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Authors

Joshi-Barr, Shivanjali
Bett, Cyrus
Chiang, Wei-Chieh
et al.

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Shivanjali Joshi-Barr, Cyrus Bett, Wei-Chieh Chiang,
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Luberski, Alex Raeber, Jonathan H. Lin, Eliezer Masliah and
Christina J. Sigurdson

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De Novo Prion Aggregates Trigger Autophagy in Skeletal Muscle

Shivanjali Joshi-Barr,^a Cyrus Bett,^a Wei-Chieh Chiang,^a Margarita Trejo,^b Hans H. Goebel,^c Beata Sikorska,^d Pawel Liberski,^d Alex Raeber,^e Jonathan H. Lin,^a Eliezer Masliah,^b Christina J. Sigurdson^{a,f}

Departments of Pathology^a and Neuroscience,^b School of Medicine, University of California, San Diego, La Jolla, California, USA; Department of Neuropathology, Johannes Gutenberg-University Medical Center, Mainz, Germany^c; Department of Pathology, Medical University of Lodz, Lodz, Poland^d; Prionics AG, Zürich, Switzerland^e; Department of Pathology, Immunology and Microbiology, University of California—Davis, Davis, California, USA^f

ABSTRACT

In certain sporadic, familial, and infectious prion diseases, the prion protein misfolds and aggregates in skeletal muscle in addition to the brain and spinal cord. In myocytes, prion aggregates accumulate intracellularly, yet little is known about clearance pathways. Here we investigated the clearance of prion aggregates in muscle of transgenic mice that develop prion disease *de novo*. In addition to neurodegeneration, aged mice developed a degenerative myopathy, with scattered myocytes containing ubiquitinated, intracellular prion inclusions that were adjacent to myocytes lacking inclusions. Myocytes also showed elevated levels of the endoplasmic reticulum chaperone Grp78/BiP, suggestive of impaired protein degradation and endoplasmic reticulum stress. Additionally, autophagy was induced, as indicated by increased levels of beclin-1 and LC3-II. In C2C12 myoblasts, inhibition of autophagosome maturation or lysosomal degradation led to enhanced prion aggregation, consistent with a role for autophagy in prion aggregate clearance. Taken together, these findings suggest that the induction of autophagy may be a central strategy for prion aggregate clearance in myocytes.

IMPORTANCE

In prion diseases, the prion protein misfolds and aggregates in the central nervous system and sometimes in other organs, including muscle, yet the cellular pathways of prion aggregate clearance are unclear. Here we investigated the clearance of prion aggregates in the muscle of a transgenic mouse model that develops profound muscle degeneration. We found that endoplasmic reticulum stress pathways were activated and that autophagy was induced. Blocking of autophagic degradation in cell culture models led to an accumulation of aggregated prion protein. Collectively, these findings suggest that autophagy has an instrumental role in prion protein clearance.

Infectious prions cause severe neurodegeneration that is ultimately fatal following a prolonged incubation period (1, 2). Prion diseases occur when the host-encoded monomeric prion protein, PrP^C, misfolds and aggregates into a β -sheet-rich multimer, PrP^{Sc}, in an autocatalytic process that leads to widespread PrP^{Sc} deposition as well as neuronal death, spongiosis, astrogliosis, and microglial activation in the brain and spinal cord (3–7).

PrP^{Sc} accumulates in neural and other tissues, including skeletal muscle, lymph nodes, spleen, gastrointestinal tract, and adrenal glands, depending on the type of prion disease (8–11). The degenerative lesions, however, are largely restricted to the nervous system. Nevertheless, infectious prions or PrP^{Sc} aggregates arise in the muscles of patients with variant, iatrogenic, and sporadic Creutzfeldt-Jakob disease (CJD) (12, 13), as well as in cervids with chronic wasting disease (CWD) and in sheep and hamster models of scrapie (14–16). In addition, skeletal muscles of transgenic mice that overexpress PrP^C develop myocyte degeneration and intracellular PrP aggregates (17, 18). Myocytes have also been shown to replicate infectious prions (19). Thus, skeletal muscle is a rare extracerebral tissue that develops prion-induced degeneration, aggregate formation, and prion infectivity.

The mature prion protein (approximately 209 amino acids) consists of an unstructured N terminus and a globular C-terminal domain composed of three α -helices and a short β -sheet (20). We previously demonstrated that transgenic mice expressing a “rigid-loop” structural variant of PrP (RL-PrP) due to two amino acid residue substitutions in the β 2- α 2 loop region (S170N, N174T) develop a transmissible prion disease characterized by hind limb

paresis, muscle atrophy, and kyphosis. Intriguingly, PrP^{Sc} aggregates accumulated as extracellular plaques in the brain and as intracellular inclusions in muscle (21), suggesting that aggregate development or clearance mechanisms differ for neurons and myocytes.

Intracellular misfolded proteins are routed for degradation by the ubiquitin-proteasome system or by autophagy, where cargo is sequestered in autophagosomes and ultimately degraded in endolysosomal compartments (22–24). Inadequate clearance can lead to the accumulation of misfolded proteins in “aggresomes” or “inclusion bodies.” In patients with sporadic inclusion body myositis (sIBM), intracellular inclusion bodies in myocytes are composed of a mixture of misfolded, β -sheet-rich proteins (25, 26). In sIBM, endoplasmic reticulum (ER) stress and induction of the unfolded protein response (UPR) occur, as well as macroautophagy, yet cells ultimately fail to clear aggregates. Impaired autophagy due to mutations in p97/valosin-containing protein (VCP) has been proposed to cause inclusion body myopathy associated with Paget’s disease of bone and frontotemporal dementia (IBMPFD) (27, 28).

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Address correspondence to Christina J. Sigurdson, csigurdson@ucsd.edu.

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The UPR is induced when secreted or membrane proteins misfold and accumulate in the ER (29). The UPR activates a transcriptional and translational program that increases the production of ER chaperones, enhances ER-associated protein degradation, and reduces the efficiency of ribosomal assembly on mRNAs, thereby attenuating protein translation (29). During a prion infection, the UPR is activated in the brain, decreasing protein translation through the phosphorylation of the α subunit of eukaryotic initiation factor 2 (eIF2 α) and leading to impaired synaptic function (30). PrP aggregates have also been shown to directly interact with and inhibit the proteasome (31). In cells expressing mutant PrP, impairment of the proteasome results in the trafficking of misfolded PrP directly to the plasma membrane (32).

The pathways of PrP aggregate clearance in the diverse cell types affected during prion disease remain unclear. To better understand how PrP aggregates are cleared from the skeletal muscle in individuals with familial or sporadic prion disease, we investigated the role of the UPR, the ubiquitin-proteasome system, and autophagy in muscles of aged transgenic mice that express RL-PrP.

MATERIALS AND METHODS

Animals. Aged *tg1020* mice (which express RL-PrP) (310 to 536 days; $n = 15$), aged *tga20* mice (which overexpress mouse PrP) (230 to 470 days; $n = 14$), and aged wild-type (WT) mice (173 to 524 days; $n = 12$) were utilized in these studies. The *tg1020* and *tga20* transgenic mice were developed using the “half-genomic” *Prnp* minigene containing the prion promoter and were on a *Prnp*^{-/-} background (21, 33). Transgene-positive mice were identified from tail or ear samples by standard PCR (21). Mice were maintained under specific-pathogen-free conditions with unlimited access to food and water. All procedures involving animals were performed in such a way as to minimize suffering and were approved by the Institutional Animal Care and Use Committee at UC San Diego. Protocols were performed in strict accordance with good animal practices, as described in the *Guide for the Care and Use of Laboratory Animals* (34).

Plasmids and constructs. Mouse *Prnp* cDNA containing the rigid-loop mutations (corresponding to S170N and N174T) were subcloned into the pcDNA3.1C vector (Invitrogen). Site-directed mutagenesis was performed to obtain WT-pcDNA3.1C and to introduce the 3F4 tag (residues 109M and 112M [human PrP numbering]) into the PrP sequence.

Cell culture, transfections, and drug treatments. C2C12 myoblasts (ATCC) were maintained in Dulbecco's modified Eagle medium (DMEM) supplemented with 20% fetal bovine serum, penicillin, and streptomycin. For transient transfections, C2C12 cells were grown in a 6-well plate and were transfected with 1 μ g plasmid by using the transfection reagent Lipofectamine 2000 (Invitrogen). Six hours posttransfection, the medium was changed to a complete medium containing dimethyl sulfoxide (DMSO) (Sigma), bafilomycin A (BafA) (160 nM; Calbiochem), or leupeptin (20 μ M; Sigma), and cells were incubated for an additional 20 h at 37°C. MG132 (50 μ M; Sigma) was added 16 h posttransfection, and incubation was performed for 6 h at 37°C.

Biochemical assays. (i) Tissue homogenization/cell lysis. Muscle tissue was homogenized (5%, wt/vol) in ice-cold lysis buffer (1% Triton X-100, 10 mM Tris-HCl [pH 7.4], 150 mM NaCl, 5 mM EDTA) using a BeadBeater tissue homogenizer. C2C12 cells were lysed in a buffer consisting of 50 mM Tris-HCl (pH 8.0), 150 mM NaCl, 1% NP-40, 0.25% sodium deoxycholate, 1 mM phenylmethylsulfonyl fluoride (PMSF), and protease inhibitor cocktail (Complete; Roche).

(ii) Western blot analysis. Fifty micrograms of total protein was resolved on 10% or 4-to-12% Bis-Tris gels and was transferred to a nitrocellulose membrane. Antibodies to Grp78/BiP (Abcam), ubiquitin (Dako), p62 (Sigma), beclin (Novus), LC3 (Novus), PrP 3F4 (Millipore), PrP POM1 (35) (a gift from Adriano Aguzzi), TDP-43 (Proteintech), transferrin receptor (Invitrogen), glyceraldehyde-3-phosphate dehydro-

genase (GAPDH) (Cell Signaling), and actin (Sigma) were used for protein detection along with the respective horseradish peroxidase (HRP)-conjugated secondary antibodies (Jackson Laboratories) and an enhanced chemiluminescence (ECL) substrate kit (Pierce). Signals were captured and quantified by using a Fujifilm LAS 4000 imager and MultiGauge software.

(iii) IP with anti-PrP antibody 15B3. Fifty milligrams of muscle was homogenized in homogenization buffer (Prionics) supplemented with protease inhibitors (PI) (1 mM PMSF and Complete) and was ultracentrifuged at 150,000 \times g for 1 h. The pellet was resuspended in immunoprecipitation (IP) buffer (Prionics) with PI. Samples were precleared with anti-IgM Dynabeads, followed by incubation with 15B3 (Prionics)-conjugated anti-IgM Dynabeads (1.5 μ g of 15B3 per tube) at 25°C for 2 h. The beads were washed three times with IP buffer, and protein was eluted by heating the beads in LDS sample buffer (Invitrogen) containing β -mercaptoethanol at 95°C for 5 min. For IP analysis of transfected C2C12 cells, the ultracentrifugation step was omitted.

(iv) PK sensitivity assay. Muscle homogenate was lysed in a Tris-based buffer (100 mM Tris-HCl [pH 7.4], 150 mM NaCl, 2% sarcosyl) for 10 min at 37°C, and aliquots containing approximately 200 μ g of protein were incubated either without proteinase K (PK) or in 1 to 10 μ g/ml PK (final PK concentration, 1, 2, 3, 4, 5, or 10 μ g/ml). The samples were incubated at 37°C for 30 min, and PK digestion was stopped by the addition of PMSF and protease inhibitors (Complete). Samples were transferred in triplicate to an enzyme-linked immunosorbent assay (ELISA) plate precoated with anti-PrP antibody POM2. PrP was detected with the anti-PrP biotinylated-POM1 antibody and streptavidin HRP-conjugated anti-mouse IgG as the secondary antibody. The signals were detected with the HRP substrate 1-Step Ultra tetramethylbenzidine (TMB) (Thermo Scientific). *tg1020*, *tga20*, and WT mice were analyzed in triplicate using 4 to 5 mice each.

(v) Deglycosylation assay. Lysates from C2C12 cells expressing WT-PrP or RL-PrP were denatured in a denaturing buffer (New England BioLabs) for 10 min at 95°C, followed by incubation with endoglycosidase H (endo H) or peptide-N-glycosidase F (PNGase F) (New England BioLabs) at 37°C for 90 min.

Histology. For PrP immunohistochemistry, paraffin-embedded tissue sections (2 μ m) were treated with 98% formic acid for 6 min, followed by treatment with proteinase K (10 μ g/ml) and autoclaving in citrate buffer (Dako). SAF84 (Cayman Chemicals) was used for labeling PrP. For beclin-1 (Abcam), LC-3 (Novus), CD68 (Novus), ubiquitin (Dako), and slow myosin heavy chain (Sigma) staining, the slides were autoclaved in Tris-EDTA (for beclin) or citrate (for LC-3, CD68, ubiquitin, and slow myosin) buffer prior to antibody incubation. Biotin-conjugated anti-mouse or anti-rabbit IgG (Jackson Laboratories) secondary antibodies were used along with peroxidase-conjugated streptavidin. Diaminobenzidine (DAB) was used for signal detection. TDP-43 (Proteintech) did not require pretreatment. For fast myosin heavy chain staining, an alkaline phosphatase-conjugated antibody (Sigma) was used without any pretreatment. The NBT (nitroblue tetrazolium)-BCIP (5-bromo-4-chloro-3-indolylphosphate) substrate was used to visualize the antibody.

Molecular biology analysis to detect ER stress. Muscle tissues were lysed, and total RNA was collected (Qiagen). cDNA was generated from total RNA using an iScript cDNA synthesis kit (Bio-Rad, Hercules, CA) and was used as the template for PCR amplification across the fragment of the *Xbp-1* cDNA bearing the intron target of IRE1. The primers used for mouse *Xbp-1* were 5'-GGCCTTGTGGTTGAGAACCAAGGAG-3' and 5'-GAATGCCAAAAGGATATCAGACTC-3'. PCR conditions were as follows: 95°C for 5 min; 95°C for 1 min; 68°C for 30 s; 72°C for 30 s; 72°C for 5 min with 35 cycles of amplification. PCR products were resolved on a 2% agarose-1 \times Tris-acetate-EDTA (TAE) gel.

For quantitative PCR, aliquots of cDNA were used as the template. The primers used included those for mouse *GAPDH* (5'-CTCAACTACA TGGTCTACATGTTCCA-3' and 5'-CCATTCTCGGCCTTGACTGT-3'), mouse *ERdj4* (5'-CAGAATTAATCCTGGCCTCC-3' and 5'-ACTAT

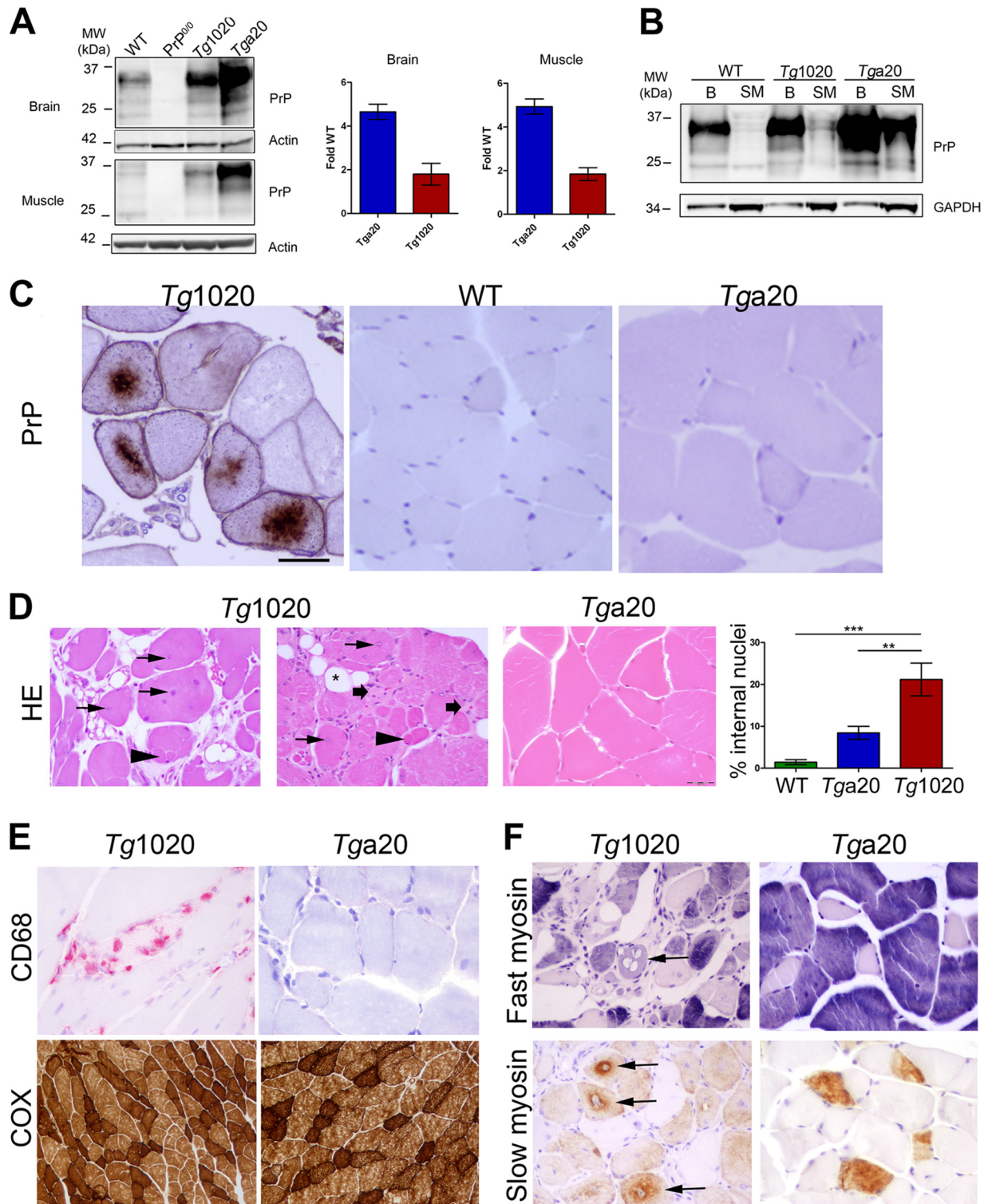


FIG 1 Aged *tg1020* mice develop a degenerative myopathy and accumulate central intracellular PrP aggregates in myocytes. (A) Western blot for PrP using equal amounts of protein from each mouse line. (B) Western blot of brain (B) and skeletal muscle (SM) shows higher PrP expression in the brain than in skeletal muscle, even with three times more protein loaded for the muscle samples (note higher GAPDH levels in muscles). POM1 was used for PrP detection. (C) Immunohistochemistry of skeletal muscle revealed coarse, dense perinuclear intracellular PrP inclusions in scattered myocytes of *tg1020* mice but not in WT or *tga20* mice. (D) Representative images of muscle show central nuclei (arrows), split fibers (arrowheads), endomysial fibrosis (broad arrows), and adipocyte replacement of myocytes (star) in *tg1020* mice. HE, hematoxylin and eosin. Asterisks in the bar graph on the right indicate significant differences in the percentages of internal nuclei among the mice by Student's *t* test (**, $P < 0.01$; ***, $P < 0.001$). (E) Focal regions of macrophage infiltration (CD68) were occasionally observed in *tg1020* muscle but not in *tga20* muscle. A cytochrome oxidase stain (COX) reveals no clear difference in mitochondrial number or pattern between *tg1020* and *tga20* muscle. (F) Slow and fast myosin heavy chain immunostains show that both type I and type II fibers are degenerating (arrows) in *tg1020* skeletal muscle.

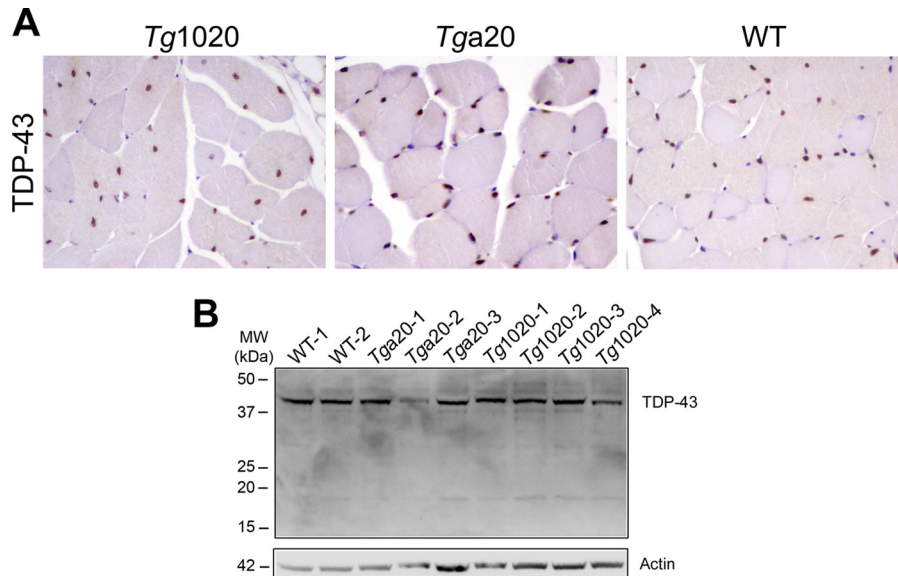


FIG 2 TDP-43 expression and localization in WT, *tg1020*, and *tga20* skeletal muscle. (A) Immunostains for TDP-43 in skeletal muscle tissue show that TDP-43 is restricted to nuclei. (B) Western blotting of skeletal muscle using an anti-TDP-43 antibody shows no marked differences in TDP-43 levels among the mice.

TGGCATCCGAGAGTG-3'), mouse *BiP* (5'-ACTCCGGCGTGAGGTA GAAA-3' and 5'-AGAGCGGAACAGGTCCATGT-3'), and mouse *Chop* (5'-ACGGAAACAGAGTGGTCAGTGC-3' and 5'-CAGGAGGTGATG CCCACTGTTC-3'). mRNA levels of *GAPDH*, a transcript whose levels are not altered by ER stress, served as an internal normalization standard. PCR conditions were as follows: 95°C for 5 min; 95°C for 10 s; 60°C for 10 s; 72°C for 10 s, with 50 cycles of amplification.

For the p62 quantitative PCR, primers and 6-carboxyfluorescein (FAM)-labeled p62 and 18S rRNA endogenous control probes (Life Technologies) were used.

RESULTS

Degenerative myopathy with PrP aggregates in *tg1020* mice. *tg1020* mice express RL-PrP^C at 2-fold WT levels, whereas *tga20* control mice express mouse PrP^C at 4- to 5-fold WT levels (Fig. 1A). The skeletal muscle expresses approximately 1/10 the PrP^C level of the brain in WT, *tg1020*, and *tga20* mice (Fig. 1B). Intracellular PrP aggregates developed in skeletal muscle from aged *tg1020* mice (more than 300 days old), but not in that from aged WT or *tga20* mice (Fig. 1C). PrP aggregates surrounded a central nucleus and occupied 10 to 40% of the sarcoplasm. Affected fibers were often clustered within the same fascicle and were interspersed with unaffected myocytes.

The muscle of *tg1020* mice showed widespread pathology, including abundant internal nuclei (approximately 23% of myocytes), split fibers, fibrosis, and myocyte replacement by adipocytes, consistent with a degenerative myopathy (Fig. 1D). Small numbers of macrophages infiltrated the interstitial space between myocytes (Fig. 1E). The mitochondria of *tg1020* and *tga20* myocytes were stained using cytochrome oxidase and did not differ significantly (Fig. 1E). Quantification of type I and II fibers by slow and fast myosin heavy chain antibody staining demonstrated that these fiber types were equally affected (Fig. 1F). Type II fibers predominated in all mice, as expected for perivertebral muscles (36).

TDP-43 was not mislocalized in affected myocytes. Mislocalization and modification of the nuclear protein TDP-43 have been

reported in several neurodegenerative and neuromuscular disorders (37–39); however, TDP-43 staining was present exclusively within nuclei, and no mislocalization was evident in *tg1020* myocytes (Fig. 2A). Additionally, no prominent cleaved fragments of TDP-43 were detected by Western blot analysis (Fig. 2B). Taken together, these results suggest that TDP-43 traffics normally in RL-PrP-expressing myocytes, in agreement with a prior study that found no TDP-43 mislocalization in brain samples from deceased patients with prion disease (40).

Biochemical properties of PrP^{Sc} in muscle. To measure the PrP aggregate load in skeletal muscle, we concentrated the insoluble PrP by ultracentrifugation and then immunoprecipitated native PrP aggregates using the PrP conformation-specific antibody, 15B3 (41). Abundant PrP aggregates were detected in the skeletal muscle of *tg1020* mice, while none were found in WT mice (Fig. 3A). Low levels of PrP aggregates were detected in the skeletal muscle of *tga20* mice (Fig. 3A), yet quantification revealed that the aggregate levels in the *tg1020* mice were 2- to 4-fold higher.

We next performed denaturing immunoprecipitation (IP) of PrP from the insoluble fractions of brain and muscle samples by using an antibody against a linear epitope (PrP antibody POM2, recognizing the octapeptide repeats in PrP residues 57 to 88 [35]). Similar levels of insoluble PrP were bound by POM2 in the *tga20* and *tg1020* homogenates (Fig. 3B and C, lower panel). In comparison, native IP using PrP conformation-specific antibody 15B3 recovered no PrP or only low levels of PrP in *tga20* muscle and brain, respectively, whereas PrP levels in the brains of *tg1020* mice were 10- to 20-fold higher, indicating that *tg1020* mice harbored abundant PrP aggregates in a PrP^{Sc}-like conformation (Fig. 3B and C, upper panel).

Aggregated PrP may be sensitive or resistant to proteinase K. We next compared the proteinase K (PK) sensitivities of PrP from WT, *tga20*, and *tg1020* muscles. Samples were incubated with increasing concentrations of PK, and PK-resistant PrP was measured by ELISA. PrP in the muscle of *tg1020* mice was significantly

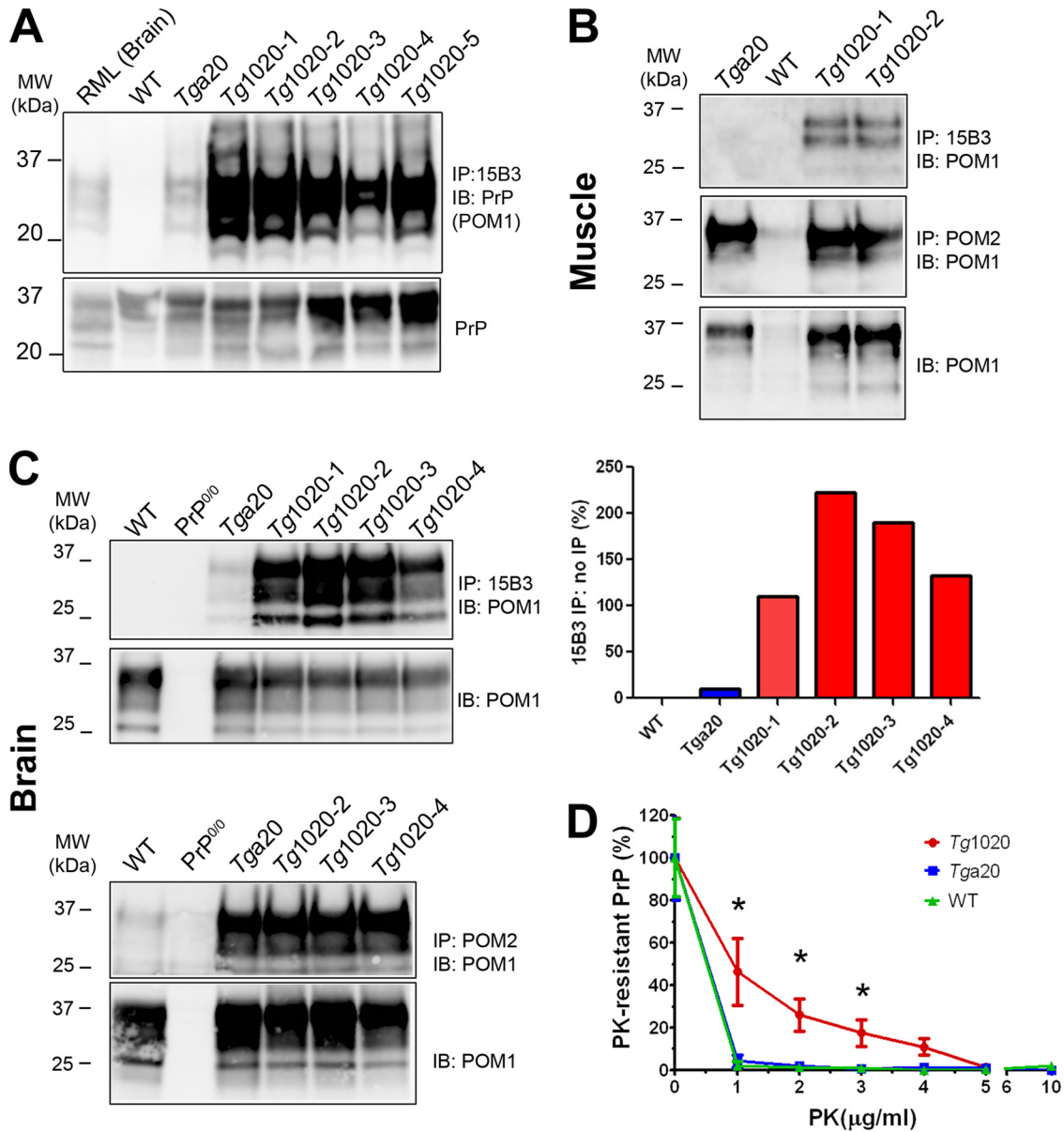


FIG 3 PrP aggregates accumulate in *tg1020* muscle. (A) (Top) Immunoprecipitation using the PrP conformation-specific antibody 15B3 reveals abundant PrP aggregates in *tg1020* skeletal muscle. (Bottom) The protein loaded for the IP contained similar amounts of PrP. Note that 4-fold-larger amounts of the WT sample were loaded. IB, immunoblot; POM1, anti-PrP antibody. (B and C) PrP in muscle and brain, respectively. Antibodies 15B3 and POM2 were used for non-denaturing and denaturing IP, respectively, of the insoluble PrP fraction obtained after ultracentrifugation. POM1 was used for detection by Western blotting. Lower panels show the PrP levels for each sample used for IP. (D) PrP in *tg1020* muscle is more PK resistant than PrP in the muscle of *tga20* or WT mice as analyzed by PrP ELISA (graphed values are normalized). Asterisks indicate significant differences (*, $P \leq 0.05$) by Student's *t* test.

more PK resistant at low PK concentrations (1 to 3 μg/ml) than PrP in the muscle of WT or *tga20* mice (Fig. 3D).

ER stress response in *tg1020* mice. Since the expression of RL-PrP may alter normal protein homeostasis, we next determined whether the UPR was induced in the muscle of *tg1020* mice. We measured levels of the ER chaperone and UPR marker Grp78/BiP (42) in the biceps femoris of aged mice. Grp78/BiP levels were significantly higher in *tg1020* mice than in WT mice (BiP/actin ratio, 9.5 ± 2 for *tg1020* mice and 2.2 ± 0.4 for WT mice [n , 10 to 16; P , 0.02 by an unpaired, two-tailed Student *t* test]) (Fig. 4A), indicating ER stress in the *tg1020* mice. The *tga20* mice showed a

slight increase in Grp78/BiP levels (BiP/actin ratio, 5.6 ± 1), which were significantly different from the levels in WT mice (P , 0.03 by Student's *t* test), suggesting that PrP overexpression causes some ER stress. Young *tg1020* mice (45 days old) showed slightly higher BiP levels than WT mice (Fig. 4B).

To assess additional UPR-related events, we searched for *Xbp-1* splicing, which is initiated by IRE1, and measured mRNA levels of *ERdj4*, an IRE1 downstream target, *BiP/Grp78*, an ATF6 downstream target, and *Chop*, a PERK downstream target, in the muscles of *tga20* and *tg1020* mice. We detected no evidence of *Xbp-1* splice events and no increase in the mRNA levels of *ERdj4* (Fig. 4C

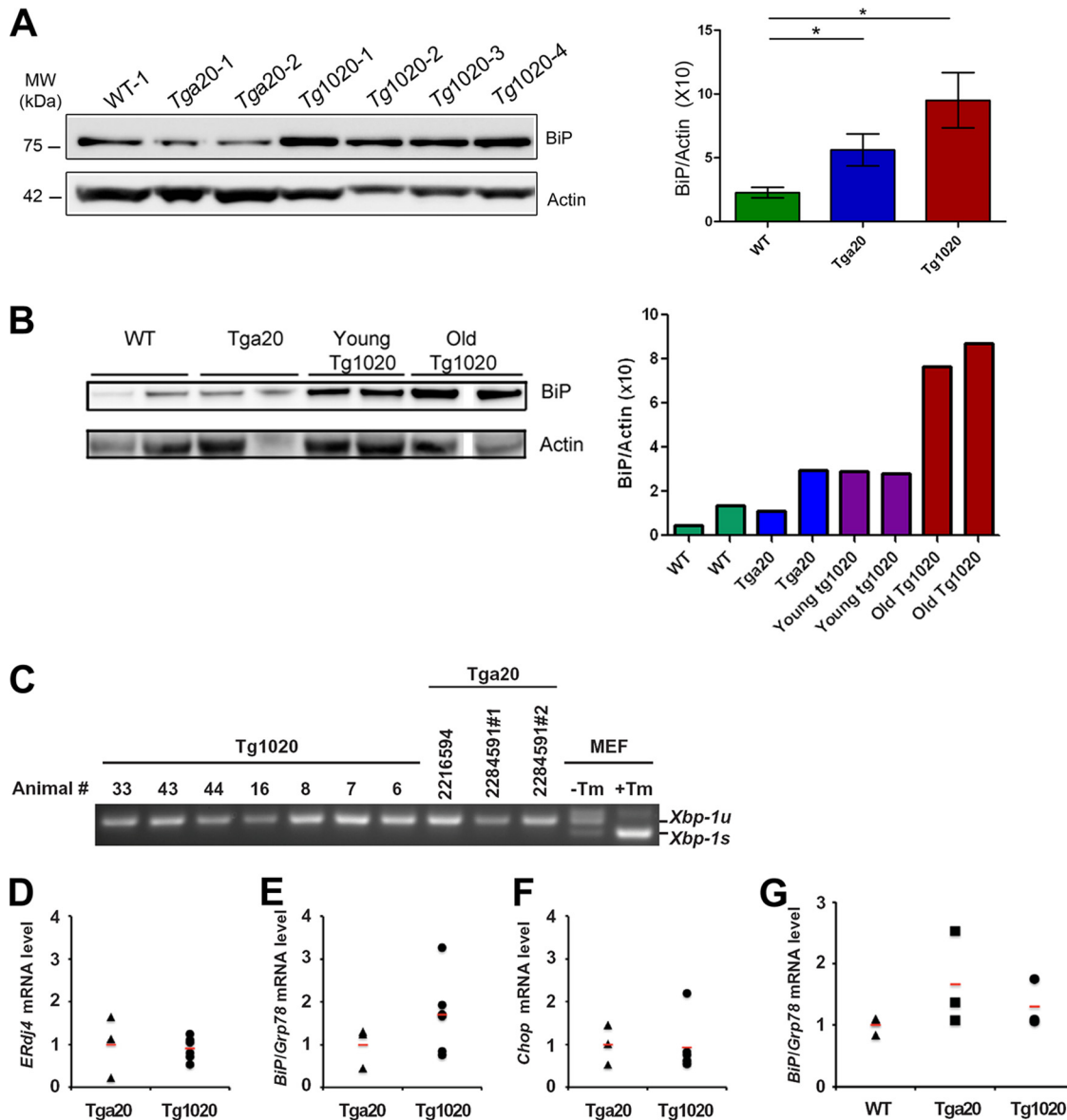


FIG 4 Induction of the unfolded protein response and an analysis of select ER stress markers in *tg1020* skeletal muscle. (A) Western blot analysis using an antibody for UPR marker Grp78/BiP shows an approximately 4-fold-higher level of Grp78/BiP in *tg1020* muscle than in WT muscle. Asterisks indicate significant differences (*, $P < 0.05$) by Student's *t* test. (B) Young *tg1020* mice had slightly higher levels of BiP than WT mice. Note: a *tg1020* sample was removed from the blot due to very low actin levels. (C) Semiquantitative real-time PCR reveals no *Xbp-1* mRNA splicing in *tg1020* skeletal muscle compared to *tga20* controls. *Xbp-1u* and *Xbp-1s* indicate the unspliced and spliced forms of *Xbp-1* mRNA, respectively. Mouse embryonic fibroblasts (MEF) treated with tunicamycin (Tm) showed strong *Xbp-1* mRNA splicing and served as a positive control. (D) Quantitative PCR analysis of mRNA levels of *ERdj4*, a downstream transcriptional target of XBP-1, showed no difference between *tga20* and *tg1020* animals. (E and F) Quantitative PCR analysis of *BiP/Grp78* and *Chop* mRNA levels, respectively, revealed higher levels of *BiP/Grp78* in the muscles of some *tg1020* mice than in *tga20* mice; however, the differences were not statistically significant. (G) Quantitative PCR analysis revealed no significant difference in *BiP/Grp78* mRNA levels between the muscles of young *tg1020* mice and those of young WT or *tga20* mice. In panels D through F, 3 *tga20* and 6 *tg1020* animals were analyzed for quantitative PCR measurements, and the values for each animal were plotted individually.

and D). Since there was variability in the levels of *BiP/Grp7* and *Chop* mRNAs among the *tg1020* mice, the differences between the levels in *tg1020* and *tga20* mice were not statistically significant. However, 3 of 6 *tg1020* mice showed elevated levels of *BiP/Grp78* mRNA, and 2 of 6 mice showed *Chop* mRNA levels more than 2-fold-higher than those of *tga20* mice (Fig. 4E and F). The induction of *BiP/Grp78* mRNA synthesis in some mice was consistent with the increase in BiP/Grp78 protein levels that was seen

(Fig. 4A and E). BiP mRNA levels were not significantly elevated in young *tg1020* mice (Fig. 4G). Together, these results are consistent with ER stress in some *tg1020* mice.

Ubiquitinated inclusions in *tg1020* muscle. Ubiquitinated inclusions occur in several protein aggregation diseases, including myopathies (43–48). Ubiquitin-positive inclusions were found histologically in the skeletal muscles of *tg1020* mice, but not in those of age-matched WT or *tga20* mice (Fig. 5A). Higher levels of

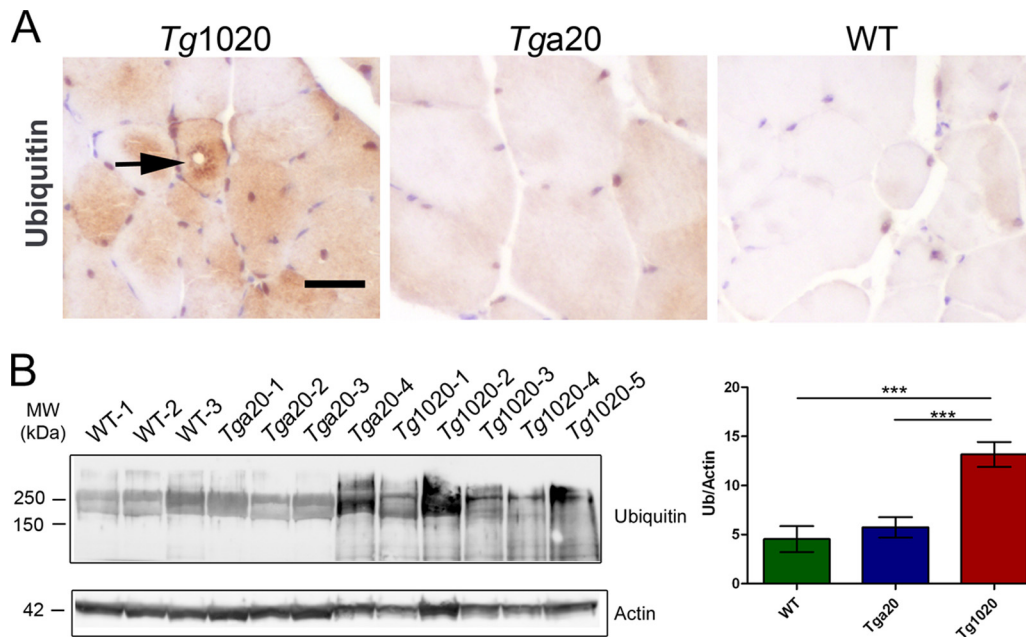


FIG 5 Accumulation of ubiquitinated proteins in skeletal muscle of *tg1020* mice (A) Immunohistochemistry shows ubiquitinated inclusions (arrow) in *tg1020* muscle. (B) Assessment of total ubiquitinated protein by Western blotting shows an approximately 3-fold-higher level of ubiquitinated proteins in *tg1020* muscle than in WT or *Tga20* muscle. Asterisks indicate significant differences (***, $P < 0.001$) by Student's *t* test.

ubiquitinated proteins were also detected biochemically in *tg1020* muscles than in *tga20* or WT muscles (Fig. 5B) (ubiquitin/actin ratios, 13.1 ± 1 for *tg1020* mice, 4.5 ± 1 for WT mice, and 5.7 ± 1 for *tga20* mice; $P < 0.001$ for *tg1020* versus WT or *tga20* mice [n , 9 to 11]).

The ubiquitinated proteins in the *tg1020* mice may consist of PrP or other proteins that were ubiquitinated due to disrupted protein homeostasis. To test for ubiquitination of PrP, we immunoprecipitated PrP aggregates using the conformation-specific antibody 15B3 under native conditions and antibody POM2 under denaturing conditions, followed by Western blot detection using the anti-ubiquitin antibody. We did not detect any ubiquitinated PrP (data not shown); however, ubiquitinated proteins can be difficult to detect. The lack of PrP ubiquitination in muscle suggests that PrP aggregates may be targeted to a nonproteasomal pathway for degradation. Alternatively, if the misfolded PrP is indeed ubiquitinated, then the turnover may be rapid and may preclude detection without the use of proteasome inhibitors *in vivo*. However, the higher levels of ubiquitinated proteins may result from a disruption of protein folding homeostasis by mutant PrP, leading to increased misfolding events and a buildup of ubiquitinated proteins, as was recently shown to occur in cells expressing a mutant huntingtin fragment (49).

Induction of autophagy in the *tg1020* muscle. Beclin-1 regulates autophagy through the formation of an autophagy initiator complex that includes Vps34 and Vps15 (50). This complex contributes to the recruitment of membranes to form autophagosomes. Beclin-1-independent pathways of autophagy may also exist (51). To determine whether beclin-1 levels were altered in *tg1020* mice, we measured beclin-1 levels in muscle homogenates by immunoblotting. Beclin-1 levels were elevated in *tg1020* skeletal muscle, suggesting that autophagy had been induced (beclin-1/actin ratios, 18.3 ± 2 for *tg1020* mice, 8.2 ± 1 for WT mice, and

7.1 ± 1 for *tga20* mice [n , 8 to 17; P , 0.01 for WT versus *tg1020* mice and 0.002 for *tga20* versus *tg1020* mice]) (Fig. 6A).

p62 levels were next assessed and were found to be elevated in *tg1020* mice, which could indicate a block in the selective degradation of ubiquitinated aggregates (52) (p62/actin ratios, 4.3 ± 0.7 for *tg1020* mice, 2.3 ± 0.3 for WT mice, and 2.6 ± 0.5 for *tga20* mice [n , 10 to 15; P , 0.04 for WT versus *tg1020* mice]) (Fig. 6B). To determine whether p62 was upregulated, we performed quantitative PCR. Here we found no differences in the p62 mRNA levels between *tg1020* mice and WT or *tga20* mice (Fig. 6C), although levels in some *tg1020* mice were 2-fold higher than those in WT mice.

As a measure of the autophagosome number (53), LC3-II protein was examined in the muscle of *tg1020* mice. LC3-II is the lipidated version of LC3-I and is involved in autophagosome formation (54). Seven of 11 (64%) *tg1020* muscle samples showed conversion of LC3-I to LC3-II (Fig. 6A and data not shown), indicating either enhanced autophagosome synthesis or a block in LC3-II degradation (53).

Consistent with the biochemical findings, immunolabeling of skeletal muscle revealed dense foci of beclin-1 and LC3 in *tg1020* mice (Fig. 6D). Taken together, these data suggest that autophagy is induced in response to PrP aggregates in the skeletal muscle.

PrP aggregates accumulate upon inhibition of autophagy and proteasomal degradation pathways *in vitro*. To investigate the clearance pathways of PrP *in vitro*, we expressed WT-PrP and RL-PrP with the 3F4 epitope tag, which is recognized by the 3F4 antibody (3F4 residues 109 to 112 [MKHM]) in mouse myoblasts (C2C12 cells). We tested the effects of two inhibitors of autophagy on the clearance of PrP: bafilomycin A (BafA), which blocks phagosome-lysosome fusion, and leupeptin, which inhibits lysosomal proteases and prevents the degradation of cargo. WT-PrP- or RL-PrP-expressing cells incubated with BafA showed 3-fold-higher

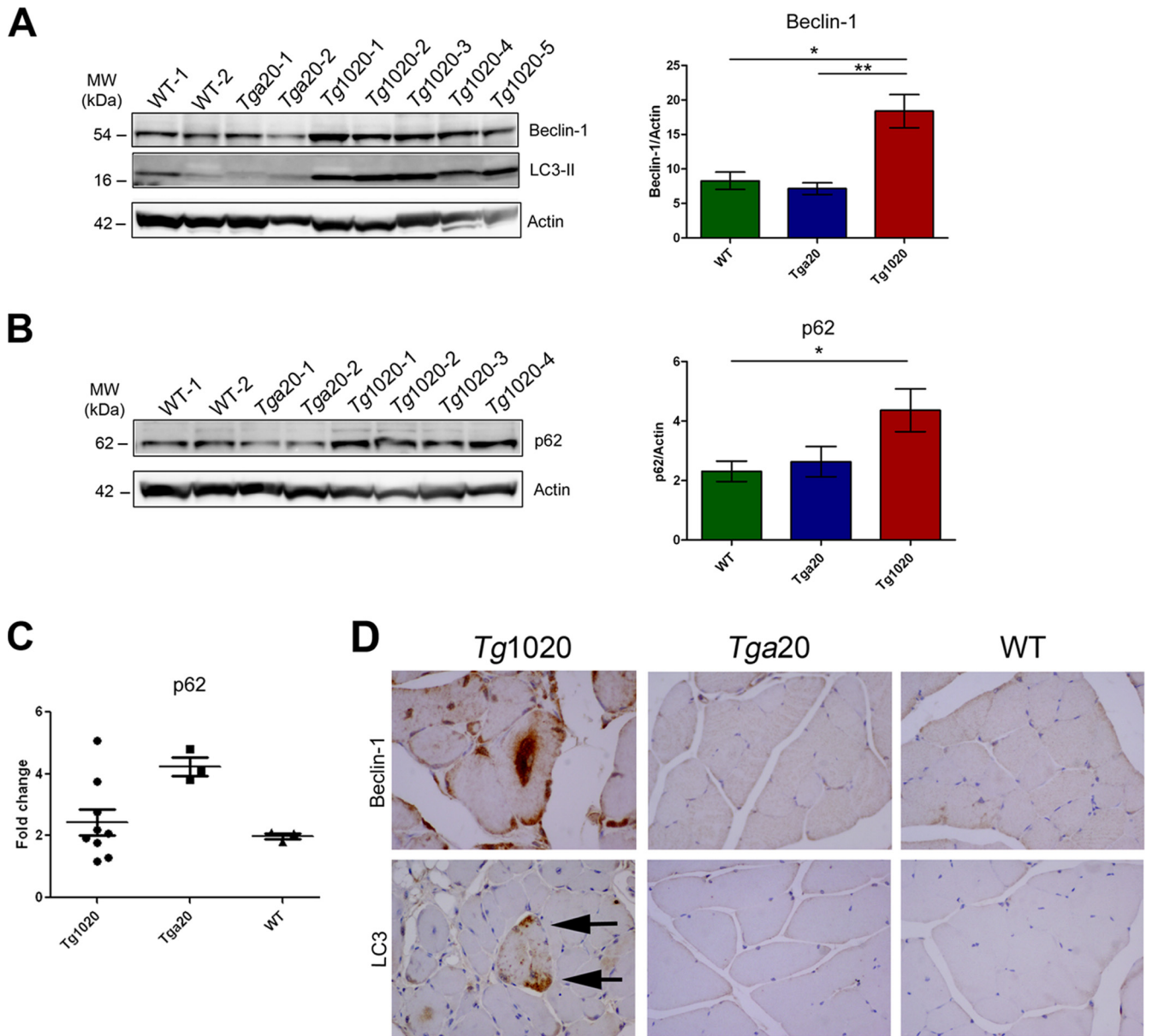


FIG 6 Increased expression of autophagy-related proteins in *tg1020* muscle. (A) Beclin-1 and LC3-II levels were higher in *tg1020* muscle than in WT or *tga20* muscle. Asterisks indicate significant differences (*, $P < 0.05$; **, $P < 0.01$) by Student's *t* test. (B) Muscle homogenates also showed increased levels of p62 in *tg1020* muscle. (C) Quantitative real-time PCR analysis of mRNA in muscles revealed that *p62* mRNA levels in *tg1020* mice do not differ from those in WT or *tga20* mice. (D) Beclin-1 immunostains revealed occasional large central aggregates in *tg1020* myocytes, whereas LC3 showed punctate peripheral staining (arrows).

levels of PrP than cells incubated with a vehicle (DMSO), whereas cells treated with leupeptin showed PrP levels 1.4- to 3-fold higher than those of vehicle-treated cells, as measured using a Fujifilm LAS 4000 imager and MultiGauge software (Fig. 7A). p62 and LC3-II levels were increased in BafA- or leupeptin-treated cells, indicating that the treatments were efficacious in inhibiting autophagy. PrP was driven by the cytomegalovirus promoter for constitutive protein expression, suggesting that the increases in PrP levels had developed due to a decrease in clearance.

To test for the formation of PrP aggregates in myoblasts, we performed IP analysis using the PrP conformational antibody 15B3. Interestingly, 15B3-reactive WT-PrP and RL-PrP accumu-

lated in BafA- and leupeptin-treated cells, but not in DMSO-treated cells (Fig. 7A, bottom). PrP overexpression may be a contributing factor underlying aggregate development in cells expressing WT-PrP, since PrP overexpression has been shown to lead to muscle and nerve degeneration (17).

We next tested the effect of the proteasomal inhibitor MG132 on WT-PrP and RL-PrP accumulation. Cells expressing WT-PrP or RL-PrP that were treated with DMSO or MG132 for 6 h showed elevated levels of ubiquitinated proteins (Fig. 7B). A previous report showed that treatment of CHO cells with proteasome inhibitors led to ubiquitination of PrP and PrP accumulation (55). Here, for both WT-PrP and RL-PrP, unglycosylated PrP levels

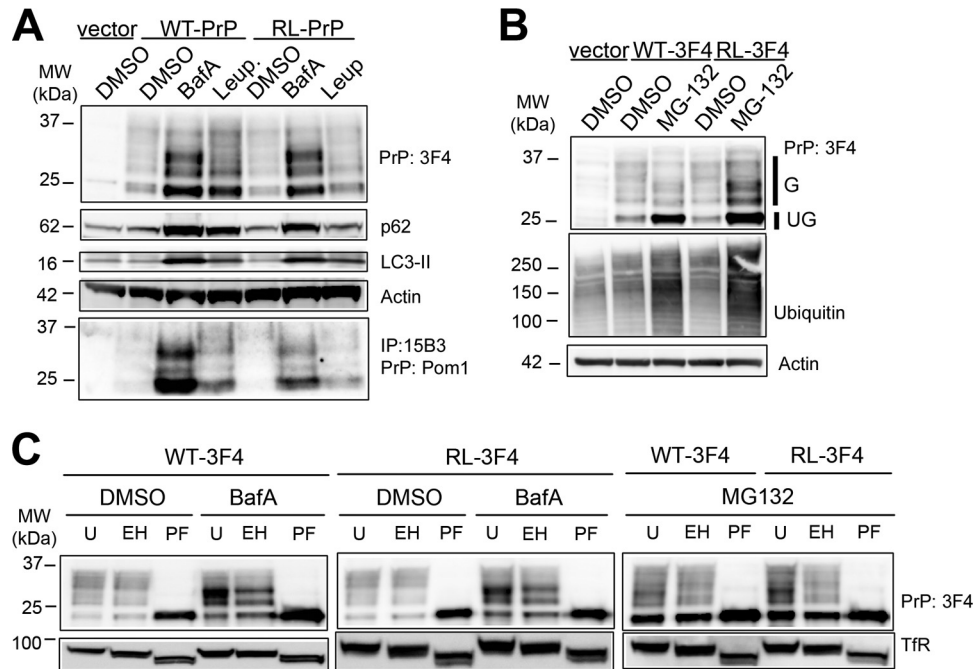


FIG 7 Inhibition of autophagy and proteasomal pathways leads to the accumulation of distinct PrP isoforms in C2C12 cells. (A) Lysates from C2C12 cells transfected either with an empty vector or with a vector encoding WT-PrP or RL-PrP show elevated levels of both unglycosylated and glycosylated PrP upon BafA or leupeptin treatment. Increased p62 and LC3-II levels show the efficacy of the BafA and leupeptin treatments in modulating autophagy. Immunoprecipitation with the PrP conformational antibody 15B3 shows that PrP aggregates accumulate in cells expressing WT-PrP and RL-PrP only upon autophagic inhibition using BafA or leupeptin. (B) Lysates from C2C12 cells show elevated levels of unglycosylated WT-PrP and RL-PrP upon MG132 treatment. Glycosylated and unglycosylated PrP signals are labeled G and UG, respectively. An increase in the level of total ubiquitinated proteins demonstrates the efficacy of MG132 treatment. Immunoblotting for actin shows that equal amounts of protein were loaded. (C) C2C12 cells were transfected with WT-PrP or RL-PrP and were treated with DMSO, BafA, or MG132. Lysates either were left untreated (U) or were treated with endo H (EH) or PNGase F (PF). Glycosylated WT-PrP and RL-PrP that accumulated upon BafA treatment were endo H resistant, yet upon MG132 treatment, they were endo H sensitive. Endo H and PNGase activities were demonstrated by shifts in the molecular weight of the transferrin receptor (TfR) corresponding to the losses of 1 and 2 glycans, respectively, compared to TfR in the untreated samples.

increased by approximately 3-fold, whereas glycosylated PrP levels increased by 2-fold, in MG132-treated cells in 3 independent experiments (Fig. 7B). Interestingly, PrP that accumulated upon proteasomal inhibition was not detected following IP with the PrP conformational antibody 15B3 in 4 independent experiments (data not shown), suggesting that PrP was monomeric or was aggregated in a conformation not recognized by the antibody.

Inhibiting the proteasome with MG132 led to relatively higher increases in levels of unglycosylated PrP than glycosylated PrP. To determine the biochemical differences between glycosylated PrP that accumulated upon autophagy and those that accumulated upon proteasome inhibition, we performed endo H and PNGase F digestion of WT-PrP and RL-PrP samples obtained after DMSO, BafA, or MG132 treatment. DMSO-treated WT-PrP and RL-PrP were primarily endo H resistant (83%) and PNGase F sensitive (Fig. 7C). After BafA treatment to inhibit autophagy, WT-PrP and RL-PrP were only slightly less endo H resistant (70 to 73%), indicating that the glycosylated species were indeed present within the Golgi apparatus or in a post-Golgi compartment. In contrast, after MG132 treatment to inhibit the proteasome, glycosylated WT-PrP and RL-PrP differed in that RL-PrP was more endo H sensitive than WT-PrP (73% of WT-PrP and 52% of RL-PrP was resistant), indicating that the accumulated RL-PrP was likely in part ER derived (Fig. 7C).

DISCUSSION

Here we report that expression of mutant PrP *in vivo* recapitulates prion aggregation and features of degenerative myopathy in a patient with spontaneous prion disease (13). Notably, the myopathy in mice is characterized by internal nuclei, split fibers, central rimmed vacuoles, cell death and replacement by adipocytes, and the accumulation of intracellular aggregates. These lesions are similar to many other protein aggregate myopathies in patients and in mouse models, including gelsolin, amyloid β , and phosphorylated tau aggregate myopathies (46, 57–59). PrP inclusions were in type I and type II fibers, suggesting that aggregation was influenced neither by the oxidative capacity nor by the glycolytic ability of the cell. In contrast to sIBM, the prion protein myopathy was not an inflammatory disease, since the inflammatory cell influx was minimal and appeared to be limited to macrophages. Additionally, these studies offer evidence of autophagy as a prion clearance mechanism, an area which has been poorly understood in prion-infected individuals, particularly in non-central-nervous-system tissues, such as muscle.

Skeletal muscles from *tg1020* mice exhibited evidence of ER stress, reflected by an increase in the levels of ER chaperones. ER stress may be directly or indirectly activated by RL-PrP and suggests that either (i) misfolded RL-PrP directly induces ER stress in the sarcoplasmic reticulum (SR) or (ii) RL-PrP disrupts general

protein folding in the SR, leading to ER stress. Chronic stress from mutant PrP may overwhelm the capacity of the SR to properly fold proteins, leading to the initiation of proapoptotic signaling pathways and cell death (37), as noted by an increase in Chop levels. Induction of apoptosis through Chop would be expected to develop asynchronously in myocytes. Here we found that not all muscle samples showed detectable increases in Chop levels, potentially due to the heterogenous distribution of the affected fibers. Further studies would be needed to clarify the pathways of myocyte death in our model. Nevertheless, our findings were consistent with reports showing ER stress in prion-expressing cell models, brains of cattle with bovine spongiform encephalopathy, and brains of mice with experimental prion infection (30, 60–65). ER stress may indirectly enhance the formation of PrP aggregates, since induction of the UPR in persistently infected scrapie cells and in cell lines expressing mutant PrP showed increased levels of PrP^{Sc} (32, 66).

To maintain proteostasis, myocytes appear to respond to the misfolded PrP by activating autophagy, suggested by the elevated levels of beclin-1 and LC3-II in the skeletal muscle *in vivo* and in C2C12 cells. Induction of autophagy has been described in other protein-misfolding myopathies. Muscle biopsy specimens from sIBM patients show an upregulation of p62 (67). Additionally, in patients with IBMPFD, mutations in VCP/p97 protein disrupted autophagy due to a defect in autophagosome maturation and led to central inclusions in myocytes similar to those seen in *tg1020* mice (28, 68). Furthermore, mutations in α -B crystallin led to protein aggregation and cardiomyopathy that were markedly exacerbated by inhibiting autophagy through heterozygous inactivation of beclin-1 (69), underscoring the importance of autophagy in aggregate clearance. Autophagy has also been shown previously to be activated in the brains of prion-infected animals (70) and in prion-infected cultured cells (71), and pharmacologic induction of autophagy has been shown to decrease PrP^{Sc} levels *in vitro* and *in vivo* (71–73). It will be of great interest to extend these studies with measurements of autophagic flux in prion models.

Although we could not detect evidence for proteasome degradation of mutant PrP in the muscles of mice, there were increased levels of unglycosylated and glycosylated PrP in cultured cells upon proteasomal inhibition. Glycosylated RL-PrP was particularly sensitive to endo H treatment, suggesting that a small fraction of RL-PrP is retained in the ER and is degraded in a proteasome-dependent manner. We failed to immunoprecipitate this species with a PrP aggregate-specific antibody, suggesting that it was monomeric or in a conformation different from the aggregated form that accumulates upon inhibition of autophagy.

Taken together, these findings highlight a role for ER stress and autophagy in the clearance of PrP from skeletal muscle in prion-infected individuals. Our data suggest that mutant PrP induces ER stress and that a small fraction of PrP is exported from the ER for degradation by the proteasome. Misfolded PrP also enters the secretory pathway, and post-Golgi PrP becomes sequestered in autophagosomes and ultimately degraded by autophagy. Additional studies may elucidate differences in aggregate clearance mechanisms in familial and transmissible prion diseases.

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