Impact of Amyotrophic Lateral Sclerosis on Slow Tonic Myofiber Composition in Human Extraocular Muscles

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PURPOSE. To analyze the proportion and cross-sectional area of myofibers containing myosin heavy chain slow-twitch (MyHCI) and myosin heavy chain slow tonic (MyHCsto) in extraocular muscles of autopsied amyotrophic lateral sclerosis (ALS) patients with either spinal or bulbar site of disease onset.

METHODS. Whole-muscle cross sections from the middle portion of the medial rectus were labeled with antibodies against MyHCI or MyHCsto and laminin. Myofibers labeled with the MyHC antibodies (MyHCI+ and MyHCsto) and the total number of myofibers were quantified in the orbital and global layer of 6 control individuals and 18 ALS patients. The cross-sectional area of myofibers labeled for either MyHC was quantified in 130 to 472 fibers/individual in the orbital and in 180 to 573 fibers/individual in the global layer of each specimen.

RESULTS. The proportion of MyHCsto myofibers was significantly smaller in the orbital and global layer of ALS compared to control individuals. MyHCsto myofibers were significantly smaller in the global layer than in the orbital layer of ALS, whereas they were of similar size in control subjects. The decreased proportion of MyHCsto fibers correlated significantly with the age of death, but not disease duration, in patients who had the bulbar-onset variant of ALS but not in patients with spinal variant.

CONCLUSIONS. ALS, regardless of site of onset, involves a loss of myofibers containing MyHCI+st. Only in bulbar-onset cases did aging seem to play a role in the pathophysiological processes underlyng the loss of MyHCsto fibers.

Keywords: amyotrophic lateral sclerosis, extraocular muscles, MyH14, slow tonic, muscle fibers

Amyotrophic lateral sclerosis (ALS) is a progressive, neurodegenerative heterogeneous syndrome characterized by adult-onset loss of primarily motor neurons in the somatic nervous system. It leads to degeneration of upper and lower motor neurons, resulting in muscle atrophy, paresis, and loss of tendon reflexes.1 ALS usually starts in the upper or lower limbs with the symptoms outlined above, and this is termed “spinal-onset” or “classic” ALS.1,2 The disease runs a progressive course, with more and more adjacent spinal levels becoming involved. In approximately 25% of cases, the first symptoms of the disease will manifest in the head and neck region, primarily affecting muscles of articulation and mastication innervated by the motor neurons in the nuclei of the cranial nerves; this is referred to as bulbar-onset ALS.1 There is no effective treatment at present and eventually the respiratory muscles become affected and the patient dies from hypercapnia and/or pneumonia. The median survival after symptom onset is 3 to 5 years, but 5% to 10% of patients survive for 10 years or more, even without treatment, usually severely incapacitated.1

One remarkable feature of ALS already recognized by Charcot and Joffroy3 is the noticeable preservation of eye motility in many ALS patients.3 As a consequence, some patients who are otherwise paralytic can communicate with eye-controlled devices. Occasionally, disturbances in eye motility do occur, especially in patients who have survived longer with the help of invasive ventilation.4,5 But even in such cases, the disturbances usually appear to be of supranuclear origin, and complete paralysis, external ophthalmoplegia, is absent in all but a small number of reported cases.4 This is further supported by neuropathologic studies reporting that motor neuron loss in the oculomotor nuclei is infrequent.6,7 We have previously demonstrated that the extraocular muscles (EOMs) of autopsied ALS patients are remarkably well preserved in comparison to their limb muscles.8 One of the findings from that study was an apparent decrease in myofibers labeled for the myosin heavy chain (MyHC) slow tonic isoform but a preserved labeling of myofibers containing the MyHC heavy chain slow-twitch (MyHC) isoform. Because of the low number of patients available and the extensive variability in MyHC changes between individual muscles and patients, a thorough quantification of MyHC slow tonic (MyHCsto) myofibers and correlation with the clinical phenotypes of the patients were not possible at that time.8

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TABLE. Clinical Characteristics of the Studied ALS Patients

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<tr>
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Age of onset, disease duration, and age at death are reported in full lived years/months. Patients 1 and 2 were siblings. Both received intermittent noninvasive ventilation for some years. All patients were DNA screened for a panel of known ALS genes (details available upon request). C9orf72 expansion implies a heterozygous large GGGGCC-hexanucleotide expansion in intron 1 of C9orf72.

Even though there are some changes in morphology and MyHC composition in the EOMs, innervation of the neuromuscular junctions is preserved both in ALS patients and in the most commonly used transgenic mouse model of ALS, the tghSOD1D90A mouse. Actual loss of the motor neuron cell body appears to be a late phenomenon in ALS. In animal studies, the motor neuron loss is preceded by a period of progressive loss of contacts between axons and myofibers at the neuromuscular junction and retraction of peripheral axons, preceding symptom onset. Therefore, studying changes at the muscle–tissue level may reveal more subtle signs of disease involvement, as well as compensatory mechanisms that may be of relevance for disease progression. Understanding the basis for the resilient traits of the EOMs and their motor neurons may provide new clues to mechanisms that could be of importance to slow down the progression of ALS in other parts of the motor system.

In the EOMs, slow myofibers are multiply innervated and also have tonic properties, reflected in their MyHCIsto content. These multiply innervated fibers (MIFs) are present in both the orbital and global layers of the EOMs and have several neuromuscular junctions along the length of each individual muscle fiber and, in the orbital layer, more than one motor neuron innervating the same myofiber. This capacity to maintain several neuromuscular junctions on a single myofiber and share synapses between different motor neurons could perhaps make these muscle fibers less vulnerable in ALS. Furthermore, the motor neurons of EOM MIFs are located in a separate neuroanatomic niche and display a different repertoire of surface markers than other motor neurons do, suggesting that they may have distinct constitutive properties.

We investigated the effect of ALS on MIFs in the orbital and global layer of ALS patients by analyzing the proportions and sizes of myofibers containing MyHCII and MyHCsto. We also investigated whether eventual changes in the MyHC composition correlated with known clinical parameters of the patients, to investigate whether the relative sparing of EOMs was manifested differently in spinal- versus bulbar-onset ALS.

METHODS

Muscle Samples

Eighteen (n = 18) medial rectus muscles were collected post mortem from nine male and nine female terminal ALS patients (ages at death 54–80). All patients were diagnosed in accordance with the EFNS criteria for managing ALS. The clinical characteristics of the ALS patients are summarized in the Table. Six (n = 6) medial rectus EOMs, collected post mortem from one female and five males (age 42–82), served as control. None of the control subjects were known to suffer from any neurologic or muscular disease. The EOMs were collected following approval of the study by the Medical Ethical Review Board. Informed consent was obtained from the patients themselves ante mortem or following the passing of the patient, from the next of kin, in accordance with relevant Swedish legislation and in accordance with the Declaration of Helsinki.

At autopsy, after removal of any orbital fat and visible connective tissue, whole EOMs were oriented and mounted longitudinally on a piece of foil-wrapped cardboard, using OCT cryomount (HistoLab Products AB, Gothenburg, Sweden). Muscle samples were quickly frozen in liquid propane chilled with liquid nitrogen. After freezing, samples were stored at −80°C until sectioning.

The longitudinally frozen muscles were placed in a Leica CM3050 cryostat (Leica Biosystems, Nussloch, Germany) set at −23°C. With a chilled razor blade, the midportion of the muscle was cut and then mounted for cross-sectioning. Whole EOM serial cross sections were cut from the middle portion, 5 μm thick, and collected on gelatin-covered glass slides. Slides were kept at −20°C until processed for immunofluorescence.

Immunofluorescence Staining. In brief, slides were brought to room temperature and left to dry for 20 minutes. Then, slides were rinsed for 3 × 5 minutes in phosphate-buffered saline (PBS), followed by 15 minutes of incubation with donkey normal serum at room temperature. Afterward, sections were incubated with a sheep polyclonal antibody against laminin (1:15,000, PC128; The Binding Site Group Ltd., Birmingham, UK) overnight at +4°C. On the next day, the...
sections were brought to room temperature, rinsed, and incubated with normal serum (as above) and incubated with a donkey anti-sheep Alexa 647 secondary antibody (1:300, No. 713-605-147; Jackson Immunoresearch, West Grove, PA, USA) for 30 minutes at 37°C. Following this, the sections were rinsed and incubated as above with a mouse monoclonal antibody against MyHC1 (1:50, A4:951; Developmental Studies Hybridoma Bank, Iowa City, IA, USA) and a rabbit polyclonal antibody against MYH14/7b20 (here referred to as MyHCsto, 1:500, generously provided by Stefano Schiaffino, University of Padova, Padova, Italy) for 60 minutes at 37°C. Subsequently, sections were rinsed and incubated as above with a donkey anti-rabbit RRX secondary antibody (1:500, No. 711-295-152; Jackson Immunoresearch) and a donkey anti-mouse FITC secondary antibody (1:100, No. 711-095-151; Jackson Immunoresearch) for 30 minutes at 37°C. Lastly, sections were rinsed in PBS and mounted with coverslips using Vectashield Antifade mounting medium (Vector Laboratories, Inc., Burlingame, CA, USA). Secondary antibody controls were used to verify the labeling of the primary antibodies.

Microscopy. All slides were analyzed with a Leica DM 6000 B microscope (Leica Microsystems, Wetzlar, Germany) equipped with a motorized stage. A set of images (1392 × 1040 pixels) together covering the whole cross section of each specimen was captured using a ×20 objective and a digital montage was automatically assembled. The boundary between the orbital and global layers was drawn on the whole-section digital montage.

From the montage image, images were randomized and quantified until at least 1800 myofibers from the global and 1500 myofibers from the orbital layer, respectively, had been labeled with regard to MyHC composition. On each image, a counting frame for unbiased quantification of variable-sized structures21 was used.

The total number of myofibers present within the counting area was recorded, together with the number of myofibers labeled with antibodies against MyHC1, MyHCsto, or both MyHCs (MyHC1-sto). From this, the proportion of each type of fiber to the total was calculated.

The cross-sectional area (CSA) of individual myofibers labeled with either myosin antibody was measured, based on the laminin-labeled outline of the myofiber, using Qwin Standard V.3.5.1 software (Leica Microsystems Ltd., Heerbrugg, Switzerland) on a randomized subset of images from each layer. CSA was calculated from 180 to 573 myosin-labeled myofibers/individual from the global layer and 130 to 472 myosin-labeled myofibers/individual from the orbital layer of each subject by tracing the perimeter of each myofiber, detected with laminin labeling, converting the enclosed pixels to a surface area (expressed in μm²).

Statistics

Statistical analyses and tests were carried out in the SPSS Statistics software (IBM, Armonk, NY, USA). Statistical comparisons between control and ALS groups were done as planned contrast tests with two levels of hypotheses: (1) Did ALS patients differ from control subjects, and (2) did ALS patients with bulbar or spinal onset differ from each other? All estimates were tested for nonnormal distributions using a Shapiro-Wilk test and Q-Q plots, with groups stratified down to the lowest expected level (controls, spinal-onset, and bulbar-onset). The test suggested that the proportion and CSA of MyHC1-sto fibers in control subjects, as well as the CSA of MyHC1-sto fibers in bulbar-onset patients, was nonnormal in the orbital layer. Based on graphic analysis, these estimates looked positively skewed with a possible heavy-tailed distribution. Since the underlying distribution for our estimates was unknown and likely to be nonnormal in some cases, we did all subsequent statistic tests of subject–group differences with a bias-corrected accelerated bootstrap sampling,22 with the sampling size set at 8000 runs. Testing of the three groups, based on the planned contrasts outlined above, was done with 1-way ANOVA with the above-mentioned bootstrap. Testing of differences between fiber types and the orbital and global layer within subject groups (controls, spinal-onset, and bulbar-onset) was performed with a paired t-test or, when appropriate due to nonnormality, a Wilcoxon matched-pair signed-rank test. Sidak correction was used with a subsequent P value adjustment for multiple comparisons where applicable. Estimates are presented as arithmetic mean ± 1 standard deviation and, where applicable, lower and upper bootstrap 95% CI (confidence interval).

A regression analysis was performed to investigate whether disease duration, age of onset, or age of death correlated with MyHC1-sto proportion in the global layer. We did the regression analysis on all three groups (controls and spinal-onset and bulbar-onset patients) separately. The residuals of cases were normally distributed in all groups, and no case had a standardized residual ≤ ± 1.96. Also, Cook’s distance was <1.0 for all cases. For estimation of P values and a 95% CI interval of the regression coefficient (β), a bias-corrected and accelerated CI was generated through an 8000-run bootstrap. To estimate generalizability of the model to the parent population, R² was adjusted with Darlington’s formula.23 P values for all tests were considered significant for P < 0.05.

RESULTS

Myofibers labeled with both MyHC antibodies (MyHC1-sto myofibers) were present across the whole EOM cross section and evenly distributed in both ALS and age-matched control subjects. Labeled myofibers did not appear to cluster in either group, but some areas in the global layer of ALS patients had very few myofibers labeled with either antibody. Those few labeled myofibers present in these areas were notably smaller than labeled myofibers in other areas.

MyHC1 and MyHCsto in Control EOMs

In the medial rectus muscle of control subjects, 12% to 18% of all myofibers were colabeled for MyHCsto and MyHC1 and evenly distributed across the whole muscle cross section. In all control individuals, a clear difference in labeling intensity with the antibodies against MyHCsto and MyHC1 was noted between the orbital and global layers. MyHCsto labeling was consistently strong in the orbital layer, but generally weak in the global layer. Conversely, MyHC1 labeling was consistently strong in the global layer but more moderate in the orbital layer (Fig. 1).

Almost every myofiber (>99.5%) labeled for MyHCsto was also labeled for MyHC1 in the middle portion of the medial rectus muscle (Fig. 1). These double-labeled MyHC1-sto myofibers corresponded to 14.0 ± 2.4% of the total number of myofibers in the global layer and 17.1 ± 3.0% in the orbital layer (Fig. 2). In addition, there were myofibers in the orbital and global layer labeled for MyHC1 but not for MyHCsto. These myofibers made up 1.9 ± 0.6% of the total number of myofibers in the global layer and 0.8 ± 0.4% of the total number of myofibers in the orbital layer (Fig. 2).

The mean CSA of MyHC1-sto myofibers was similar (P < 0.712) in the global layer (186 ± 72 μm²) and orbital layer (188 ± 39 μm²) (Fig. 3). The mean CSA of MyHC1-only myofibers was also similar (P < 0.712) in the global layer (157 ± 17 μm²) and orbital layer (144 ± 45 μm²). However, the global layer
myofibers in the orbital layer than in the global layer of both ALS and control subjects. In contrast, labeling with the antibody against MyHCsto (left) was stronger in the orbital layer than in the global layer of both ALS and control subjects. In control subjects, a higher proportion of MyHCsto myofibers in the orbital (mid) was consistently stronger in the global layer than in the orbital layer. Sporadic myofibers in both layers were labeled for MyHCl but were completely unlabeled with the antibody against MyHCsto (white arrows). In the orbital layer of ALS patients, a minority of myofibers had a labeling pattern where MyHCl reactivity was strong and MyHCsto reactivity was weak (open arrows). Notice the small CSA of myofibers labeled with the antibody against MyHCsto in (d1-d3), and that labeling with the antibody against MyHCsto is seen in only two myofibers (arrowheads). The antibody against laminin labels the basal lamina of myofibers, capillaries, and nerve fascicles. Lipofuscin inclusions (exemplified with open arrowheads) are visible in red and green filters due to their autofluorescent properties, but are easily distinguished from true double-labeling by their distinct color in the merged pictures and by their morphology. Scale bar: 50 µm.

MyHCsto myofibers were significantly larger than the MyHCl myofibers in that layer (P < 0.004), whereas in the orbital layer there was only a trend (P < 0.055) for similar differences between CSAs.

MyHCl and MyHCsto in the ALS EOMs

As in control individuals, a difference in MyHCsto and MyHCl labeling was noted between the orbital and global layers. However, in ALS patients, a minority of myofibers (estimated to be less than 5%) in the orbital layer exhibited a different labeling pattern where MyHCsto labeling was weak and MyHCl labeling was strong (Fig. 1).

In the medial rectus muscle of ALS patients, the proportion of double-labeled MyHC+sto myofibers was significantly lower than in control subjects both in the global layer (9.8 ± 3.1 vs. 14.0 ± 2.4% in controls, P < 0.002) and in the orbital layer (13.0 ± 4.3 vs. 17.1 ± 3.0% in controls, P < 0.014) (Fig. 2). There was no statistical difference between ALS patients with spinal-onset or bulbar-onset, but bulbar-onset patients had generally lower proportions of MyHC+sto myofibers in both the global layer (8.6 ± 3.0 vs. 10.7 ± 3.0% in spinal-onset patients, P < 0.13) and orbital layer (11.1 ± 2.6 vs. 14.6 ± 4.8% in spinal-onset patients, P < 0.08). The proportion of MyHC-only myofibers was significantly (P < 0.007) higher in ALS patients than in age-matched control individuals in the global layer (3.7 ± 1.7% in ALS vs. 1.9 ± 0.6% in controls), whereas in the orbital layer, there was only a trend (P < 0.053) in the same direction (1.6 ± 1.3% in ALS versus 0.8 ± 0.4% in controls).

As in healthy control subjects, a higher proportion of myofibers in the orbital layer than in the global layer was double-labeled for MyHC+sto (P < 0.002). As in controls, the proportion of MyHC-only myofibers in the global layer was higher than in the orbital layer (P < 0.001) in the ALS patients.

The mean CSA of MyHC+sto myofibers in the orbital layer did not differ between controls and ALS individuals (P < 0.371; Fig. 3). In contrast to controls, where the mean CSA of MyHC+sto fibers was very similar in the global and orbital layers, both spinal- and bulbar-onset ALS patients had significantly smaller MyHC+sto myofibers in the global layer.
The clinical course varied notably between them, with patient 1 (with a longer survival length) having a higher proportion of MyHCΙ+ surviving twice as long as patient 2. Patient 1 (with a longer survival length) had a considerably lower proportion of MyHCΙ+ sto fibers in the global layer (10.0%) that were notably smaller (131 μm²) as other ALS patients with spinal onset (patients 3 and 4) had very short survival (<1 year) and normal proportions (13.2% and 13.3%, respectively) of MyHCΙ+ sto fibers in the global layer that were of similar mean CSA as in control subjects (201 and 179 μm², respectively). Another spinal-onset C9orf72 ALS patient (patient 7) had slower progression and signs of cognitive involvement had a somewhat lower proportion of MyHCΙ+ sto fibers in the global layer (10.0%) that were notably smaller (131 μm²) compared to the orbital layer (155 ± 33 vs. 234 ± 73 μm², P < 0.004) in bulbar-onset patients and 153 ± 23 vs. 200 ± 32 μm², P < 0.010 in spinal-onset patients). The difference in mean CSA between ALS and control subjects did not reach statistical significance (P < 0.063), but suggested that global layer MyHCΙ+ sto myofibers may be smaller in ALS than in control individuals.

Two ALS patients with lower leg disease onset (patients 1 and 2) were siblings and homozygous carriers for the SOD1 D90A mutation. Despite their high genetic similarity (>50%), the clinical course varied notably between them, with patient 1 surviving twice as long as patient 2. Patient 1 (with a longer survival length) had a higher proportion of MyHCΙ+ sto fibers (12.4%) with a similar mean CSA (151 μm²) as other ALS patients, whereas patient 2 (with a shorter survival length) had a considerably lower proportion of MyHCΙ+ sto fibers (8.3%) that were also smaller in size (138 μm²).

Four ALS patients in the cohort had very long trinucleotide expansions in the gene C9orf72, a known genetic cause of ALS that can also give rise to frontotemporal dementia. Four C9orf72 patients constituted too few subjects for a statistical analysis, but some differences were noted. Two C9orf72 ALS patients with spinal onset (patients 3 and 4) had very short survival (<1 year) and normal proportions (13.2% and 13.3%, respectively) of MyHCΙ+ sto in the global layer that were of similar mean CSA as in control subjects (201 and 179 μm², respectively). Another spinal-onset C9orf72 ALS patient (patient 7) with slower progression and signs of cognitive involvement had a somewhat lower proportion of MyHCΙ+ sto fibers in the global layer (7.7%) with the smallest mean CSA of all subjects studied in the cohort (118 μm²). The data on fiber proportion and mean CSA in the orbital layer of these four patients and the two D90A patients mentioned above were more varied and did not segregate clearly into different clinical phenotypes.

**Correlations With Patient Age**

ALS patients as a group had significantly lower proportions of MyHCΙ+ sto fibers, especially in the global layer, this being the finding that most robustly separated ALS from control subjects. Simultaneously, there was a remarkable variation in proportions between individuals, varying between almost normal (13.1% in global layer of patient 11) and extremely low (4.0% in global layer of patient 18). Therefore, we investigated factors that could explain this variation in a regression analysis. We found that in bulbar-onset patients, but no other group, age of death (Fig. 4) correlated with a significant (P < 0.019) amount of variation in MyHCΙ+ sto fibers (R² = 0.669 for age of death) but practically none of the variation in the other two groups (Supplementary Fig. S1). Age of onset was borderline significant with a similar coefficient to the age of death regression. Interestingly, disease duration explained no variation in MyHCΙ+ sto proportion as a function of age of death in bulbar-onset patients (Supplementary Fig. S2). The regression coefficient (β) for MyHCΙ+ sto proportion as a function of age of death in years was –0.003 (bootstrap CI95%; –0.004 to –0.001). The population-adjusted R² was ≈0.46, implying that MyHCΙ+ sto proportion and age of death should correlate well in bulbar-onset patient populations beyond our study cohort. In the case of estimating P values of age of onset as a dependent of MyHCΙ+ sto proportion, convergence was difficult to achieve; even when performing in excess of 46,000 runs, the bootstrapped P value was inconsistent (range: 0.046–0.052) between individual runs.

![Figure 3. Bar graph showing the mean CSA of MyHCΙ+ sto myofibers (gray bars) and MyHCΙ-only (striped bars) myofibers in the orbital and global layer, respectively, of control and ALS subjects.](image1)

![Figure 4. Proportion of MyHCΙ+ sto myofibers in the global layer, as a function of age of death in bulbar-onset patients (crosses).](image2)
DISCUSSION

The major findings of the present study were (1) loss of MyHC<sup>+</sup> fibers in both layers of the medial rectus in ALS patients; (2) increase in the proportion of myofibers labeled only for MyHC<sub>1</sub> and lacking MyHC<sub>sto</sub> in the global layer of ALS patients; (3) high correlation of age of death, but not disease duration, with loss of MyHC<sub>1</sub>-sto myofibers in the global layer of patients with bulbar onset of symptoms, but not in patients with spinal onset nor in controls.

A previous study<sup>14</sup> reported the normal fiber type distribution in the superior, lateral, and inferior rectus along with the superior oblique of healthy human subjects with a mean age of 37.5 years, whereas the current study focused solely on the medial rectus of older subjects (mean age 66.0 years). Despite these differences, the data from the two studies are similar, with the mean proportion of MyHC<sub>sto</sub> fibers in the previous study being 15% in the orbital layer and 16% in the global layer<sup>14</sup> and the proportions in the current study being 14% and 17%, respectively. Comprising a larger myofiber sampling and with an older group of subjects, the current study showed that myofibers labeled with only MyHC<sub>sto</sub> were few, but approximately twice as prevalent in the global layer compared to the orbital layer, and that myofibers labeled with only MyHC<sub>1</sub> were approximately 20% smaller in size than myofibers labeled with both MyHC isoforms. The labeling patterns of MyHC<sub>1</sub> and MyHC<sub>sto</sub> were different in the orbital and global layers, with MyHC<sub>1</sub> labeling being particularly strong in the global layer and MyHC<sub>sto</sub> being particularly strong in the orbital layer. This could reflect a difference in epitope accessibility but could also reflect differences in the relative amount of MyHC isoforms present in these myofibers. The slow tonic MIFs of the global and orbital layers have previously been shown to have different electrophysiological properties<sup>24</sup> and different ultrastructural characteristics; for example, the orbital layer slow tonic myofibers have more mitochondria, more sarcoplasmic reticulum, and smaller myofilbrils than slow tonic fibers in the global layer.<sup>25</sup> These functional differences could possibly also be reflected in different ratios of MyHC isoforms in the sarcosomes of these myofibers.

In addition to the significant loss of MyHC<sub>1</sub>-sto fibers in the global and orbital layers of ALS patients, there was a tendency for decrease in CSA in MyHC<sub>1</sub>-sto fibers in the global layer, suggesting that the global layer MyHC<sub>1</sub>-sto fibers of EOMs may be more severely affected in ALS. Although the patient group included a number of familial cases with known genetic causes of ALS (namely SOD1 missense mutations and C9orf72 nucleotide expansions), the numbers of patients were too low for a statistical comparison between groups with different forms of familial ALS. However, low proportions of global layer MyHC<sub>1</sub>-sto fibers were noted in individual patients from both familial forms of ALS, suggesting that loss of slow tonic fibers is a shared feature across different types of ALS.

Orbital MyHC<sub>1</sub>-sto fibers have been reported to react to acute mechanical denervation with a noticeable hypertrophy that can last for at least 3 months.<sup>26</sup> Similar results have also been demonstrated in chicken limb slow tonic myofibers.<sup>27</sup> However, hypertrophy of orbital MIFs has not been reported in response to bilateral botulinum toxin treatment, where only singly innervated fibers in the orbital layer undergo a transient hypertrophy.<sup>28</sup> This suggests that denervation-induced hypertrophy in the MyHC<sub>sto</sub> myofibers of EOMs may be dependent on the mechanism of denervation. Due to the transient and mechanism-dependent hypertrophy of orbital MyHC<sub>sto</sub> myofibers in EOMs, it is difficult to study these changes in the context of ALS. The finding of an altered labeling of some MyHC<sub>1</sub>-sto fibers in the orbital layer raises the question whether these myofibers have maintained their innervation but shifted myofiber phenotype, or whether they reflect a new phenotype conveyed by reinnervation by another motor neuron. Motor neurons innervating slow type myofibers may exhibit a high sprouting competence and seem to be the most resilient motor neurons of the spinal cord in ALS,<sup>29,30</sup> but the present results suggest that this may not be the case for motor neurons in the oculomotor nucleus that innervate slow myofibers in the EOMs.

The present study supports previous findings that EOMs are only partially spared in ALS<sup>8</sup> and raises questions regarding the impact of loss of MyHC<sub>1</sub>-sto myofibers on eye motility. Although we lack quantitative data on eye motility, the patients studied here had not reported difficulties with eye motility during the course of their disease. MyHC<sub>1</sub>-sto myofibers are thought to be responsible for fine-tuning of ocular movements, regulation during vergence, and possibly also smooth pursuit movements.<sup>3</sup> Two recent studies conclude that the majority of ALS patients have either no eye motility disturbance or show deficits only in the executive functions of eye movements.<sup>52,53</sup> One of the studies reported that maximum saccadic velocity is slightly decreased in a subgroup of ALS patients,<sup>52</sup> suggesting some involvement of the oculomotor nucleus. Those patients also display catch-up saccades during smooth pursuit movements, indicating damage to the neurons involved in these movements.<sup>52</sup> It remains to be determined whether saccade replacement of smooth pursuit movements may be linked to the loss of MyHC<sub>1</sub>-sto myofibers. The neuroanatomic segregation<sup>16,17,34</sup> of singly innervated fiber (SIF) motor neurons and MIF motor neurons in the oculomotor nucleus allows for future studies to link the changes in MIFs in the EOMs more directly to possible changes at the motor neuron level.

The strong association between age of death and loss of MyHC<sub>1</sub>-sto myofibers in bulbar-onset patients warrants some consideration of its validity before interpreting the results. First, all groups passed the statistical assumptions set up for the linear regression, and a visual inspection of Figure 4 reveals that the bulbar-onset patients line up in an orderly way around the regression slope. Secondly, while the correlational coefficient was very small in the control and spinal-onset groups ($R^2 = 0.037$ and $R^2 = 0.019$, respectively), it was unexpectedly large ($R^2 = 0.669$) in the bulbar-onset group, suggesting that this correlation is a distinct characteristic of this group. The notion that this age-dependent decline is inherently different from normal aging is supported by studies of the oculomotor system that show no decline in neurons with aging.<sup>55,56</sup> It is further supported by the minimal difference in proportion of myofibers labeled for MyHC<sub>sto</sub> between a previous study on younger human subjects<sup>14</sup> and those older control subjects reported here, suggesting that both MIFs of the EOMs and their motor neurons undergo very limited degenerative changes with normal aging. Thirdly, while ALS pathophysiology is complex and poorly understood, it has been noted previously that the correlation between aging and ALS is more pronounced in bulbar-onset cases.<sup>37-39</sup>

In summary, this study has established the degree of loss of MyHC<sub>sto</sub> myofibers in ALS with different types of onset, duration, and genetics. This could possibly explain the deterioration of smooth pursuit movements previously reported in some ALS patients. Also, this study strengthens the notion that bulbar-onset ALS is more intimately associated with aging processes than spinal-onset ALS. Importantly, it suggests that while eye motility is generally spared in ALS, subtle degenerative changes do occur at the terminal stage of the disease.
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