Muscle stem cell adaptations to cellular and environmental stress

Maria Vittoria Gugliuzza1,2 and Colin Crist1,2*

Abstract

Background: Lifelong regeneration of the skeletal muscle is dependent on a rare population of resident skeletal muscle stem cells, also named 'satellite cells' for their anatomical position on the outside of the myofibre and underneath the basal lamina. Muscle stem cells maintain prolonged quiescence, but activate the myogenic programme and the cell cycle in response to injury to expand a population of myogenic progenitors required to regenerate muscle. The skeletal muscle does not regenerate in the absence of muscle stem cells.

Main body: The notion that lifelong regeneration of the muscle is dependent on a rare, non-redundant population of stem cells seems contradictory to accumulating evidence that muscle stem cells have activated multiple stress response pathways. For example, muscle stem cell quiescence is mediated in part by the eIF2α arm of the integrated stress response and by negative regulators of mTORC1, two translational control pathways that downregulate protein synthesis in response to stress. Muscle stem cells also activate pathways to protect against DNA damage, heat shock, and environmental stress. Here, we review accumulating evidence that muscle stem cells encounter stress during their prolonged quiescence and their activation. While stress response pathways are classically described to be bimodal whereby a threshold dictates cell survival versus cell death responses to stress, we review evidence that muscle stem cells additionally respond to stress by spontaneous activation and fusion to myofibres.

Conclusion: We propose a cellular stress test model whereby the prolonged state of quiescence and the microenvironment serve as selective pressures to maintain muscle stem cell fitness, to safeguard the lifelong regeneration of the muscle. Fit muscle stem cells that maintain robust stress responses are permitted to maintain the muscle stem cell pool. Unfit muscle stem cells are depleted from the pool first by spontaneous activation, or in the case of severe stress, by activating cell death or senescence pathways.

Keywords: Muscle stem cell, MuSC, Stress response pathways, Translational control of gene expression

Background

The skeletal muscle efficiently regenerates after acute injury in part due to a population of resident adult muscle stem cells (MuSCs), also named 'satellite cells' for their anatomical position sandwiched between the myofibre and the basal lamina [1]. Normally mitotically quiescent (G0 phase), MuSCs express members of the paired box (Pax) family of transcription factors Pax7, and in a subset of muscle, Pax3. MuSCs are primed to rapidly enter the myogenic programme, in part because they accumulate transcripts for myogenic regulatory factors myogenic factor 5 (Myf5) and myogenic determination protein (MyoD), along with cell cycle genes like Dek, which remain repressed by the action of microRNA and ribonucleic acid (RNA) binding proteins [2–6]. Some accumulating messenger RNA (mRNA), like those for Myf5, are translated inefficiently and further sequestered in cytoplasmic RNA granules [2] (Fig. 1). The formation of RNA granules in quiescent MuSCs requires

*Correspondence: colin.crist@mcgill.ca

2 Lady Davis Institute for Medical Research, Jewish General Hospital, 3755 chemin de la Côte Ste. Catherine, Montréal H3T 1E2, Canada

Full list of author information is available at the end of the article
the phosphorylation of eukaryotic initiation factor 2α (P-eIF2α) [7] (Fig. 1), a component of the integrated stress response (ISR) [8] (Fig. 2). In response to injury, the earliest stages of MuSC activation include the rapid dissociation of RNA granules and the accumulation of MYF5 and MYOD protein [2, 7, 9, 10]. Within hours after injury but prior to the first cell division, activated MuSCs also initiate a new transcriptional response to stress, rapidly upregulating core stress genes in ontologies for the stress-activated p38-mitogen-activated protein kinase (p38MAPK) and transcription in response to stress [11]. Activated MuSCs re-enter the growth 1 (G1) phase of the cell cycle, proliferate extensively to expand the population of myogenic progenitors that are required to efficiently regenerate muscle, and self-renew to replace the endogenous MuSC pool for future rounds of regeneration.

The maintenance of the MyoSC pool is critical for lifelong regeneration of the skeletal muscle, and yet, MuSCs are under pressure by, and have adapted to, numerous sources of cellular stress (Fig. 1). Not unlike yeast and all microorganisms, MuSCs spend most of their lifetime in a prolonged state of quiescence, which is also an evolutionary conserved response of cells that encounter environmental stress, poor nutrient availability, limited oxygen, and poor sources of cellular energy [12, 13]. The skeletal muscle is considered a hypoxic environment, and in particular, the MuSC niche sandwiched between the myofibre and basal lamina is hypoxic [14]. Although the majority of MuSCs reside in close proximity to the blood vessels, up to 15% of MuSCs reside away from the vessels where hypoxia may be even greater [15]. Quiescent MuSCs also have poor sources of cellular energy. They have few mitochondria and generate low amounts of cellular adenosine triphosphate (ATP) compared to activated MuSCs [16]. Quiescent MuSCs exhibit low metabolism, which may preserve MuSC regenerative potential by limiting the production of reactive oxygen species (ROS). Low metabolism is dependent on fatty acid oxidation and oxidative phosphorylation, which in turn promotes epigenetic modifications that silence the myogenic programme [17].

Upon acute injury, MuSCs break quiescence and reactivate the cell cycle. Activation is associated with a metabolic shift to anaerobic glycolysis, which is needed to support the increased cellular demands of cell growth and proliferation [17, 18]. Activated MuSCs potentially encounter proliferative stress during their critical expansion phase needed to rapidly generate a population of myogenic progenitors required for muscle regeneration. The proliferative stress encountered by MuSCs in the
normal regeneration of the muscle is illustrated by the accelerated decline in proliferation potential exhibited by myoblasts isolated from the skeletal muscle of Duchenne muscular dystrophy (DMD) patients [19]. Moreover, the critical expansion phase of MuSCs responding to muscle injury takes place within a regenerative environment characterised by pro-inflammatory cytokines with potential for cytotoxicity [20] and increased oxidative stress [21].

In this review, we highlight the consequences of a decline in MuSC fitness that is evident in muscle disease and summarise mechanisms of stress resistance and adaptation that MuSCs use to maintain the fidelity of the MuSC pool. We provide examples by which MuSCs not only survive, but possibly thrive under stressful conditions to regenerate the skeletal muscle. We propose that MuSCs have not only adapted to stress, but also thrive under stressful conditions, using basal levels of cellular and environmental stress to ensure their fitness.

**Main text**

**Maintenance of the MuSC pool is essential for regeneration of the skeletal muscle**

MuSCs are a rare population of cells in the skeletal muscle, making up less than 5% of all nuclei in the skeletal muscle. The importance of the integrity of the MuSC pool was elegantly illustrated by genetic approaches, whereby the Pax7-expressing MuSC pool was conditionally ablated by the expression of diphtheria toxin (DTA) after tamoxifen administration to $\text{Pax7}^{\text{CreERT2/}}$; $\text{R26}^{\text{DTA}}$ mice, with consequent collapse of the skeletal muscle regeneration after acute injury [22–24]. These studies
demonstrate that the skeletal muscle does not regenerate without MuSCs and also confirm that MuSCs are the only cell population with myogenic potential that regenerates muscle in vivo [25]. With only a single and rare cellular source fueling lifelong regeneration of the skeletal muscle, it is critical that this population is maintained.

**A decline in skeletal muscle regeneration occurs when the integrity of the MuSC pool is compromised**

The importance of maintaining the MuSC pool is also illustrated in muscle disease, the progression of which often coincides with a loss of MuSC numbers and function. DMD is a devastating X-linked skeletal muscle degenerative disease affecting approximately 1 in 5000 boys [26] with 100% mortality by early adulthood. The disease is caused by mutations in the *DMD* gene, which lead to impaired synthesis of full-length dystrophin protein, the absence of which causes myofibre fragility. In DMD patients, cycles of muscle degeneration and regeneration lead to the exhaustion of the MuSC pool, in part because chronically activated MuSCs exhibit severe proliferation defects and undergo premature senescence [19]. Dystrophin protein is now understood to be expressed in activated MuSCs, where it regulates MuSC senescence [19]. Dystrophin protein is now understood to be expressed in activated MuSCs, where it regulates MuSC polarity and asymmetric cell divisions that are required to maintain the MuSC pool [27, 28]. In the *Dmd*mdx mouse model of DMD, dystrophin deficiency also leads to chronic degeneration of skeletal muscle. However, the phenotype is mild and *Dmd*mdx mice have a normal lifespan, in part due to greater proliferation capacity of mouse MuSCs that fuels regeneration of the muscle [29] and likely in part due to their shorter (27 months) lifespan. In addition to DMD, mutations in *PAX7* have also been linked to the pathology of a new myopathy with variable severity in humans. The lack of *PAX7*-expressing MuSCs in the human muscle may lead in part to muscle atrophy, hypotonia, scoliosis, and mild dysmorphic facial features that are present in individuals with these mutations [30].

The progression of the muscle wasting associated with sarcopenia is also accompanied by a decrease in the fitness and numbers of MuSCs [31], with consequent loss of skeletal muscle regeneration [32, 33]. Compared to MuSCs isolated from young adult muscle, MuSCs isolated from old mice are prone to apoptosis and senescence when placed in culture [32, 34]. Aged MuSCs appear to activate a number of stress response pathways associated with p38MAPK [34–36], which is activated in response to a variety of cell stress and inflammation [37], Jak-Stat3 [38, 39], a pro-survival pathway activated in response to stress [40] and p16INK4A [32], which negatively regulates the cell cycle in response to cell stress. In young adult MuSCs, p16INK4A expression is epigenetically silenced. In geriatric mice, ubiquitination of H2A leads to permissive chromatin marks that enable p16INK4A expression. MuSCs with elevated p16INK4A expression do not activate and transit into the G1 phase, but instead irreversibly become senescent [32], with consequent depletion of the functional MuSC pool and impaired muscle regeneration.

Altogether, genetic ablation strategies and myopathies that are characterised by a loss of MuSCs number and function illustrate the importance of maintaining the MuSC pool to fuel lifelong regeneration of the muscle. The notion that the tissue microenvironment, or niche, protects MuSCs from cellular and environmental stress is challenged by evidence that quiescent MuSCs actively initiate multiple stress response pathways (Fig. 1). Next, we review the cellular responses to stress utilised by MuSCs and further discuss the fate of MuSCs when these stress response pathways are compromised.

**MuSC adaptations to cellular stress**

**MuSC responses to stress by reprogramming mRNA translation**

Regulation of mRNA translation contributes to many aspects of cell physiology, including cell growth, proliferation, differentiation, and cell survival when exposed to stress. The coordinated regulation of transcription and translation provides optimal levels of required proteins that is balanced against the energy expenditure of protein synthesis [41, 42]. Under conditions of stress, the survival of cells depends on the rapid reprogramming of translation to selectively translate mRNAs required to initiate a stress response, while globally repressing mRNA translation to reduce the energy requirements of protein synthesis [43, 44].

The arrest of translation initiation, the rate limiting step of protein synthesis, is a major hallmark of stress-induced translational control. Two translation initiation factors play central roles in the regulation of mRNA translation in response to stress. These are eukaryotic initiation factor 2 (eIF2), which is central to the ISR [45] (Fig. 2) and eukaryotic initiation factor 4E (eIF4E), which is a key component of a stress response regulated by the mechanistic target of rapamycin complex 1 (mTORC1) signalling pathway (Fig. 3) via eIF4E binding proteins (4E-BPs) [46].

**The integrated stress response**

In response to a broad range of cellular stress, eukaryotes activate the ISR [8, 45] (Fig. 2). The central event in this pathway is the phosphorylation of eukaryotic initiation factor 2α (P-eIF2α) by one of four members of the eIF2α kinase family. General control nonderepressible 2 (GCN2) responds to amino acid starvation [8], protein
kinase R (PKR) responds to the presence of viral double-stranded RNA [47], heme-regulated inhibitor (HRI) responds to the absence of heme in erythroid cells [48, 49], and PKR-like endoplasmic reticulum kinase (PERK) is activated in response to endoplasmic reticulum stress [50]. Additional environmental stresses that induce eIF2α phosphorylation for which the specific kinase remains unknown are exposure to arsenite, osmotic stress, heat shock, and nutrient starvation (Fig. 2).

The eIF2 complex (eIF2α, eIF2β, and eIF2γ) is a trimeric protein complex that is essential for protein synthesis and responsible for recycling the methionine loaded tRNA (Met-tRNA) initiation complex to the 40S ribosomal subunit to form the 43S preinitiation complex. P-eIF2α turns eIF2 into a competitive inhibitor of the guanine nucleotide exchange factor eIF2B, to prevent recycling of the eIF2-GTP-initiator methionyl tRNA ternary complex needed to initiate translation [51]. The resultant block in translation initiation has two important consequences to initiate a stress response (Fig. 2). First, translation reprogramming occurs in the cell whereby a global arrest in translation of mRNA is countered by selective translation of specific mRNAs required for the initiation of a stress response. Selective mRNA translation is mediated in part by inhibitory upstream ORFs (uORFs) in the 5'UTRs of transcripts, exemplified by transcripts for activating transcription factor 4 (Atf4) [52] (Fig. 2). P-eIF2α-dependent readthrough of inhibitory uORFs in the 5'UTR of Atf4 enables the initiation of translation at the main ORF encoding for ATF4, and ATF4 in turn activates the expression of genes required for cell recovery in response to stress [53]. Although the ISR is a pro-survival pathway, exposure to severe or prolonged stress leads to the induction of cell death pathways [54–56]. Cells that have genetic modifications to remove the phosphorylated serine residue at position...
51 of eIF2α (S51A) are unable to cope with acute stress. Moreover, the importance of eIF2α phosphorylation in mammals is illustrated by perinatal lethality in eIF2αS51A/S51A mice [57].

Second, P-eIF2α leads to a pool of mRNAs paused at the initiation step of translation, which through liquid-liquid phase separation seed the assembly of stress granules, membrane-less organelles of ribonucleoprotein complexes composed of RNA binding proteins and stalled mRNAs [48, 49] (Fig. 2). When the elf2-GTP-initiator methionyl tRNA ternary complexes are reduced, RNA binding proteins TIA1 and TIAR promote the assembly of non-canonical preinitiation complexes that lack the methionine loaded tRNA. TIA1 and TIAR dynamically triage translationally incompetent mRNAs into stress granules [58]. Despite that stress granule composition, assembly and disassembly have been studied for many years, their true function in the cell remains unclear. They presumably serve as sites of mRNA triage, help the cell cope with stress, and possibly facilitate the recovery and rapid reinitiation of translation after stress removal and stress granule disassembly [43, 59].

The phosphorylation of eIF2α is a translational control mechanism regulating MuSC quiescence and self-renewal

Quiescent MuSCs maintain low levels of protein synthesis, by PERK phosphorylation of eIF2α. The activity of PERK and P-eIF2α are both essential for MuSC quiescence and self-renewal [7]. Upon MuSC activation, eIF2α is rapidly dephosphorylated, coincident with translation and rapid accumulation of myogenic regulatory factors MYF5 and MYOD. When cultured ex vivo, rare MuSCs expressing only PAX7 maintain P-eIF2α, while the bulk of proliferating MuSCs that activate the myogenic programme dephosphorylate eIF2α. Like all cells, MuSCs require P-eIF2α to initiate a pro-survival stress response when challenged with an acute stress, for example brief exposure to ER stress inducer thapsigargin. However, MuSCs do not require P-eIF2α for cell survival under physiological conditions, nor is P-eIF2α required for MuSC survival during a regenerative response after acute injury. Instead, MuSCs that are unable to phosphorylate eIF2α are prone to spurious activation, proliferation, and contribution to new or existing myofibres in vivo [7].

P-eIF2α is also required for the assembly of RNA granules within the cytoplasm of quiescent MuSCs [7] (Fig. 1). These RNA granules are similar to size and RNA binding protein composition to stress granules, marked by RNA binding proteins DDX6, TIAR, FMRP, and GW182 [2, 3, 6]. They do not contain mRNA decapping enzyme DCP1, which is a marker of P bodies that are considered sites of mRNA decay. Instead, DCP1-positive P-bodies predominate in activated MuSCs [6]. Quiescent MuSC RNA granules are thought to be sites of storage for transcripts required for activation of the myogenic programme and proliferation. For example, Myf5 transcripts colocalize to RNA granules and can be immunoprecipitated with antibodies against DDX6 [2]. Upon MuSC activation, the dissolution of RNA granules and rapid accumulation of MYF5 protein are amongst the earliest markers of MuSC activation, which coincides with reengagement of Myf5 mRNA with translating ribosomes and rapid accumulation of MYF5 protein. Therefore, RNA granules possibly ‘prime’ quiescent MuSCs for rapid activation by their disassembly and rapid initiation of Myf5 mRNA translation [2, 7].

MuSCs appear not only to activate the P-eIF2α stress response pathway to maintain quiescence and self-renewal, but also thrive under ex vivo conditions that promote eIF2α phosphorylation [7, 60]. Under normal culture conditions, a subset of PAX7-expressing MuSCs maintain P-eIF2α, while activated MuSCs that express MYOD dephosphorylate eIF2α. Fresh isolated MuSCs that are cultured in the presence of the eIF2α phosphatase inhibitor sal003 expand ex vivo as a population of PAX7(+), MYOD(-) cells. These cells retain their stem cell properties to regenerate muscle and self-renew, illustrated by their engraftment into the Dmdmdx preclinical mouse model of Duchenne muscular dystrophy [7].

How MuSCs expand under culture conditions that promote the eIF2α phosphorylation stress response and lower global rates of protein synthesis is an interesting paradox that is potentially resolved by translational reprogramming. Culture of MuSCs in the presence of sal003 revealed hundreds of genes that are upregulated at the level of protein, without a corresponding increase in mRNA levels [61], suggesting post-transcriptional regulation. The most significantly represented class of genes were for those involved in spindle assembly, suggesting that Pax7-expressing MuSCs use eIF2α phosphorylation to maintain the fidelity of cell division. For example, P-eIF2α enables the translation of an mRNA for the mitotic spindle assembly gene transforming acidic coiled coil protein 3 (Tacc3) by virtue of inhibitory uORFs present in the 5’ untranslated region (5’UTR) of Tacc3 mRNA. In the absence of Tacc3, MuSCs expand poorly, with consequent depletion of the integrity of the MuSC pool and compromised regeneration of the skeletal muscle after acute injury [61].

There are a number of remaining questions related to the eIF2α pathway in MuSCs. What are the identity and fate of mRNAs that localise to P-eIF2α dependent RNA granules? Conversely, which mRNAs are translated in a P-eIF2α-dependent manner in quiescent MuSCs? These questions are potentially addressed with next generation RNA-seq technologies.
compatible with low amounts of mRNA isolated from ribosomes [60] or new strategies to isolate and determine the RNA component of RNA granules [61, 62]. Another important question is to what extent is P-eIF2α dependent changes in mRNA translation modified in muscle disease. While in normal healthy muscle, MuSC quiescence is maintained by PERK phosphorylation of eIF2α, the extent to which eIF2α phosphorylation is modified by kinases responding to other forms of cellular stress, for example PKR or GCN2, within the context of aging or chronically degenerating muscle, remains unclear.

**MuSC quiescence is mediated in part by the stress response pathway regulated by mTORC1**

Mechanistic target of rapamycin (mTOR) is a serine threonine kinase belonging to the family of phosphatidylinositol 3-kinase (PI3K)–related kinase (PIKKs) and is a main activator of the cellular biosynthesis machinery needed for increase cell growth and proliferation [62]. Mechanistically, mTOR functions in multiprotein complexes mTORC1 (Fig. 3) and mTORC2 and is activated by growth factors, nutrients, and energy [63]. The two most extensively studied downstream effectors of mTORC1 signalling are p70 S6 kinase (p70S6K; RPS6K1/2) and the eIF4E binding protein 1/2/3 family (4E-BP) (Fig. 3). p70S6K regulates cell growth by phosphorylation of ribosome protein S6 to increase rates of ribosome biogenesis and protein synthesis [63, 64] (Fig. 3). Phosphorylation of 4E-BPs regulates cell proliferation by disrupting their inhibition of eIF4E to enable 7-methylguanosine 5-triphosphate (m7GTP) cap-dependent translation of mRNAs encoding cell cycle regulators [46] (Fig. 3). Translational reprogramming of mRNA is also a feature of the mTORC1 pathway, since 4E-BPs regulate the translation of specific mRNAs that have established 5′ terminal oligopyrimidine (TOP) motifs [65].

The activity of mTOR is sensitive to complex signalling networks, including those that are activated in response to cell stress. The bulk of mTORC1 inhibition is channelled through the tuberous sclerosis (TSC) proteins TSC1 and TSC2 [62] (Fig. 3), which together serve to promote inactivating GTP hydrolysis of components of mTORC1. Cellular stresses that activate TSC1/TSC2 include growth factor deficiencies, low cellular energy, hypoxia, ROS, and DNA damage [62]. The resultant decrease in p70S6K activity leads to decreased ribosome biogenesis and reduced cell growth. The decrease in phosphorylated 4E-BP enables 4E-BP binding to eIF4E, leading to inhibition of cap dependent mRNA translation, and reduced cell proliferation (Fig. 3).

**Quiescent Galert MuSCs activate mTORC1 signalling to transition to Galert**

An important role for mTORC1 signalling has been elucidated in the early activation of MuSCs, termed Galert [16]. Tissue injury at distal sites leads to the accumulation and circulation of growth factors like hepatocyte growth factor (HGF) that activate mTORC1 to increase rates of protein synthesis [66]. The Galert phase is characterised by increased mitochondria, more ATP and an increase in cell size, but not by an increase in cell proliferation. Mechanistically, the Galert phenotype, characterised by an increase in MuSC size, is associated with an increase in S6 kinase phosphorylation. Moreover, genetic inactivation of Tsc1 specifically in Pax7-expressing MuSCs leads to acquisition of the Galert phenotype independent of distal injury, suggesting that MuSC quiescence is also regulated by cellular or environmental stresses that together inhibit mTORC1 signalling via TSC1 (Fig. 1). Altogether, the inhibition of mTORC1 signalling by TSC1 maintains MuSC quiescence, while the activation of mTORC1 by circulating growth factors like HGF is an early stage of MuSC activation [16, 66].

How the cell growth arm of the mTOR pathway, regulated by S6 phosphorylation, is specifically activated in Galert, while the cell proliferation arm of the mTOR pathway, potentially regulated by 4E-BP, remains resistant, is unknown. Positive mTORC1 regulation of cell proliferation potentially becomes the dominant response in activated MuSCs, since inactivation of Raptor, a specific component of the mTORC1 signalling pathway, limits MuSC proliferation, with consequent perturbation in muscle differentiation and regeneration [67]. Lastly, mTORC2 is a second mTOR complex that responds to growth factors to regulate cell proliferation, but the study of mTORC2 has lagged behind mTORC1 and has also not yet been investigated within the context of MuSC quiescence and activation.

**The DNA damage response**

When challenged with irradiation induced genotoxic stress, MuSCs resist apoptosis compared to non-myogenic cells and differentiated muscle present in the skeletal muscle. Mechanistically, quiescent MuSCs more accurately and efficiently repair DNA double-stranded breaks (DSBs) than activated MuSCs and committed progeny. Resistance to DNA damage is mediated in part due to the activity of DNA-PKcs [68] (Fig. 1), which is a central effector of the DNA damage response (DDR), a stress response pathway that senses DNA damage and replication stress to activate a protective response. DNA-PKcs is another member of the PIKK kinase family that function to phosphorylate a large number of substrates.
that are required for efficient and accurate DNA repair and also coordinate DNA repair with stalls on transcription, replication, and cell proliferation. While all quiescent MuSCs exhibit increased DNA damage repair compared to their activated and differentiated progeny, the subset of Pax3-expressing MuSCs is particularly resistant to genotoxic stress (Fig. 1). These cells have reduced levels of ROS, exhibit low levels gamma histone family member X (γH2A.X) foci, and reduced DNA damage in response to irradiation than MuSCs that only express Pax7. These cells are rare, exhibit limited contribution to normal regeneration and repair, but exhibit stress tolerance and are capable of clonal expansion and contribution to repair under stress [69].

Environmental stress
Quiescent MuSCs have developed resistance to xenobiotics, genotoxics, and oxidative stress. Toxic substances may be pumped out of the quiescent MuSC by virtue of high-level expression of genes for efflux channels Abcb1a, Abca5, and Abcc9. Moreover, quiescent MuSCs may have developed strategies to solubilise toxic substances. The aryl hydrocarbon receptor (Ahr) is also expressed at high levels in MuSCs, where it plays a role to sense toxic molecules like dioxin derivatives or polycyclic aromatic hydrocarbons [70]. Of significant interest, exposure of mice to environmental stress by the injection of the environmental pollutant 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD) leads directly to MuSC activation and fusion to myofibres in an AhR dependent manner, while only a minor fraction of MuSCs exhibit impaired survival. In contrast, the subset of Pax3-expressing MuSCs express low levels of AhR and resist TCDD exposure (Fig. 1). When adult mice are treated with TCDD for a period of 10 weeks, numbers of MuSCs expressing Pax7 only decrease, while Pax7/Pax3-expressing MuSCs remain unchanged [71].

The heat shock response
The heat shock response is a pro-survival pathway first described as a signalling response to elevations in temperature. However, many stresses activate the heat shock response, including the accumulation of protein aggregates caused by oxidative stress which in turn can be caused by exposure to heavy metals and pollutants [72–74]. To counteract the presence of protein aggregation, cells upregulate the expression of chaperone proteins that help fold nascent proteins correctly, refold misfolded proteins, and clear protein aggregation. Overall rates of transcription and mRNA translation are reduced to alleviate the burden of misfolded proteins, while protective genes are selectively expressed such as the heat shock factors (HSFs) HSF1, HSF2, and HSF4 [72]. HSFs translocate to the nucleus where they activate the expression of chaperones designated as the heat shock proteins (Hsp) such as Hsp27 and Hsp70. These HSPs promote cell survival by inhibiting apoptosis pathways and by refolding proteins [72, 73, 75].

Transcripts for heat shock proteins in the HSP40, HSP70, and HSP90 family of chaperones increase in fresh isolated, early activated MuSCs, although these transcripts are also abundant in quiescent MuSCs in vivo [76] (Fig. 1). Although it remains unclear the extent to which HSPs are involved in MuSC quiescence and activation [77], a mild heat shock to donor derived myoblasts was sufficient to induce HSP70 expression, increase cell survival after exposure to an acute stress, and improve the engraftment of these myoblasts into the Dmd<sup>mdx</sup> mouse model of DMD [78].

Perspective: cellular stress tests maintain the fitness of the MuSC pool
Environmental and cellular stress cause the cell to activate pathways that allow the cell to cope with the stress and activate an appropriate protective response. Conversely, if the stress is too severe or prolonged, stress response pathways eventually lead to senescence or the activation of cell death pathways that lead to apoptosis, autophagic cell death, and necrosis [73, 79, 80]. These cell fate decisions, cell survival if stress is resolved versus cell death if stress is too severe, are cellular responses to a variety of stress including DNA damage, heat shock, oxidative stress, and endoplasmic reticulum stress. They are considered essential to protect the tissue from accumulating damage [73]. The essence of this bimodal response to stress is there exists a threshold; the cell mounts a protective response if the stress stimulus does not go above the threshold, while severe stress leads to the activation of stress signalling cascades that fuel into cell death pathways.

A trimodal response to stress: less adaptive MuSCs spontaneously activate, differentiate, and/or fuse to the myofibre in vivo
Within the physiological context of skeletal muscle regeneration, we propose a trimodal response to stress that maintains both the integrity of the MuSC pool and the tissue from accumulating damage (Fig. 4). In addition to pro-cell survival and pro-cell death pathways described above, their exist multiple lines of evidence that MuSCs that have reduced capacity to respond to stress have eliminated themselves from the MuSC pool by spontaneous activation and/or differentiation (Fig. 4). One example is the PERK eIF2α arm of the integrated stress response that is a short-term pro-survival pathway [80, 81]. However, in MuSCs made less resistant to stress
by the genetic inactivation of Perk or eIF2α phosphorylation, cell death pathways were only activated in response to acute stress induced by thapsigargin treatment in MuSCs cultured ex vivo. Under normal physiological conditions or in response to injury, P-eIF2α defective MuSCs activated and contributed to differentiation, but they did not contribute to the MuSC pool by self-renewal, nor did they undergo apoptosis [7]. A second example is the TSC1/TSC2 inhibitor complex of mTORC1 signalling that is implicated in pro-survival pathways. Mouse embryonic fibroblasts deficient for TSC1/TSC2 exhibit increased levels of apoptosis in response to DNA damage or TNFα stimulation [82]. In contrast, MuSCs made less resistant to stress by the genetic inactivation of Tsc1 enter the Galert state, the earliest stage of their activation [16, 66]. Third, spontaneous activation and differentiation are also the mechanisms that leads to the depletion of the less stress-resistant, PAX7-only subset of MuSCs that express AhR, when challenged with the environmental stress TCDD [71], while the more stress-resistant Pax3-expressing MuSC population enter a Galert state, and may remain present to potentially restore the MuSC pool, as they have been shown to do in response to irradiation [69]. Although these potential activation/differentiation responses to decreased stress resistance are illustrated by genetic perturbations or exposure to environmental stress, it remains unclear whether spontaneous differentiation to protect the stem cell pool is a normal physiological response. Using genetic lineage marker analyses, MuSCs spontaneously activate and fuse to myofibres at unexpected rates [83, 84]. It would be of significant interest to determine whether MuSC activation and fusion to myofibres increase in response to additional stressors below the threshold that would lead to the activation of cell death pathways.

We also highlight that stimulation of stress response pathways enhances MuSC self-renewal and expansion ex vivo. This concept is illustrated by MuSCs that are cultured under low oxygen, which promotes self-renewal and delays differentiation pathways [85].
Pharmacologically maintaining the stress response pathway mediated by P-eIF2α is also effective to expand MuSCs that retain their stem cell regenerative properties ex vivo [7, 60]. Mechanistically, MuSCs may reprogramme translation to P-eIF2α dependent mRNAs, to favour self-renewal or expansion, for example by P-eIF2α-dependent translation of mRNAs like Tacc3 [61].

Conclusions
MuSCs are comfortable being uncomfortable. Cellular and environmental stress tests maintain the MuSC pool. MuSCs activate multiple stress response pathways during quiescence and activation, which seems counterintuitive to the lifelong requirement for these cells to fuel muscle regeneration. These differences may be reconciled if we repurpose the Yerkes-Dodson law, which states that there is an empirical relationship between stress and performance, to stem cell biology (Fig. 4). Quiescent MuSCs might be considered not as a population of cells that exist in comfort to protect them from stress, but rather as a specialised cell population that is continuously challenged with stress as a selective pressure to maintain their fitness, or ‘ability to serve’ in a regenerative response (Fig. 4). Unfit MuSCs may escape stress by spontaneous activation of the myogenic programme and differentiation, if damage is not severe [7, 16, 66, 71] (Fig. 4). Alternatively, if MuSCs encounter severe stress for example in aging and geriatric individuals, or within the context of proliferative stress, they are removed by activation of cell death pathways and/or irreversible senescence [19, 32] (Fig. 4).

A ‘stress test’ is a process to confirm the integrity of critical process by pushing a system to failure. Entities that fail stress tests are removed, while those that pass stress tests are selected to safeguard the system. We propose a cellular stress test for stem cell fitness, whereby quiescent and self-renewing MuSCs pass stress tests administered by their unique prolonged quiescence and microenvironment, which serve as a selective pressure to maintain MuSC fitness and safeguard the lifelong regeneration of the skeletal muscle.

Abbreviations
MuSCs: Muscle stem cells; G0: G zero phase, quiescence; Pax: Paired box; Myf5: Myogenic factor 5; MyoD: Myogenic determination protein; RNA: Ribonucleic acid; mRNA: Messenger RNA; eIF2: Eukaryotic initiation factor 2; eIF2α: Eukaryotic initiation factor 2a; ISR: Integrated stress response; p38MAPK: p38-mitogen-activated protein kinase; G0: Growth one phase; ATP: Adenosine triphosphate; DMD: Duchenne muscular dystrophy; DTA: Diptheria toxin; Dmd<sup>ts</sup>: X-chromosome linked muscular dystrophy mouse model of DMD; eIF4E: Eukaryotic initiation factor 4E, mTORC1: Mecha-nistic target of rapamycin complex 1; GCN2: General control nondepressible 2; PKB: Protein kinase B; HRI: Heme-regulated inhibitor; PERK: PKR-like endoplasmic reticulum kinase; met-tRNA: Methionine loaded tRNA; uORF: Upstream open reading frame; 5'UTR: 5' Untranslated region; Atf4: Activating transcription factor 4; Tacc3: Transforming acidic coiled coil protein 3; PI3K: Phosphatidylinositol 3-kinase; 4E-BP: eIF4E binding protein; TOP: 5' Terminal oligopyrimidine; TSC: Tuberous sclerosis; ROS: Reactive oxygen species; DNA: Deoxynucleobonic acid; HGF: Hepatocyte growth factor; DSB: Double-stranded break; DDR: DNA damage response; γH2AX: Gamma histone family member X; Ahr: Aryl hydrocarbon receptor; TCDD: 2,3,7,8-Tetra-chlorodibenzo-p-dioxin; HSF: Heat shock factor; HSP: Heat shock protein.

References
1. Mauro A. Satellite cell of skeletal muscle fibers. J Biophys Biochem Cytol. 1961;9:493–5.
2. Crist CG, Montarras D, Buckingham M. Muscle satellite cells are primed for myogenesis but maintain quiescence with sequestration of Myf5 mRNA targeted by microRNA-31 in mRNP granules. Cell Stem Cell. 2012;11(1):118–26.
3. Fujita R, Zismanov V, Jacob J-M, Jamet S, Asiev K, Crist C. Fragile X mental retardation protein regulates skeletal muscle stem cell activity by regulating the stability of Myf5 mRNA. Proc Natl Acad Sci USA. 2017;114(31):8190–5.
4. Hausburg MA, Doles JD, Clement SL, Cadwallader AB, Hall MN, Blackshear PJ, et al. Post-transcriptional regulation of satellite cell quiescence by TTP-mediated mRNA decay. Elife. 2015;4:e03390.
5. De Morree A, van Velthoven CTJ, Janq Q, Salvi JS, Klein JDD, Akimenko I, et al. Staufen1 inhibits MyoD translation to actively maintain muscle stem cell quiescence. Proc Natl Acad Sci USA. 2017;114(43):E8996–9005.
6. Roy N, Sundar S, Pillai M, Patell-Socha F, Ganesh S, Aloysius A, et al. mRNP granule proteins Fmrp and Dcx1a differentially regulate mRNP complexes to contribute to control of muscle stem cell quiescence and activation. Skeletal Muscle. 2021;11(1):1–28.
7. Zismanov V, Chichkov V, Colangelo V, Jamet S, Wang S, Syme A, et al. Phosphorylation of eIF2α is a translational control mechanism...
38. Price FD, Maltzahn von J, Bentzinger CF, Dumont NA, Yin H, Chang NC, et al. mTORC1-mediated cell proliferation, but not cell growth, controlled by the 4E-BPs. Science. 2010;328(5982):1172–6.

39. Tierney MT, Aydogdu T, Hall MJ, Kelly Tanaka K, Carter TA, Olwin BB. p38 MAPK signaling underlies a cell-autonomous loss of stem cell self-renewal in skeletal muscle of aged mice. Nat Med. 2014;20(3):265–71.

40. Troy A, Cadwallader AB, Fedorov Y, Tyner K, Tanaka KK, Olwin BB. Coordination of satellite cell activation and self-renewal by Par-complex dependent asymmetric activation of p38αβ MAPK. Cell Stem Cell. 2012;11(4):541–53.

41. Coulthurst L, White DE, Jones DI, McDermott MF, Burchill SA. p38(MAPK): stress responses from molecular mechanisms to therapeutics. Trends Mol Med. 2009;15(8):369–79.

42. Tahmasebi S, Amiri M, Sonenberg N. Principles of translational control: an overview. Cold Spring Harb Perspect Biol. 2012;4(12):1–10.

43. Hershey JWB, Sonenberg N, Mathews MB. Principles of translational control. Nature. 2015;510(7505):393–6.

44. Advani VM, Ivanov P. Translational control under stress: reshaping the translome. BioEssays. 2019;41(5):1–10.

45. Pakos-Zebrucka K, Koryga I, Mchnik K, Ljucic M, Samali A, Gorman AM. The integrated stress response. EMBO Rep. 2012;13(10):1174–95.

46. Hersey JWB, Sonenberg N, Mathews MB. Translational control. Front Genet. Frontiers. 2018;9:709.

47. Ivanov P, Kedersha N, Anderson P. Stress granules and processing bodies in translational control. Cold Spring Harb Perspect Biol. 2019;11(5):a032813.

48. Vattem KM, Wek RC. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

49. McEwen E, Kedersha N, Anderson P. Stress granules and process- ing bodies in translational control. Cold Spring Harb Perspect Biol. 2012;4(12):1–10.

50. Harding HP, Novoa I, Zhang Y, Zeng H, Wek R, Schapira M, et al. Regulated eIF2α翻译osomes in the control of translation initiation. Cell Stem Cell. 2012;10(3):255–44.

51. Bennett JO, Dodes JD, Hall MJ, Kelly Tanaka K, Carter TA, Olwin BB. p38 MAPK signaling underlies a cell-autonomous loss of stem cell self-renewal in skeletal muscle of aged mice. Nat Med. 2014;20(3):265–71.

52. Vattem KM, Wek RC. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

53. Harding HP, Novoa I, Zhang Y, Zeng H, Wek R, Schapira M, et al. Regulated eIF2α translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

54. Kazemi S, Papadopoulou S, Li S, Su Q, Wang S, Yoshimura A, et al. Control of alpha myosin of eukaryotic translation initiation factor 2 (eIF2alpha) by the human papillomavirus type 18 E6 oncoprotein. Science. 2012;334(6059):1074–5.

55. McEwen E, Kedersha N, Anderson P. Stress granules and process- ing bodies in translational control. Cold Spring Harb Perspect Biol. 2012;4(12):1–10.

56. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

57. McEwen E, Kedersha N, Anderson P. Stress granules and process- ing bodies in translational control. Cold Spring Harb Perspect Biol. 2012;4(12):1–10.

58. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

59. McEwen E, Kedersha N, Anderson P. Stress granules and process- ing bodies in translational control. Cold Spring Harb Perspect Biol. 2012;4(12):1–10.

60. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

61. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

62. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

63. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

64. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

65. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

66. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

67. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

68. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

69. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.

70. Reinitiation involving upstream ORFs regulates translation initiation controls stress-induced gene expression in mammalian cells. Proc Natl Acad Sci USA. 2005;102(17):5982–7.
implications for eIF2α-dependent gene expression and cell death. Mol Cell Biol. 2004;24(8):3415–29.
55. Fritsch RM, Schneider GN, Saur D, Scheibel M, Schmid RM. Translational repression of MCL-1 couples stress-induced eIF2α phosphorylation to mitochondrial apoptosis initiation. J Biol Chem. 2007;282(31):22551–62.
56. Jiang H-Y, Wnek RC. Phosphorylation of the α-subunit of the eukaryotic initiation factor 2 (eIF2α) reduces protein synthesis and enhances apoptosis in response to proteasome inhibition. J Biol Chem. 2005;280(14):14189–202.
57. Back SH, Scheuner D, Han J, Song B, Ribick M, Wang J, et al. Translation attenuation through eIF2α phosphorylation prevents oxidative stress and maintains the differentiated state in b cells. Cell. 2009;10(1):113–26.
58. Anderson P, Kedersha N. Visibly stressed: the role of eIF2, TIA-1, and stress granules in protein translation. Cell Stress Chaperones. 2002;7(2):213.
59. Campos-Melo D, Hawley ZCE, Droppelmann CA, Strong MJ. The integral role of RNA in stress granule formation and function. Front Cell Dev Biol. 2021;9:621779.
60. Lean G, Halloran MW, Mareshal O, Jamet S, Lumb J-P, Crist C. Ex vivo expansion of skeletal muscle stem cells with a novel small compound inhibitor of eIF2α dephosphorylation. Regen Med Front. 2019;1:11–16.
61. Fujita R, Jamet S, Lean G, Cheng HCM, Herbert S, Kleinman CL, et al. Satellite cell expansion is mediated by P-eIF2α-dependent Tacc3 translation. Development. 2021;148(2):dev194480–11.
62. Santon RA, Sabatini DM. mTOR signaling in growth, metabolism, and disease. Cell. 2017;168(6):960–76.
63. Magnuson B, Ekim B, Fingar DC. Regulation and function of ribosomal protein S6 kinase (S6K) within mTOR signaling networks. Biochem J. 2012;441(1):1–21.
64. Laplante M, Sabatini DM. mTOR signaling in growth control and disease. Cell. 2012;149(2):274–93.
65. Thoreen CC, Chantranupong L, Keys HR, Wang T, Gray NS, Sabatini DM. A unifying model for mTORC1-mediated regulation of mRNA translation. Nature. 2012;485(7396):109–13.
66. Rodgers JT, Schroeder MD, Ma C, Rando TA. HGFA is an injury-regulated systemic factor that induces the transition of stem cells into GA1ert. Cell Reports. 2017;19(9):1479–86.
67. Rion N, Castets P, Lin S, Enderle L, Reinhard JR, Eickhorst C, et al. mTOR controls embryonic and adult myogenesis via mTORC1. Development. 2019;146(7):dev172460–15.
68. Ferdousi LV, Rocheteau P, Chayot R, Montagne B, Chaker Z, Flamant R, et al. More efficient repair of DNA double-strand breaks in skeletal muscle stem cells compared to their committed progeny. Stem Cell Res. 2014;4(Suppl. 3):S9A–507.
69. Scaramozza A, Park D, Kollu S, Beerman I, Sun X, Rossi DJ, et al. Lineage tracing reveals a subset of reserve muscle stem cells capable of clonal expansion under stress. Cell Stem Cell. 2019;24(6):944–5.
70. Tierney MT, Stec MJ, Rulands S, Simons BD, Sacco A. Muscle stem cells exhibit distinct clonal dynamics in response to tissue repair and homeostatic aging. Cell Stem Cell. 2018;22(1):119–127.e3.
71. Liu W, Wen Y, Bi P, Bai X, Liu XS, Liu X, et al. Hypoxia promotes satellite cell self-renewal and enhances the efficiency of myoblast transplantaion. Development. 2012;139(16):2857–65.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.