #7. C-I30 and tubulin serve as mitochondrial and cytosolic markers.

(M) Dot blot assays showing the reduction of GR but not GA level by nigericin treatment in additional patient fibroblast cell lines #4 and #7. Actin serves as loading control.

\*, \*\*, and \*\*\* indicate  $p < 0.05$ ,  $p < 0.01$  and  $P < 0.001$ , respectively, in one-way ANOVA test followed by Student–Newman–Keuls test (SNK test) plus Bonferroni correction. Immunoblots are representative of at least two independent repeats.