A chemical chaperone improves muscle function in mice with a RyR1 mutation

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Mutations in the \textit{RYR1} gene cause severe myopathies. Mice with an I4895T mutation in the type 1 ryanodine receptor/Ca\(^{2+}\) release channel (RyR1) display muscle weakness and atrophy, but the underlying mechanisms are unclear. Here we show that the I4895T mutation in RyR1 decreases the amplitude of the sarcoplasmic reticulum (SR) Ca\(^{2+}\) transient, resting cytosolic Ca\(^{2+}\) levels, muscle triadin content and calsequestrin (CSQ) localization to the junctional SR, and increases endoplasmic reticulum (ER) stress/unfolded protein response (UPR) and mitochondrial ROS production. Treatment of mice carrying the I4895T mutation with a chemical chaperone, sodium 4-phenylbutyrate (4PBA), reduces ER stress/UPR and improves muscle function, but does not restore SR Ca\(^{2+}\) transients in I4895T fibres to wild type levels, suggesting that decreased SR Ca\(^{2+}\) release is not the major driver of the myopathy. These findings suggest that 4PBA, an FDA-approved drug, has potential as a therapeutic intervention for RyR1 myopathies that are associated with ER stress.

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Core myopathies are the most common form of congenital myopathies and more than half of these myopathies arise from mutations in the gene (RYR1) for the skeletal muscle Ca$^{2+}$ release channel or type 1 ryanodine receptor (RyR1). RYR1 mutations in the human population occur at frequencies as high as 1 in 2000 (ref. 2). In mice some mutations in the RyR1 gene increase SR Ca$^{2+}$ leak$^1$, while others decrease Ca$^{2+}$ permeation through RyR1 (ref. 4), raising the question of how mutations with opposing effects on both the amplitude of the Ca$^{2+}$ transient and resting cytosolic Ca$^{2+}$ concentrations produce similar disease symptoms. We have shown that a mutation in RyR1 that leads to the replacement of tyrosine with a serine at amino acid 524 in RyR1 (Y522S in humans, Y524S in mice), which cause both malignant hyperthermia and myopathy with cores, increases heat sensitivity, temperature-dependent SR Ca$^{2+}$ leak, mitochondrial damage and oxidative/nitrosative stress.$^5$ Despite substantial progress towards developing interventions for myopathies arising from Ca$^{2+}$ leak in RyR1 (ref. 6), there are currently no FDA-approved drugs available for any RyR1 myopathy. The heat-induced life-threatening response in transgenic mice carrying the Y524S mutation is prevented by treatment with 5-aminoimidazole-4-carboxamide ribonucleoside$^7$ and the myopathy is alleviated by anti-oxidants.$^5$ Since, as described in this manuscript, the IT mutation decreases RyR1 Ca$^{2+}$ permeation and Ca$^{2+}$ leak, it is unlikely that interventions that further decrease Ca$^{2+}$ leak via RyR1 will be beneficial (and may even be harmful) in patients with mutations that either decrease Ca$^{2+}$ permeation through RyR1 or affect its interaction with SR-luminal proteins.

To develop interventions for RyR1 myopathies arising from RyR1 mutations that do not produce Ca$^{2+}$ leak, we created mice with the I4895T mutation (equivalent to I4898T in humans). This mutation, identified initially by Lynch et al.$^8$, is one of the common RyR1 mutations in humans and has a variable presentation and penetrance ranging from myopathies such as Central Core Disease, which can present extremely mild to severe muscle impairment in a single family$^9$–$^{11}$ to a lethal early onset congenital core-rod myopathy.$^{12}$ Mice with the I4895T mutation on a different genetic background$^{13}$ display a highly variable phenotype ranging from a severe muscle phenotype that dramatically increases in severity with age to a mild, non-progressive myopathy. The ultrastructural changes detected in these mice were found to be consistent with premature muscle ageing.$^{14}$ However, for the most part, RyR1 myopathies are early onset and non-progressive or slowly progressive$^{15}$ suggesting that the subset of I4895T mice with the severe, age-dependent phenotype are not truly representative of human early onset, slowly progressive RyR1 myopathy. In the present study we demonstrate that persistent endoplasmic reticulum (ER) stress/UPR and excessive mitochondrial reactive oxygen species (ROS) production contribute to muscle weakness and atrophy in mice with the I4895T mutation in RyR1 and that both the weakness and the atrophy are reversed by the treatment of these IT mice with a chemical chaperone, 4-phenylbutyrate (4PBA).

Results

Generation and phenotyping of IT mice. We generated mice with the I4895T mutation in RyR1 as described in Supplementary Fig. 1. These mice were backcrossed $>10$ generations onto a C57B6/J background. The I4895T mutation was homozygous lethal and, therefore, all experiments were performed with the heterozygous I4895T male mice (designated IT) and wild type (WT) littermates. Parallel studies with female mice are ongoing. When an adequate number of WT littermates were not available we used carefully aged matched WT mice from our colony on a C57B6/J background.

IT mice on a normal chow diet weighed less than their WT littermates with the greatest difference manifesting at young ages (Fig. 1a). However, after 6 weeks on a high fat diet, the IT mice gained more weight (Fig. 1b) than WT mice. IT mice ran significantly less on monitored exercise wheels (Fig. 1c) and performed poorly in wire hang tests compared to WT littermate controls (Fig. 1d). We also assessed muscle-specific (soleus, extensor digitorum longus (EDL) and diaphragm) changes in function. For clarity, we present primarily soleus data in Figs 1–8. Similar results were obtained in EDL and diaphragm muscles and these data are shown in Supplementary Figs 2 and 3, respectively.

In younger mice the force-frequency curves were shifted to the right (~9 Hz, 10-week-old mice, Fig. 1e) without a major change in maximal specific force production. In older IT mice (30 weeks of age) maximal force was decreased (see Fig. 8d). However, fatigue was not significantly altered by the IT mutation (Fig. 1f).

While the soleus from mice with the IT mutation displayed slightly more IIb/x fibres (Supplementary Fig. 4a–c), we found a significant decrease in the cross-sectional area (CSA) of all fibre types at three different ages (Fig. 1g–j). The frequency distributions of fibres with different CSAs from soleus, EDL and diaphragm muscles of IT and WT mice at different ages are shown in Supplementary Fig. 5.

Ca$^{2+}$ handling and oxidative stress. Loy et al.$^4$ found that the IT mutation reduced Ca$^{2+}$ permeation through RyR1 without changing SR luminal Ca$^{2+}$ levels. We evaluated the effects of the IT mutation on Ca$^{2+}$ handling in isolated flexor digitorum brevis (FDB) fibres using the low affinity Ca$^{2+}$ sensor, Mag-Fluo 4. We found a small reduction in the Ca$^{2+}$ transients at all stimulation frequencies in fibres from IT compared to WT littermate mice (Fig. 2a). The resting cytosolic Ca$^{2+}$ levels (measured with the high affinity Ca$^{2+}$ sensor, Fura-2) were lower in FDB fibres from IT compared to WT littermate mice (Fig. 2b), suggesting that this mutation does not lead to Ca$^{2+}$ leak. However, the relative difference in the amplitudes of the Ca$^{2+}$ transients in IT and WT fibres increased with repetitive, fatiguing stimulations (100 Hz), suggesting the possibility of activity-dependent modifications of RyR1 activity (Fig. 2c).

ER-mitochondrial-associated membranes (MaMs) are sites of Ca$^{2+}$, phospholipid and ROS exchange between the ER and mitochondria.$^{16,17}$ Similar structures designated SR mitochondrial-associated membranes (SR-MaMs) are found in skeletal muscle$^{18}$ and are thought to play a role in matching contraction-associated energy demands with ATP production via Ca$^{2+}$ signalling to mitochondria.$^{19}$ Decreased resting cytosolic Ca$^{2+}$ levels and reduced Ca$^{2+}$ transient amplitudes would be expected to decrease mitochondrial Ca$^{2+}$ uptake in IT compared to WT fibres. To determine if this is indeed the case, we attempted to assess the effects of the IT mutation on Ca$^{2+}$ uptake by mitochondria at the SR-MaMs using Rhod2 (refs 20,21). However, any changes in passive Ca$^{2+}$ uptake into mitochondria in IT and WT fibres were below the detection limits of Rhod2. Since even small increases in mitochondrial Ca$^{2+}$ uptake are associated with increased mitochondrial ROS production$^{22,23}$, we used the mitochondrial ROS sensor, MitoSOX$^{34}$; RU360, an inhibitor of the mitochondrial Ca$^{2+}$ uniporter (MCU); and Xestospongin C, an inhibitor of inositol 1,4,5-trisphosphate receptors$^{25}$ (IP3R), to determine if mitochondrial production was different in WT and IT FDB fibres. The interfibrillar mitochondria in the FDB fibres of the IT mice displayed markedly increased ROS production and this increase in ROS in the IT fibres was blocked by RU360 (Fig. 2d,e) and
Figure 1 | Phenotype of I4898T mice. (a) Body weight of IT and WT littermates on a normal chow diet as a function of age. (b) Body weight of mice on a high fat diet as a function of age. (c) Monitored running in WT and IT mice at two ages. (d) Holding impulse (Body weight (g) \times \text{maximal length of time that mice (29 weeks) hang from a wire without falling.} (e) Force frequency for the soleus muscle of 10-week-old WT and IT mice. (f) Fatigue in the soleus of 10-week-old WT and IT mice (plotted as % initial force) with 15 Hz stimulation train. (g) Representative fibre type immunostaining images in soleus muscles of IT and WT at different ages using specific myosin heavy chain antibodies. Figures shown are pseudo-coloured magenta for type I fibres, green for type IIa and black for type IIb/x. Scale bars are 100 μm. (h–j) Analysis of fibre CSA of different fibre types in the soleus of IT and WT. Distributions of CSAs for these muscles are shown in Supplementary Fig. 5. n is the number of mice used in each experiment. Data are shown as mean ± s.e.m. *P < 0.05, **P < 0.01, ***P < 0.001. For statistical analysis we used a two-tailed Student’s t-test, with differences considered significant at P < 0.05.
Xestospongin C (Fig. 2f). Since resting cytosolic Ca^{2+} levels were decreased and mitochondrial ROS production was blocked by both RU360 and Xestospongin C in IT fibres, the increased ROS production is likely to be occurring as the result of increased mitochondrial Ca^{2+} uptake at ER/SR-MaMs. The lack of an effect of RU360 and Xestospongin C in WT fibres suggests that, under normal conditions in WT fibres, Ca^{2+} release to mitochondria at the ER/SR-MaMs is minimal. The absence of a major contribution of mitochondrial Ca^{2+} uptake via the microdomain at the ER/SR MaMs in WT skeletal muscle is consistent with the finding that skeletal muscle intermembil lar mitochondria primarily take up Ca^{2+} in response to large global myoplasmic Ca^{2+} changes during repetitive stimulation. Our data suggest that persistent ER stress drives Ca^{2+} uptake and ROS production by mitochondria at the ER/SR-MaMs. IP3Rs are the Ca^{2+} release channels enriched in ER-MaMs and in other cell types and our finding of blockade of mitochondrial ROS production by Xestospongin C suggests that IP3Rs are also the Ca^{2+} release channels at the ER/SR-MaMs in skeletal muscle. The IT mutation appears to be increasing Ca^{2+} uptake, leading to increased ROS production by mitochondria at the ER/SR-MaMs despite decreases in both resting bulk cytosolic Ca^{2+} and voltage gated Ca^{2+} release. The IT mutation in RyR1 did not greatly alter the overall structure of the SR-MaMs (Fig. 2g), but the tethers between the SR and mitochondria were significantly shorter at the IT SR-MaMs (Fig. 2h,i). The decreased distance from the SR to the mitochondria could contribute to the increased mitochondrial Ca^{2+} uptake in IT muscle; however, further analyses of the structure and function of the ER/SR MaMs in IT muscle is needed.

We also assessed the expression levels of Ca^{2+} handling proteins in the IT mice. We found that the levels of Ca^{2+} handling proteins were decreased in both WT and IT soleus compared to WT control mice. The decrease was more pronounced in IT soleus, indicating that the IT mutation affects Ca^{2+} handling in skeletal muscle. This finding is consistent with the observed decrease in Ca^{2+} release and increased ROS production in IT fibres.
handling proteins (RyR1, CaV1.1, SERCA2, calsequestrin (CSQ)) in the EDL, diaphragm and soleus were, for the most part, not significantly different in muscles of WT and IT mice (Supplementary Figs 2f,g; 3f,g; 4d,e), but, as expected from the small fibre type shifts, there was a small increase in SERCA1 in the diaphragm (Supplementary Fig. 3g). The IT mutation in RyR1 is located in the selectivity filter but is close to the lumenal binding sites for triadin30,31. Triadin is required to maintain CSQ at the junctional SR in close proximity to RyR1 (ref. 32) and to optimize excitation–contraction coupling30,33. We found a decrease in the amount of triadin in IT muscle (Fig. 3a,b) at three different age groups (4, 20 and 29 weeks old) and an alteration in the localization of CSQ to the junctional SR (Fig. 3c–f). Our data suggest that the IT mutation is altering the interactions of RyR1 with proteins in the SR lumen.

Consequences of persistent Ca\(^{2+}\) uptake into the mitochondria.

Persistent elevations in mitochondrial Ca\(^{2+}\) uptake and ROS production at the ER/SR-MaMs would be expected to cause mitochondrial damage. Consistent with increased mitochondrial damage, mitochondrial protein content was decreased (20-week-old mice, Fig. 4a,b; Supplementary Figs 2h,i; 3h,i) and cytosolic cytochrome c was increased in IT muscle (Fig. 4c,d), suggesting that there has been mitochondrial permeability transition pore opening (mPTP).

Excessive mitochondrial Ca\(^{2+}\) uptake at the ER-MaMs also drives the activation of proapoptotic pathways34–36. In support of this, we found: (a) increased apoptotic nuclei (Fig. 4e,f), (b) increased cleaved caspases 3, 9 and 12 (Fig. 4g,h), (c) increased caspase 3 activity (Fig. 4i) and (d) marked elevation of the tumour suppressor and proapoptotic protein, p53, compared to WT littermates (Fig. 4j,k). These findings suggest that the persistent increases in mitochondrial Ca\(^{2+}\) and ROS production cause both mitochondrial damage and elevation of proapoptotic pathways, thereby contributing to the muscle dysfunction.

The IT mutation causes ER stress. The above findings raise the question of what drives increased mitochondrial Ca\(^{2+}\) uptake/ROS production at the ER/SR-MaMs, leading to damaged mitochondria. The SR is an extensive, specialized domain of the ER with a continuous intralumenal space37. Physiological stressors38 such as oxidative stress, glucose deprivation, inflammatory cytokines, free fatty acids, and other factors that regulate ER function, can cause the accumulation of unfolded/misfolded proteins that trigger an unfolded protein response (UPR) and ER stress. ER stress/UPR activated signalling, in turn, decreases protein synthesis, upregulates molecular chaperones to facilitate protein folding, and strengthens the contact between the ER and mitochondria to alleviate ER stress, increase ATP production and promote survival39. However, persistent or
unresolved ER stress further increases mitochondrial Ca\(^{2+}\) uptake\(^{34}\) and ROS production, induces mPTP, and drives apoptosis\(^{46}\). Both ablation of triadin and a mutation in CSQ that leads to its mislocalization are associated with ER stress in cardiac muscle\(^{41,42}\). The decreased triadin, CSQ mislocalization, increased mitochondrial ROS production, and elevation of proapoptotic markers suggested that the muscle of IT might be displaying persistent ER stress/UPR and this was further explored.

We found that proteins associated with ER stress were significantly elevated in muscles (Fig. 5a–d) of IT compared to WT littermate mice at all ages tested. ER stress-mediated UPR is orchestrated by the action of three signalling proteins, IRE1\(\alpha\) (inositol requiring enzyme 1), PERK (protein kinase RNA (PKR)-like ER kinase) and ATF6 (activating transcription factor 6\(^{43}\)). Cleaved ATF6\(\alpha\) (the active form of this transcription factor) and the downstream target of ATF6, CHOP (transcriptional factor C/EBP homologous protein), were both elevated. eIF2\(\alpha\) (eukaryotic translation initiation factor 2\(^\alpha\)) is the downstream target of PERK and the ratio of p-eIF2\(\alpha\) to eIF2\(\alpha\) was increased in IT muscle. We also analysed the alternative splicing (regulated by IRE1\(\alpha\)) of XBPI (X box-binding protein 1,
is the number of mice used in each experiment. Data are shown as mean ± s.e.m. *P < 0.05, **P < 0.01, ***P < 0.001 (two-tailed Student’s t-test).

Persistent ER stress decreases protein synthesis, causes mitochondrial damage and activates proapoptotic pathways\(^4\). As discussed above, both mitochondrial damage and activation of proapoptotic pathways are occurring in IT muscle (Fig. 4).

Figure 5 | Evidence of ER stress in IT muscle. (a) Representative western blot of proteins involved in ER stress in the soleus of 4, 8 and 29-week-old IT and WT mice. (b) Analysis of ER stress markers in soleus muscle of 4-week-old IT compared to WT littermate mice. (c) Analysis of ER stress markers in soleus muscle of 8-week-old IT compared to WT littermate mice. (d) Analysis of ER stress markers in soleus muscle of 29-week-old IT compared to WT littermate mice. (e) Analysis of XBP1 splicing by qRT-PCR. (f) Representative western blots for newly synthesized puromycin labelled proteins using anti-puromycin antibody for SUnSET analysis. (g) Analysis of changes in proteins downstream of mTORC1 activation and REDD1. (h–k) Analysis of ER stress markers in soleus muscle of 29-week-old IT compared to WT littermate mice.

Mitigation of ER stress with a chemical chaperone. If ER stress is a major contributor to the myopathy associated with the IT muscle fibre size in the IT mice.

a transcriptional activator for many of the UPR target genes), and found an increase in XBP1 splicing in the soleus muscle (Fig. 5e). These findings suggest that all three arms of the ER stress/UPR response are persistently elevated in the muscle of the IT mice.

WT mice (Fig. 5f,g). Both p-S6/S6 (ribosomal protein S6) and p-4EBP/4EBP1 (eukaryotic translation initiation factor 4E-binding protein), markers of mTORC1 activation, were decreased while REDD1 (a negative regulator of mTORC1 activity) was increased in IT muscle (Fig. 5h–k). Decreased protein synthesis is, therefore, likely to contribute to decreased muscle fibre size in the IT mice.
mutation, then interventions that alleviate ER stress should normalize the altered pathways and improve muscle function. Chemical chaperones aid in protein folding and reduce protein aggregation and oxidative stress (reviewed in ref. 47). Since ER stress, mitochondrial damage and apoptosis are defining features of IT muscle pathology, we sought to determine if restoration of protein folding with a chemical chaperone, 4PBA, would reverse muscle pathology through reduction of ER stress/UPR and oxidative stress. The data shown in Figs 6 and 7 were obtained with mice treated for 4 weeks with 4PBA in the drinking water.

Figure 6 | 4PBA reverses the signalling changes in IT muscle. (a) ProteoStat aggresome staining in the soleus of IT and WT littermates treated for 2 weeks with or without 4PBA. Lower panels are enlarged images boxed in upper panels. Scale bar is 100 μm. (b) Analysis of aggresome intensity in the soleus with or without treatment of 4PBA for 2 weeks. (c) Representative western blots on ER stress markers from the soleus of WT and IT mice treated for 2 weeks with 4PBA. (d) Effects of 4PBA on ER stress markers. (e) Representative western blot of cytosolic cytochrome c in the cytosolic fraction of soleus from WT and IT mice treated with and without 4PBA for 2 weeks. (f) Analysis of cytosolic cytochrome c in muscle from IT and WT littermate mice treated with and without 4PBA. (g) Representative western blot showing the effect of 2-week treatment of 4PBA on cleaved caspases in the soleus. (h) Effects of 4PBA on cleaved caspases in the soleus. (i) Representative western blots of p53 in the soleus of IT and WT littermates without or with 4PBA treatment for 2–3 weeks. (j) The effects of 4PBA on p53 levels in soleus. (k) Representative western blot of proteins downstream of mTORC1 in the soleus of IT and WT littermates ± 4PBA treatment for 2–3 weeks. (l) The effects of 4PBA on proteins downstream of mTORC1 from > 3 pairs of IT and WT littermates. Data are shown as mean ± s.e.m. *P < 0.05, **P < 0.01, ***P < 0.001 (two-tailed Student’s t-test).
We found that protein aggregates (stained with ProteoStat) (Fig. 6a,b), ER stress markers (Fig. 6c,d), cytosolic cytochrome c (Fig. 6e,f), cleaved caspases (Fig. 6g,h), p53 (Fig. 6i,j), all elevated in IT muscle, were markedly reduced by treating the IT mice with 4PBA. Markers of mTORC1 activation, reduced by the IT mutation, were restored to WT levels by 4PBA (Fig. 6k,l). Treatment with 4PBA also reduced muscle protein ubiquitination (Supplementary Fig. 6a,b) and autophagy markers (Supplementary Fig. 6c,d). Collectively, these findings suggest that alleviating ER stress/UPR normalizes the pathways that are altered by the IT mutation.

We next addressed the critical question of whether 4PBA affects ROS production by the interribular mitochondria or restores Ca$^{2+}$ release via RyR1 (perhaps by facilitating its refolding). Treatment of mice with 4PBA reduced both ROS production in FDB fibres (Fig. 7a) and oxidized proteins in muscle homogenates of IT mice (Fig. 7b,c). However, 4PBA did not ‘fix’ the IT associated decrease in SR Ca$^{2+}$ release via RyR1 (Fig. 7d), suggesting that the 4PBA is not just functioning to refold RyR1 with the IT mutation.

4PBA improves muscle function in mice with the IT mutation. Clinically, the most important question is whether the treatment of the IT mice with 4PBA improves muscle function and increases muscle fibre size. In 10-week-old mice 4PBA significantly improved wheel running compared to WT mice (Fig. 8a). In 30-week-old mice, 4PBA treatment dramatically improved wire-hang performance in the IT mice (Fig. 8b,c), fully restoring IT function to WT levels. Maximal force generation in the soleus was decreased in muscle isolated from the 30-week-old IT mice compared to their WT littermate controls, but after 4PBA treatment, the muscle force of the IT mice significantly improved and force generation was similar to the muscle of WT mice (Fig. 8d). Serial muscle cross-sections (stained with H&E, NADH tetrazolium reductase and MHC antibodies) from the 29-week-old IT mice, their WT littermates, and both IT and WT mice treated with 4PBA for 2–3 weeks are shown in Supplementary Fig. 7. If ER stress-driven changes in protein synthesis and protein turnover underlie the decreased fibre size in the IT mice, then chronic treatment with 4PBA should restore fibre size of IT mice to a WT level. 4PBA produced an impressive increase in fibre size in all muscles tested (Fig. 8e; Supplementary Figs 2n; 3n). The effect of 4PBA on the CSA distributions in the soleus (29-week-old mice) of IT mice is shown in Supplementary Fig. 8.

We also tested the effect of 4PBA on wire hang performance in old (50 week) IT mice and again found significant improvement (Fig. 8f). 4PBA also increased fibre size in 50-week-old mice (Fig. 8g). We did not detect major ultrastructural defects in IT muscle either with or without 4PBA (Fig. 8h), but 4PBA reduced ER stress markers even in these older mice (Fig. 8i,j) and increased mitochondrial protein content (Fig. 8k,l).

### Discussion

Defining the mechanisms underlying the pathology of RyR1 myopathies is a critical first step towards developing interventions. We explored the mechanisms by which the I4895T mutation, associated with a severe myopathy in humans (I4898T), alters muscle function in mice. In marked contrast to our previous findings in mice with a RyR1 Ca$^{2+}$ leaky mutation (Y522S in humans, Y524S in mice$^5$), the IT mutation decreases voltage-gated Ca$^{2+}$ release and resting cytosolic Ca$^{2+}$ levels but increases ER stress/UPR, enhances Ca$^{2+}$ uptake/ROS production by interfibrillar mitochondria, activates proapoptotic pathways and decreases protein synthesis. Since persistent ER stress/UPR decreases protein synthesis, strengthens the contact between the ER and mitochondria, increases mitochondrial Ca$^{2+}$ uptake and ROS production$^{34}$, induces mPTP and drives apoptosis$^{40}$, our findings suggested the possibility that the IT myopathy arises from a persistent elevation in ER stress/UPR. Indeed, we found ER stress/UPR markers were persistently elevated in the muscle of IT mice at different ages.

A critical question is how the IT mutation in RyR1 drives ER stress/UPR. The IT mutation in RyR1 is in close proximity to the amino acids (D4878, D4907 and E4908) involved in the binding of triadin to RyR1 (refs. 30). We found a decrease in triadin and a mislocalization of CSQ in muscle from the IT mice. CSQ polymerization and its interactions with triadin and junctin play a critical role in its localization to the junctional SR where it is in close contact with RyR1 (refs 30–32). Both triadin deficiency and CSQ mislocalization are associated with ER stress in cardiac muscle$^{41,42}$. Hence, we propose that alterations in lumenal SR proteins (triadin and CSQ) occurring as a result of alterations in their interactions with IT mutated RyR1 drive ER stress/UPR in skeletal muscle.

Another consequence of the IT mutation was increased ROS production, which was blocked by Xestospongin C and RU360 (blockers of IP3R and the MCU, respectively), indicating that...
 increased mitochondrial Ca$^{2+}$ uptake is driving increased ROS production. Since this is occurring under conditions with reduced cytoplasmic Ca$^{2+}$ levels, the increased mitochondrial Ca$^{2+}$ uptake is likely to be occurring at the highly specialized microdomains of the ER/SR-MaMs. Consistent with this possibility unresolved ER stress has been shown to increase mitochondrial Ca$^{2+}$ uptake$^{34}$. In addition, Csordas et al.$^{48}$ found that the spacing between the ER and mitochondria modulates mitochondrial Ca$^{2+}$ uptake. Shortened distances between the ER and mitochondrial membranes are associated with increased Ca$^{2+}$ uptake by mitochondria at ER-MaMs and mPTP$^{49,50}$. The observed decrease in the length of the tethers (tightening of tethers) between the SR and interfibrillar mitochondria in IT fibres could contribute to the increased mitochondrial Ca$^{2+}$

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Figure 8 | 4PBA improves muscle CSA and function in IT mice. (a) Wheel turns (%WT) in 10–12-week-old IT and WT mice treated with 4PBA. (b) Number of falls (% before treatment) in a wire hang test (29-week-old mice). Performance is normalized to performance before receiving 4PBA. (c) Holding impulse (%before treatment, holding time × body weight) for 29-week-old WT and IT mice. (d) Force frequency for WT, IT and IT mice treated with 4PBA for 2–3 weeks (age 25–35 weeks). (e) CSA of the soleus muscle fibres from 29-week-old mice WT and IT mice ± chronic 4PBA. (f) Holding impulse of 50-week-old IT and WT mice ± 4PBA (% pretreatment). (g) Diaphragm fibres CSA in 50-week-old IT mice treated ± 4PBA for 4–6 weeks. (h) Negatively stained transmission electron microscopy images of soleus muscle sections from 50-week-old IT mice treated ± 4PBA. Scale bars are 2 µm. (i) Representative western blot of ER stress proteins in 50-week-old IT mice treated ± 4PBA. (j) Analysis of the effects of 4PBA on ER stress markers (plotted as % untreated IT). (k) Western blot of mitochondrial proteins from 50-week-old IT mice ± 4PBA treatment. (l) Relative changes in mitochondrial protein content in muscle of 50-week-old IT mice ± 4PBA treatment. For the data in panels A, B, C and F, performance was compared to performance before 4PBA and used a paired Student's t-test. For the data in panels A, B, C and F, performance was compared to performance before 4PBA and used a paired Student's t-test.
uptake/ROS production, but additional studies of the structure/number of ER-SR-MaMs and direct measurement of mitochondrial Ca$^{2+}$ uptake are required. Alterations in mitochondrial function could contribute to the metabolic changes detected in the IT mice (increased weight gain on a high fat diet).

MUSCLE ATROPHY IS A SERIOUS PROBLEM IN RYR1 MYOPATHIES$^{11,52}$. The CSA of Type I and IIa fibres in the IT mice are decreased at least in part due to decreased muscle protein synthesis. Reduced protein synthesis is likely to result from ER stress-driven reductions in mTORC1 activity$^{44}$. Increased REDD1 (secondary to elevated p53)$^{53}$, decreased phosphorylation of Akt at S473 by mTORC2, and reduced translational initiation due to the increased phosphorylated eIF2 are likely to contribute to decreased mTORC1 activity.

The finding of persistent ER stress/UPR in the muscle of the IT mice led us to test the effects of a chemical chaperone, 4PBA, for ability to improve muscle function in the IT mice. 4PBA reduces markers of ER stress/UPR, but also increases protein synthesis and muscle mitochondrial protein content and reduces oxidative stress and proapoptotic markers, suggesting that mitochondrial damage and elevation of proapoptotic pathways are secondary to the ER stress/UPR. 4PBA, however, did not restore the amplitude of SR Ca$^{2+}$ release via RyR1, suggesting that reduced Ca$^{2+}$ release is not the primary driver of the myopathy. 4PBA is an FDA-approved drug that facilitates protein folding, in turn, suppressing ER stress-mediated apoptosis by inhibiting eukaryotic initiation factor 2α (eIF2α) phosphorylation, CCAAT (highly conserved promoter region of the Grp genes)/enhancer-binding protein homologous protein (CHOP) induction, and caspase-12 activation$^{54}$. 4PBA also increases gene expression of SR Ca$^{2+}$ leaky mutations in RyR1. Drugs such as N-benzyl-p-toluene sulfonamide (BTS) was purchased from Tocris Biosciences and 4-chloro-m-cresol (4CmC) from PFALTZ & BAUER INC. The calcium dyes fura-2 AM and mag-fluo4 AM and the mitochondrial ROS sensor, Mitotracker Red were purchased from ThermoFisher. Xeometric C was obtained from Abcam. RU-360 was purchased from Calbiochem. 4-phenylbutyric acid (4PBA) and all other chemicals were purchased from Sigma-Aldrich unless otherwise stated. Acrylamide solution (30%), protease inhibitor cocktail and acrylamide solutions were also purchased from GenDEPOT. All antibodies used are listed in Supplementary Table 1.

**Fibretyping with cryosections and immunostaining.** Muscles (soleus, EDL and diaphragm) were dissected, embedded in OCT compound (Tissue-Tek Cryomold (SAKURA)), and frozen in precooled 2-methylbutane. Frozen muscles were sectioned on 9 μm thickness using a cryostat microtome (SHANDON Cryotome E, Thermo Electron Corporation). For fibre type staining, sections were rehydrated with PBS for 10 min and incubated at 4°C overnight with anti-MHCI (dilution 1:150 dilution) and anti-MHCII (dilution 1:150 dilution) antibodies (see Supplementary Table 1 for details$^{45}$). Sections were washed twice in PBS and incubated at room temperature for 30 min AlexaFluo-594-conjugated goat anti-mouse IgG<sub>2a</sub> and AlexaFluo-488-conjugated goat anti-mouse IgG<sub>1</sub> (Life Technologies) secondary antibodies diluted 1:200 in PBS. After washing with PBS, muscle sections were mounted with Fluoromount-G mounting medium (SouthernBiotech) supplemented with DAPI for nuclear staining. Imaging was performed under a fluorescence microscope (Olympus). Fibre type number was quantified and normalized by the total number of muscle fibres per field.

**Fibre cross-sectional area.** Fibre type images were imported into Photoshop CSE v. 10.0 (Adobe Systems, San Jose, CA) for analysis. After establishing the measurement scale by tracing a scale bar in each image (μm) CSA of muscle fibres was measured by tracing the external border of individual muscle fibre using the Magnetic Lasso tool$^{56}$. For accurate analysis muscle fibres exhibiting evidence of damage or processing artefacts were excluded from the analysis. Recorded values were exported into a spreadsheet program (Excel, Microsoft Inc.) for analysis and CSA values were described as μm$^2$.

**Wire hang test.** A wire hang test$^{57}$ was employed to assess whole-body muscle strength and endurance. Mice were suspended by the forelimbs at the centre of a metallic wire (1 mm in diameter, 65 cm in length), which is elevated at 40 cm above the surface level. As the primary outcome, the holding impulse (g*s) is calculated as the product of the body mass (g) and the longest suspension time (s) during the 3-min test period. These studies were blinded to testers and the mice being analysed were randomised.

**Mouse activity.** Briefly, to assess the food intake and cage activity, mice were placed in the Comprehensive Laboratory Animal Monitoring System (CLAM System, Columbus Instruments) equipped with running wheel at room temperature and monitored for up to 2 weeks before and after drug treatment$^{58}$.

**Materials.** N-benzyl-p-toluene sulfonamide (BTS) was purchased from Tocris Biosciences and 4-chloro-m-cresol (4CmC) from PFALTZ & BAUER INC. The calcium dyes fura-2 AM and mag-fluo4 AM and the mitochondrial ROS sensor, Mitotracker Red were purchased from ThermoFisher. Xeometric C was obtained from Abcam. RU-360 was purchased from Calbiochem. 4-phenylbutyric acid (4PBA) and all other chemicals were purchased from Sigma-Aldrich unless otherwise stated. Acrylamide solution (30%), protease inhibitor cocktail and acrylamide solutions were also purchased from GenDEPOT. All antibodies used are listed in Supplementary Table 1.
**Figure 9 | Model for effects of the IT mutation and 4PBA.** The RyR1 IT mutation alters the interactions of RyR1 with SR luminal proteins to drive ER stress/UPR, leading to decreased protein synthesis, increased mitochondrial Ca\(^{2+}\) uptake and ROS production (which could feed forward to further increase ER stress), mitochondrial damage and activation of proapoptotic pathways. 4PBA reverses pathological signalling pathways and improves muscle functions in IT mice.

**Muscle force frequency and fatigue.** Strips of diaphragm muscle and intact soleus and EDL muscles were removed and immersed in Kreb’s ringer solution (137 mM NaCl, 5 mM KCl, 1 mM NaH\(_2\)PO\(_4\), 24 mM NaHCO\(_3\), 5 mM Glucose, 2 mM CaCl\(_2\), 1 mM MgSO\(_4\), pH 7.4) oxygenated with a 95/5% mixture of O\(_2\)/CO\(_2\) (ref. 56). Muscles were suspended from a force transducer and anchored in a test chamber filled with Kreb’s solution. After a 20-min equilibration period, muscle optimal length (L\(_o\)) and optimal stimulation voltage was determined via single twitch force generation measurements applied using platinum electrodes attached to a Grass S48 stimulator and recorded within Chart\(_\text{\textregistered}4\) (version 5.2) software. Force frequency measurements were obtained using frequencies from 1–300 Hz at 200 ms/train followed by a fatigue protocol performed over 5 min per muscle. The specific fatigue protocol for each muscle used was for the soleus: 15 Hz, 200 ms duration, 1 s intervals and for EDL: 60 Hz, 200 ms duration, 1 s intervals. All experiments were performed at 35 °C.

**Immunohistochemistry.** Intact FDB fibres were plated on extracellular matrix and cultured overnight at 37 °C. Muscles were fixed for 10 min in PBS containing 4% formalin and 100 μM EGTA and then washed 3 × 5 min in PBS. Fibres were then permeabilized for 30 min in 1% Triton X-100 and blocked for 1 h in blocking buffer (2% goat serum, 0.1% Triton X-100 and 0.5% BSA). Antibodies to RyR1 (Thermo Scientific MAB3925, 1:300 dilution) and CQ (Thermo Scientific PA1-913, 1:400 dilution) were diluted in blocking buffer and applied overnight at 4 °C. After washing 3 × 5 min in PBS, fibres were incubated with Alexa Fluor conjugated secondary antibodies for 2 h at room temperature before being mounted using VECTASHIELD mounting media. Fibres were imaged in the Optical Imaging and Vital Microscopy Core at BCM on a Zeiss LSM 780 confocal microscope using the x60 oil objective and images were quantified in ImageJ.

**Intracellular Ca\(^{2+}\) handling.** FDB fibres were dissected and fibres were isolated by collagenase digestion\(^\text{77}\). Fibres were imaged within 24 h of isolation. Fibres were loaded with 5 μM Mag-Fluo 4 AM for 40 min at room temperature followed by 30 min washout with Tyrode’s solution containing: 121 mM NaCl, 5 mM KCl, 1.8 mM CaCl\(_2\), 500 μM MgCl\(_2\), 400 μM NaH\(_2\)PO\(_4\), 100 μM EDTA, 5.5 mM glucose and 24 mM NaHCO\(_3\), pH regulated with bubbling of CO\(_2\)/O\(_2\) 5:95%, supplemented with 40 μM BtS to inhibit muscle contraction. To assess the frequency dependence of Ca\(^{2+}\) release, electrical stimulation was applied using two platinum electrodes positioned longitudinally either side of the fibre at the following stimulation frequencies: 1 Hz, 20 Hz, 60 Hz, 100 Hz (250 ms train duration), separated by a 1-min rest period. Fibres were imaged using a Zeiss LSM 510 meta confocal microscope in line scan mode with a ×20 objective (EC Plan-Neofluar). One line was acquired every 1.13 ms (3.66 μm/pixel/time). Mag-fluo 4 was excited at 488 nm and emitted light was detected with a 505 nm long-pass filter. Analyses of the area under the curve and Ca\(^{2+}\) transient kinetics were made using Sigmaplot 13.

In other experiments Mag Fluo 4 and Fura 2 loaded FDB fibres were imaged with an inverted Nikon TE-200 microscope equipped with a Lambda DG5 illumination system and a Rolera MG1-plus CCD camera (Photometrics). Mag Fluo 4 loaded FDB fibres were stimulated with 100 Hz trains (250 ms duration, every 1.5 s, 0.17 duty cycle) for 300 s and Mag Fluo 4 was excited at 500 nm and emitted light was collected continuously at >510 nm. To measure baseline cytoplasmic Ca\(^{2+}\) isolated FDB fibres were loaded with 10 μM Fura-2/AM for 1 h at room temperature followed by a 30-min washout with Tyrode’s solution. Fibres were excited at 340/380 nm and fluorescence emission was measured at 510 nm. Data were acquired using MetaFluor (Version 7.6) and analysed in Sigmaplot 13.

**Western blotting.** Briefly, muscles were lysed with bead homogenizer (Precelly 24 lysis & homogenization, Bertin technologies) in ice-cold RIPA buffer: 25 mM Tris pH 7.6, 150 mM NaCl, 1 mM Na\(_3\)VO\(_4\), 10 mM NaPyroPO\(_4\), 10 mM β-glycerophosphate, 10 mM NaF, 1 mM PMSF, 1X protease inhibitor cocktail (Santa Cruz), 1% NP40, 1% sodium deoxycholate and 0.1% SDS. After quantification by BCA protein assay (Thermo Scientific) equal amounts of muscle homogenates were resolved by SDS-PAGE, transferred to PVDF membrane (Millipore, Billerica, United States) and probed using primary antibodies from the vendors and at the dilutions shown in Supplementary Table 1 and visualized with Alexa Fluor conjugated secondary antibodies for 2 h at room temperature before being mounted using VECTASHIELD mounting media. Analyses of the area under the curve and Ca\(^{2+}\) transient kinetics were made using Sigmaplot 13.

**Transmission electron microscopy for ultrastructure.** Electron microscopy was performed by using methods modified from Boncompagni\(^\text{14}\). Muscle segments were postfixed in 1% OsO\(_4\), for 1 h at 4 °C, en block-stained in saturated aqueous uranyl acetate for 1 h and embedded in Spurr’s Low Viscosity resin. Ultrathin sections (50–60 nm) were cut with an ultramicrotome Leica U7 (Leica Microsystems, Austria) using a Diatome diamond knife (Diatome Ltd, CH-2501 Biel, Switzerland). Sections were stained in saturated aqueous uranyl acetate and lead citrate solutions. Sections were imaged at ×3500.

**TUNEL and caspase activity assay.** Diaphragm, EDL and soleus muscle tissues were lysed in lysis buffer (Triton X-100 0.1%, Tris HCl pH 8, 5 mM, EGTA 20 mM, 0.5% Triton X-100) and homogenized by sonication. Western blotting was performed using antibodies to terminal deoxynucleotidyl transferase dUTP nick-end labelling (TUNEL, clone TUNEL, cat. no. 452355, BD Biosciences, San Jose, CA) and anti-caspase-3 (V-F2, cat. no. 14980-1-AP,ℎ) as described in Materials and Methods. Blots were imaged with ImageJ. Western blotting was performed using antibodies to terminal deoxynucleotidyl transferase dUTP nick-end labelling (TUNEL, clone TUNEL, cat. no. 452355, BD Biosciences, San Jose, CA) and anti-caspase-3 (V-F2, cat. no. 14980-1-AP, h) as described in Materials and Methods. Blots were imaged with ImageJ.
EDTA 20 mM) and homogenized with polytron homogenizer (Precellys 24, Bertin Technologies). Protein concentrations were measured using Lowry assay. Cell apoptosis ELISA analysis was performed according to manufacturer’s instructions after appropriate dilutions (Roche). For the TUNEL assay, frozen sections of soleus were fixed with fixation solution (4% paraformaldehyde in PBS pH 7.4) for 20 min, washed with PBS for 30 min and then incubated with permeabilization solution (0.1% Triton X-100, 0.1% sodium citrate). Fixed and permeabilized slides were treated with TUNEL reaction mixture (terminal deoxynucleotidyl transferase from calf thymus, recombinant in E. coli and nucleotide mixture – Roche). Nuclear stain DAPI was used to label the nuclei. For caspase activation assay, acetyl-Asp-

**Oxyblot for oxidized proteins.** Oxidized proteins were assessed by the OxyBlot Protein Oxidation Detection Kit (Millipore) in 10 μg of soleus, EDL and diaphragm lysates following the manufacturer’s protocol.

**Mitochondrial and cytosolic fractionation.** Fresh muscles were homogenized using a glass homogenizer with ice-cold mitochondrial isolation buffer (10 mM Tris–HCl pH 7.2, 320 mM sucrose, 1 mM ethylenediamine tetra acetic acid, 1 mM dithiothreitol, 1 mM dithiothreitol, 1 mM Na3VO4, 10 mM NaPyroPO4, 10 mM β-glycerophosphate, 10 mM NaF, 1 mM PMSF, 1X protease inhibitor cocktail (Santa Cruz), 1% NPG, 1% sodium deoxycholate, and 0.1% SDS (reagents from Sigma Aldrich) and protein concentrations of each fraction were determined by the BCA Protein Assay (Thermo Scientific), according to the manufacturer’s protocol.

**XB1P1 splicing.** Total RNA was extracted from muscle tissue using TRIzol reagent (Life Technologies), and reverse transcribed into cDNA using iScript cDNA Synthesis Kit (BIO-RAD). Quantitative real-time polymerase chain reaction (qRT-PCR) was performed with iQ SYBR Green Mix (BIO-RAD) using primer sets specific for target and reference genes (PrimerBank, MGH). The relative splicing level of XB1P1 was determined by normalizing the expression of spliced XB1P1(5′) to unspliced XB1P1 (U) mRNA. The relative transcription levels of all other genes were determined by normalizing the expression of target genes to the reference 18 s rRNA. Relative gene expression was calculated based on the reference 18 s rRNA. Relative gene expression was calculated based on the reference 18 s rRNA. Relative gene expression was calculated based on the reference 18 s rRNA.

**Mitochondrial ROS.** Mitochondrial ROS production was measured using methods modified from Cheng et al.40,41 and Aydin et al.42. Briefly, single FDB fibres were incubated at room temperature with 5 μM MitoSOX Red in Tyrodes solution for 15 min, followed by a 45-min washout in indicator-free Tyrodes solution. MitoSOX loaded fibres were imaged on a Zeiss LSM 510 meta laser scanning confocal microscope (×20 objective). After 488 nm laser excitation at 488 nm and emission was collected using a 560LP filter. XY images of baseline mitochondrial ROS fluorescence were always captured within 20 min of completion of mitochondrial ROS loading. Antimycin A (20 μM) was added to the chamber at the end of each experiment to verify the rise in mitochondrial ROS fluorescence was mitochondrial. To determine the effect of blocking mitochondrial Ca2+ uptake or IP3R mediated Ca2+ release on mitochondrial ROS production, in some experiments fibres were incubated for 1 h with 10 μM RU-360, or with 1 μM Xestospongin C for 20 min before mitochondrial ROS loading.

**SunSET assay for detection of puromycin-labelled proteins.** For assessment of newly synthesized proteins we used in vivo puromycin labelling44,45. Briefly, mice (14–15 weeks of age) were deprived of food for 6 h and refed for 2 h. Propofol (18 mg/kg) was intraperitoneally injected in mice 15 min after refed. After 10 min, mice were intraperitoneally injected with puromycin (0.04 mg/kg – 2 body weight) and killed 35 min later. Excised muscles were processed for western blotting with anti-puromycin antibody (KeraFAST EQ-001, 1:1,000 dilution). Puromycin-labelled proteins were normalized to total proteins of the same blot stained with the Swift Membrane Stain kit (G-Biosciences).

**Chronic treatment with 4BPA.** Age-matched IT and WT littermates were used for chronic (up to 6 weeks) treatment. For chronic treatment, 4BPA (Sigma) was dissolved in dimethyl sulfoxide (DMSO) at a concentration of 200 mg/ml – 2 treated ad-libitum. Approximate water consumption was 3.5–3 ml per day per mouse. Therefore, each mouse consumes about 6–10 mg per day per mouse.

**Aggresome detection.** All procedures were performed as described in the manufacturer’s protocol (Enzo Life Sciences). Briefly, muscle frozen sections from animals were stained with the Swift Membrane Stain kit (G-Biosciences). Imaging was performed under ×20 magnification through an Olympus DP70 camera (Olympus America, Center Valley, PA). Fluorescence intensity of each image was quantified using Image-J software.

**Statistical analyses.** Differences between means of two groups were analysed for significance using two tailed Student’s t-tests. Data are also subjected to a normality test. If data fail the normality test they are then subjected to Mann–Whitney rank sum test. Only those data with *P*<0.05, **P*<0.01 or ***P*<0.001 are considered significant. Outliers, defined at room temperature for 10 min and rehydrated in 1X PBS. After removing excess buffer dual detection reagent (1 μl of ProteaseStat Aggresome detection reagent and 2 μl of Hoechst 33342 in 200 μl of 1X assay buffer) was added to sections and incubated for 2 h at room temperature. After washing with 1X PBS, slides were slides were mounted with coverslip using Fluoromount-G (SouthernBiotrace). Imaging was performed under ×20 magnification through an Olympus DP70 camera (Olympus America, Center Valley, PA). Fluorescence intensity of each image was quantified using Image-J software.

**Data availability.** The data that support the findings of this study are available from the corresponding author upon reasonable request.

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