Altered skeletal muscle glucose–fatty acid flux in amyotrophic lateral sclerosis

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Amyotrophic lateral sclerosis is characterized by the degeneration of upper and lower motor neurons, yet an increasing number of studies in both mouse models and patients with amyotrophic lateral sclerosis suggest that altered metabolic homeostasis is also a feature of disease. Pre-clinical and clinical studies have shown that modulation of energy balance can be beneficial in amyotrophic lateral sclerosis. However, the capacity to target specific metabolic pathways or mechanisms requires detailed understanding of metabolic dysregulation in amyotrophic lateral sclerosis. Here, using the superoxide dismutase 1, glycine to alanine substitution at amino acid 93 (SOD1G93A) mouse model of amyotrophic lateral sclerosis, we demonstrate that an increase in whole-body metabolism occurs at a time when glycolytic muscle exhibits an increased dependence on fatty acid oxidation. Using myotubes derived from muscle of amyotrophic lateral sclerosis patients, we also show that increased dependence on fatty acid oxidation is associated with increased whole-body energy expenditure. In the present study, increased fatty acid oxidation was associated with slower disease progression. However, within the patient cohort, there was considerable heterogeneity in whole-body metabolism and fuel oxidation profiles. Thus, future studies that decipher specific metabolic changes at an individual patient level are essential for the development of treatments that aim to target metabolic pathways in amyotrophic lateral sclerosis.

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Introduction

Amyotrophic lateral sclerosis (ALS) is a fatal neurodegenerative disease that is characterized by the degeneration of motor neurons in the brain, brainstem and spinal cord. The progressive loss of neurons in ALS results in muscle denervation, weakness and paralysis. Death usually occurs within 3–5 years from diagnosis (Brown and Al-Chalabi, 2017). ALS is variable in terms of age of onset, site of symptom onset and rate and pattern of disease progression. Underlying the clinical variability of ALS is complexity of genetic contribution and the pathogenic pathways that lead to the death of neurons (Brown and Al-Chalabi, 2017; Ghasemi and Brown, 2018).

While multiple pathogenic mechanisms, including neuronal hyperexcitability (Vucic et al., 2008; Kiernan, 2009), glutamate excitotoxicity (Rothstein et al., 1992), protein aggregation (Arai et al., 2006) and oxidative stress (Blasco et al., 2017), are proposed to contribute to the pathogenesis of ALS, studies also indicate that metabolic dysfunction is associated with disease progression (Dupuis et al., 2004; Ahmed et al., 2016; Ioannides et al., 2016).

In human ALS, an increase in resting energy expenditure (i.e. hypermetabolism) has been observed in approximately one-third to more than half of all patients (Desport et al., 2001, 2005; Bouteloup et al., 2009; Funalot et al., 2009; Vaisman et al., 2009; Jesus et al., 2018; Steyn et al., 2018a). More recently, we have shown that hypermetabolism in patients with ALS is associated with lower motor neuron dysfunction, more rapid disease progression and increased risk of earlier death (Steyn et al., 2018a). Despite this evidence of the importance of metabolism, the capacity to target metabolism to slow disease progression is restricted by our limited understanding of the underlying biochemistry of altered metabolism in ALS.

Previous studies in superoxide dismutase 1, glycine to alanine substitution at amino acid 93 (SOD1G93A) and superoxide dismutase 1, glycine to arginine substitution at amino acid 86 (SOD1G86R) mouse models of ALS suggest that altered metabolic balance is a feature of the disease (Dupuis et al., 2004), with early and persistent perturbations in the blood levels of metabolic hormones such as leptin, the presence of circulating ketone bodies and increased energy expenditure (Dupuis et al., 2004).
Reports of increased peripheral clearance of circulating lipids (Fergani et al., 2007) and increased expression of fatty acid metabolism genes in glycolytic muscle (Palamiuc et al., 2015) suggest that altered skeletal muscle metabolism could contribute to the altered metabolism observed in ALS. However, whether this altered profile of gene expression is associated with functional changes in glucose–fatty acid metabolism in the muscle of ALS mice, and how this might relate to metabolic changes at the whole-body level remains to be determined. Moreover, whether similar metabolic changes occur in the skeletal muscle of ALS patients and the relevance of these changes to the clinical and metabolic presentations in human ALS is unknown.

In this study, we aimed to address this gap in knowledge by characterizing whole-body metabolism in the SOD1<sup>G93A</sup> mouse model of ALS throughout the course of disease, and by assessing energy substrate utilization in intact glycolytic muscle fibres isolated from SOD1<sup>G93A</sup> mice when compared to wild-type (WT) controls. We also assessed fatty acid and glucose oxidation in myotubes derived from well characterized ALS patients and healthy controls to determine whether substrate metabolism in ALS muscle is linked to the energy expenditure profiles and clinical features. We report evidence of a functional alteration in glucose–fatty acid flux in glycolytic muscle of SOD1<sup>G93A</sup> mice. In ALS patient-derived myotubes, changes in glucose and fatty acid oxidation are associated with whole-body energy expenditure and disease progression, but not hypermetabolism.

**Materials and methods**

**Animal studies**

**Mice**

Experiments at the University of Queensland were approved by the University of Queensland Animal Ethics Committee and conducted in accordance with the Queensland Government Animal Care and Protection Act 2001, associated Animal Care and Protection Regulations (2002 and 2008) and the Australian Code of Practice for the Care and Use of Animals for Scientific Purposes (2004). Experiments at the University of Notre Dame were conducted in accordance with the Freimann Life Science Center Guidelines and the Notre Dame Institutional Animal Care and Use Committee Policy on Humane Endpoints for animal use. Transgenic mice overexpressing the human SOD1<sup>G93A</sup> mutation [B6-Cg-Tg (SOD1-G93A) 1Gur/J] (Gurney et al., 1994) were purchased from the Jackson Laboratory (Bar Harbor, ME, USA) and bred on a C57 black 6 (C57BL/6j) background. Male SOD1<sup>G93A</sup> mice and litter- or age-matched WT control mice (n = 3–26/group) were randomly assigned for experiments at ages that correspond to pre-defined stages of disease in SOD1<sup>G93A</sup> mice; presymptomatic (5 weeks of age, no symptoms), onset (9–11 weeks of age, early signs of hindlimb tremor and weakness), mid-stage (16–19 weeks of age, pronounced hindlimb weakness) and end-stage (21–25 weeks of age, significant hindlimb weakness leading to paralysis and euthanasia due to loss of the righting reflex) (Ngo et al., 2012; Lee et al., 2013). In line with ethical requirements, SOD1<sup>G93A</sup> mice were monitored twice daily after the onset of pronounced hindlimb weakness (i.e. mid-stage of disease). All mice were group-housed (3–4 mice per cage) in filter top cages, or in individually ventilated cages when maintained in a specific pathogen-free environment. For the assessment of energy expenditure, mice were single-housed in cages designed specifically for the Phenomaster metabolic phenotyping system (TSE-Systems, Bad Homburg, Germany). Mice were maintained on a 12 h light, 12 h dark cycle (ON at 0600 h and OFF at 1800 h) and had free access to food (20% protein, 4.8% fat; Specialty Feeds, WA, Australia) and water. Room temperature was maintained at 22 ± 2°C. Assessments in live mice or collection of tissue from mice occurred between 1000 and 1400 h on any given day. Prior to tissue collection, all mice were anaesthetized with an intraperitoneal injection of sodium pentobarbitone (32.5 mg/kg, Virbac Animal Health, NSW, Australia), Following complete loss of the pedal withdrawal reflex and eye-blink reflex, mice were killed by cervical dislocation. All animal work was conducted in accordance with the Animal Research: Reporting of In Vivo Experiments guidelines (Kilkenny et al., 2010).

**EchoMRI assessment of fat and fat free mass**

Imaging took place in the Notre Dame Integrated Imaging Facility Freimann Life Sciences Center. Whole-body fat and fat free mass was measured with an EchoMRI-130<sup>TM</sup> Quantitative Magnetic Resonance (EchoMRI, TX, USA) (Metzinger et al., 2014). Two X-ray images were taken at different energy levels to assess soft tissue density and bone. Mass measurements for fat and fat free tissue were produced for all scans in the EchoMRI Body Composition Analyzer EMR-184 software (EchoMRI).

**Indirect calorimetry**

Energy expenditure was measured with a Phenomaster open-circuit indirect calorimetry system housed within a temperature (22°C) and 12 h light, 12 h dark cycle (ON at 0600 h and OFF at 1800 h) controlled chamber (TSE-Systems), as we have done previously (Steyn et al., 2018b). All mice were acclimated for single-housing before collection of individualized measures using the Phenomaster system. Experimental cages (n = 16) were sampled at 60 min intervals for 3.5 min/cage, with concentrations of O<sub>2</sub> and CO<sub>2</sub> in the outgoing air being measured sequentially within each interval. One vacant cage was included to obtain a reference concentration for ambient gas. Activity (x- and y-plane), food intake and
body weight were recorded synchronously with metabolic data. Measurements were performed continuously over 72 h, with analysis restricted to the final 24 h assessment period (allowing 48 h of acclimation). For data analysis, measures of total energy expenditure and food intake were adjusted for body weight. Food intake measures are representative of hourly food intake for each mouse.

**18F-deoxyglucose PET/single-photon emission computed tomography/CT imaging**

Animals were anaesthetized using isoflurane inhalation for 2–5 min in a vaporized-controlled tank to allow sufficient anaesthesia followed by recovery. Animal breathing was checked once every minute by visual inspection. 18F-deoxyglucose at a dose of 0.200mCi activity was administered to mice intravenously through the tail vein (volume < 100 μl). PET images were acquired on a trimodal Alibra PET/single-photon emission computed tomography/CT image station (Carestream Health, Woodbridge, CT, USA) to produce high-resolution PET images that were reconstructed for analysis. 18F-deoxyglucose uptake in brown adipose tissue (BAT) was quantified as the mean voxel value within a visually determined volume of interest as described previously (van der Veen et al., 2012).

**Ex vivo lipolysis**

Epididymal white adipose tissue (WAT) was excised and rinsed in 1x phosphate buffered saline supplemented with 0.1% fatty acid-free bovine serum albumin. All epididymal WAT explants were placed in plastic vials containing 1 ml of modified Krebs–Henseleit buffer (in mM): 4.7 KCl, 1.2 KH2PO4, 1.2 MgSO4·7H2O, 1.25 CaCl2·2H2O, 2.5 NaHCO3, 5 glucose, 118 NaCl and 4% fatty acid-free bovine serum albumin. Buffer was gassed with 95% O2/5% CO2 for 45 min to reach a pH of 7.4. All procedures were conducted in a shaking water bath at 37°C. For the assessment of non-esterified fatty acids (NEFA), 6 μl of buffer was collected at 0-, 30-, 60-, 90- and 120-min time points, placed immediately on dry ice and stored at −80°C. Samples were assayed on a NEFA-C kit (Wako Chemicals, Osaka, Japan). For glycerol, buffer was collected after 2 h of incubation. Glycerol content was assessed using a free glycerol determination kit with glycerol standard solution (Sigma-Aldrich, MO, USA). Final glycerol content and NEFA-C levels were expressed relative to the weight of the respective epididymal WAT explant.

**Plasma NEFA**

Following sacrifice, terminal blood samples were collected from mice via cardiac puncture. Samples were transferred into ethylenediaminetetraacetic acid-precoated tubes and centrifuged for 3 min. Plasma was aliquoted and stored at −80°C until use. NEFA levels in plasma were determined using a NEFA-C test kit (Wako Chemicals).

**Oil Red O staining**

Extensor digitorum longus (EDL) muscles were embedded in optimum cutting temperature compound and rapidly frozen in liquid nitrogen cooled isopentane. Oil Red O (ORO) staining was performed on muscle cryosections (10 μm) to visualize neutral lipids using an ORO kit (Abcam, Cambridge, UK). Briefly, sections were fixed with 10% neutral buffered formalin for 15 min and rinsed three times with distilled water for 30s at room temperature. ORO solution was added onto the sections and incubated for 10 min, and the slides were differentiated in 85% propylene glycol solution for 1 min at room temperature. After two rinses, slides were air dried and mounted with aqueous mounting agent (Aquatex, EMD Millipore, CA, USA). Bright-field images (20× magnification) were taken with an Aperio ScanScope system (Leica, Mannheim, Germany). ORO labelling was quantified using 10+ randomly selected sections of muscle per animal (n=5 per group). Representative images of the muscle section were processed using ImageJ to identify the mean ORO intensity for each animal, following thresholding to remove non-specific labelling [threshold set to 225 (0–255)].

**Assessment of cellular respiration in muscle fibre bundles**

EDL muscle fibre bundles were chemically dissociated as previously described (Li et al., 2016). Muscle fibre bundles were seeded onto Seahorse XF®96 microplates in culture media (low glucose Dulbecco’s Modified Eagle Medium supplemented with 10% foetal bovine serum and 1% Antibiotic-Antimycotic; ThermoFisher, MA, USA) and maintained overnight at 37°C with 5% CO2. Prior to the commencement of metabolic assays, muscle fibre viability was assessed using an alamarBlue cell viability assay (ThermoFisher) and used for the data normalization. Real-time assessment of bioenergetic parameters in EDL fibre bundles was performed on the XF®96 Extracellular Flux Analyzer (Agilent Technologies, CA, USA).

The dependence and capacity of EDL fibres to use glucose and fatty acid as fuel substrates were determined using the Seahorse XF Mito Flex Test Kit (Agilent Technologies). Prior to the assay, culture media was replaced with pre-warmed assay media (pH 7.4) consisting of extracellular flux base media (Agilent Technologies), 10 mM D-glucose (Sigma-Aldrich), 1 mM sodium pyruvate (Sigma-Aldrich) and 2 mM L-glutamine (ThermoFisher). Carnitine palmitoyltransferase 1A inhibitor etomoxir (ETO, 4.0 μM), mitochondrial pyruvate carrier inhibitor UK5099 (2.0 μM) and glutaminase inhibitor BPTES (3.0 μM) were prepared with assay media and loaded into the XF®96 sensor cartridge following the manufacturer guidelines. Following the first three cycles of baseline measurement of oxygen consumption rate (OCR), the decrease of OCR levels upon inhibition of one or more pathways was continuously recorded for the
following six cycles. Each cycle consisted of 3 min mix, 30 s wait and 3 min measurement. The dependence and capacity of each fuel pathway relative to total fuel oxidation were calculated according to the manufacturer guidelines.

Substrate induced maximal respiration was tested in EDL fibre bundles as previously described (Li et al., 2016). Briefly, plates containing muscles fibres were changed into assay media (pH 7.4) containing 120 mM NaCl, 3.5 mM KCl, 1.3 mM CaCl₂, 0.4 mM KH₂PO₄, 1 mM MgCl₂, 2.5 mM d-glucose and 0.5 mM L-carnitine prior to the assay run. OCR was continuously measured for six cycles after sequential injections of sodium pyruvate (10 mM) or palmitate-bovine serum albumin (100 µM) with carbonyl cyanide-p-trifluoromethoxyphenylhydrazone (0.4 µM), followed with antimycin/rotenone (1 µM).

**Patient studies**

**Subjects**

Eighteen ALS patients who met the revised El-Escorial criteria for ALS (Brooks et al., 2000) were enrolled from the Royal Brisbane and Women’s Hospital ALS clinic for the collection of skeletal muscle biopsies. Eleven healthy control participants were also enrolled. These control individuals were the spouses, friends or family members of ALS participants. For all participants, exclusion criteria were history of a metabolic condition (e.g. Hashimoto’s disease) and diabetes mellitus. Participant details are shown in Table 1 and Supplementary Table 1. For ALS patients, the ALS functional rating scale-revised (ALSFRS-R) score, King’s stage and deltaFRS were obtained from clinical records. All participants provided written informed consent; participant consent was obtained according to the Declaration of Helsinki. Work performed in this study was approved by the Royal Brisbane and Women’s Hospital and University of Queensland human research ethics committees.

**Assessment of body composition and energy expenditure**

Body composition (fat mass and fat free mass) was determined by whole-body air displacement plethysmography using the BodPod system (Cosmed USA, Rome, Italy) (Ioannides et al., 2017). Values of fat mass and fat free mass were used to predict resting energy expenditure (Steyn et al., 2018a). Energy expenditure in fasting and alert ALS and control participants at rest (i.e. resting energy expenditure) was then measured by indirect calorimetry using a Quark RM respirometer (Cosmed) as per our established methodology (Steyn et al., 2018a). Controls were matched to patients with ALS by age, sex, weight, body mass index and body composition. The metabolic index (MI) of each individual was derived by calculating measured resting energy expenditure as a percentage of predicted resting energy expenditure. An MI (i.e. measured energy expenditure relative to predicted resting energy expenditure) ≥120% was defined as hypermetabolism, as we have done previously (Steyn et al., 2018a).

**Muscle biopsy and culture**

Muscle biopsies were collected from the vastus lateralis of one leg. Lignocaine (1%; 5 ml) was injected to anaesthetize the skin and underlying fat and muscle tissue. A 10-mm incision was made and advanced through the fascia of the muscle. A ~200 mg sample of muscle was collected using a sterile 6-mm hollow Bergstrom biopsy needle (Pelomi, Albertslund, Denmark) modified for suction (Tarnopolsky et al., 2011) and placed in holding media (Dulbecco’s Modified Eagle Medium/Ham’s F-12 with 0.5% gentamicin; ThermoFisher).

Primary myoblasts were isolated and cultured using a modified muscle tissue explant method (Tarnopolsky et al., 2011) and frozen as low passage cell stocks. Primary myogenic cells were maintained in Dulbecco’s Modified Eagle Medium/Ham’s F-12 medium supplemented with 20% foetal bovine serum, 10% AminoMAX C-100 and 0.5% gentamicin (ThermoFisher). Culture media was changed every second day and cells were passaged when they reached ~70% confluence. For experiments, primary myogenic cells were seeded at a density of 15 000 cells/well into Seahorse XF96 cell culture microplates. At 80% confluence, cells were differentiated into myotubes by replacing maintenance media with differentiation media consisting of Dulbecco’s Modified Eagle Medium/Ham’s F-12 medium with 2% non-inactivated horse serum and 0.5% gentamicin (ThermoFisher). Fresh differentiation medium was fed every 2 days until mature multinucleated myotubes formed. Primary myotubes underwent assessment of key parameters of glucose

### Table 1 Characteristics of ALS patients and healthy controls at the time of muscle biopsy collection

<table>
<thead>
<tr>
<th></th>
<th>Control (n = 11)</th>
<th>ALS (n = 18)</th>
<th>P</th>
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<tbody>
<tr>
<td><strong>Demographics</strong></td>
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<tr>
<td>Age (years)</td>
<td>58.9 ± 10.1</td>
<td>55.4 ± 7.2</td>
<td>0.277</td>
</tr>
<tr>
<td>Sex, female</td>
<td>3 (27.27)</td>
<td>4 (22.2)</td>
<td>0.758</td>
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<tr>
<td><strong>Anthropometric and metabolic measures</strong></td>
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<tr>
<td>Body mass index (kg/m²)</td>
<td>25.9 ± 2.7</td>
<td>26.6 ± 4.2</td>
<td>0.644</td>
</tr>
<tr>
<td>Fat free mass (kg)</td>
<td>53.4 ± 10.5</td>
<td>54.7 ± 10.8</td>
<td>0.745</td>
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<tr>
<td>Fat mass (%)</td>
<td>31.1 ± 9.4</td>
<td>32.4 ± 11.5</td>
<td>0.759</td>
</tr>
<tr>
<td>EEkc (kcal/day)</td>
<td>1596 ± 331.1</td>
<td>1809 ± 336.2</td>
<td>0.107</td>
</tr>
<tr>
<td>MI (%)</td>
<td>108.4 ± 15.41</td>
<td>119.5 ± 9.6</td>
<td>0.025</td>
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<tr>
<td><strong>Clinical scores</strong></td>
<td></td>
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<td></td>
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<tr>
<td>Time since onset (months)</td>
<td>29.2 ± 21.5</td>
<td></td>
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<tr>
<td>ALSFRS-R</td>
<td>36.7 ± 6.1</td>
<td></td>
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<tr>
<td>deltaFRS</td>
<td>−0.5 ± 0.3</td>
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<tr>
<td>King’s stage</td>
<td>2.4 ± 1.0</td>
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<tr>
<td>FVC (% of predicted)</td>
<td>88.5 ± 22.1</td>
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Data presented as mean (SD) and n (%) for categorical data. Means were compared by independent t-test and proportions with the chi-square test. ALSFRS-R = ALS functional rating scale-revised; deltaFRS = change in ALSFRS-R since symptom onset; EEkc = energy expenditure kilocalories; FVC = forced vital capacity; MI = metabolic index (measured EEkc/predicted EEkc).
and fatty acid oxidation dependency and capacity using the Seahorse Mito Fuel Flex Test Kit (Agilent Technologies). Parameters of glycolytic flux (glycolysis, glycolytic capacity and glycolytic reserve) were determined using the Seahorse XF Glycolysis Stress Test Kit (Agilent Technologies).

**Statistical analysis**

Statistical differences were assessed using Prism 8.0a. (GraphPad Software Inc., La Jolla, CA, USA). Statistical comparisons were performed following testing for normal distribution, and unless otherwise indicated, outcomes were compared using unpaired Student’s t-test, non-parametric t-test or two-way ANOVA followed by Bonferroni multiple comparison. Percentage data were log transformed prior to statistical comparison. Linear relationships were assessed by Pearson correlation. All graphical data are presented as mean ± SD. Values of *P* < 0.05 were considered to be statistically significant.

**Data availability**

The data from this study are available from the corresponding author, upon reasonable request.

**Results**

**Disease progression in SOD1<sup>G93A</sup> mice is associated with a decline in both fat free mass and fat mass**

To determine the impact of disease progression on body composition and total body weight in SOD1<sup>G93A</sup> mice, we conducted a serial assessment of body weight, and whole-body fat free mass and fat mass by EchoMRI (Fig. 1A–C). Total body weight in SOD1<sup>G93A</sup> mice was significantly lower than that of litter-matched WT controls by 19 weeks of age (Fig. 1A). Loss of body weight was, in part, to a loss in total fat free mass (Fig. 1B). Tibialis anterior and gastrocnemius weight were reduced by disease onset (Fig. 1D and E), and EDL mass was reduced by the mid-stage of disease (Fig. 1F). When compared to age-matched WT controls, fat accumulation in SOD1<sup>G93A</sup> mice slowed following disease onset, and whole-body fat mass was significantly lower from 15 weeks of age (Fig. 1C). Epididymal and inguinal fat mass was lower in SOD1<sup>G93A</sup> mice from the mid-stage of disease (Fig. 1G and H). Consistent with this, circulating levels of leptin were stable in SOD1<sup>G93A</sup> mice throughout the disease course while the WT controls showed increased levels of leptin with time (Fig. 1I).

**Total energy expenditure in SOD1<sup>G93A</sup> mice increases by the mid-stage of disease**

We next sought to determine whether energy expenditure in SOD1<sup>G93A</sup> mice increases relative to disease progression. When compared to WT controls, SOD1<sup>G93A</sup> mice at the mid-stage of disease had higher levels of energy expenditure during the light and dark cycle, resulting in an overall increase in 24 h total energy expenditure (Fig. 2A and B). SOD1<sup>G93A</sup> mice had decreased activity by disease onset (Fig. 2C and D).

We also investigated whether reductions in body weight and fat mass in SOD1<sup>G93A</sup> mice were due to a decline in food intake. We observed no change in food intake between SOD1<sup>G93A</sup> mice at disease onset and litter-matched WT controls. SOD1<sup>G93A</sup> mice at the mid-stage of disease consumed more food than litter-matched WT controls (Fig. 2E and F). Thus, reductions in weight are not associated with decreased food consumption, and despite an increase in food intake and a decline in activity-dependent energy expenditure, SOD1<sup>G93A</sup> mice are unable to offset increased energy expenditure to prevent the depletion of energy stores.

**Lipolytic rate is maintained in SOD1<sup>G93A</sup> mice**

BAT regulates non-shivering thermogenesis, which itself can contribute to total energy expenditure in mice (Even and Blais, 2016). We found no difference in BAT weight (Fig. 3A) or glucose uptake in BAT between SOD1<sup>G93A</sup> and WT control mice (Fig. 3B and C). Thus, the increase in energy expenditure in SOD1<sup>G93A</sup> at the mid-stage of disease is unlikely to be due to increased non-shivering thermogenesis.

WAT stores lipids, providing energy reserves that are available during periods of increased energy demand. Given the slowing of accumulation and eventual reduction of fat mass in SOD1<sup>G93A</sup> mice, we assessed the release of lipids (a proxy measure for lipolysis) from epididymal WAT explants. The lipolytic rate of ex vivo explants of epididymal WAT decreased over the lifespan of WT mice, while the lipolytic rate in SOD1<sup>G93A</sup> mice was maintained throughout the course of disease. The lipolytic rate in SOD1<sup>G93A</sup> mice was significantly higher by the mid-stage of disease when compared to WT controls (Fig. 3D and E). This corresponded to a sustained rate of appearance of cumulative NEFA and glycerol from WAT explants (Fig. 3F). Circulating plasma NEFA in WT and SOD1<sup>G93A</sup> mice did not differ (Fig. 3G). Overall, results suggest that the rate of lipolysis in epididymal WAT explants from SOD1<sup>G93A</sup> is sustained throughout disease, and that it does not decrease with age, as is seen in WT mice.
SOD1<sup>G93A</sup> mice exhibit a functional preference for fat oxidation in glycolytic EDL muscle

Previously, a decrease in the expression of glucose handling genes and an increase in the expression of lipid handling genes in glycolytic muscle of SOD1<sup>G86R</sup> mice have been suggested to drive a switch towards the use of lipid as an energy substrate (Palamiuc et al., 2015). Thus, sustained lipolysis in SOD1<sup>G93A</sup> mice could serve to mobilize fatty acids for use as an energy source in skeletal muscle. We used ORO staining to quantify intramuscular lipid accumulation in the glycolytic EDL muscle and conducted real-time assessment of substrate utilization in EDL muscle fibre bundles isolated from SOD1<sup>G93A</sup> and age-matched WT mice. There was no difference in intramuscular lipid content in the EDL muscle of SOD1<sup>G93A</sup> mice when compared to WT controls (Fig. 4A and B). We observed no difference in glucose oxidation dependence (Fig. 4C and D), and following the inhibition of mitochondrial fatty acid and glutamine uptake, EDL muscle fibres from SOD1<sup>G93A</sup> mice exhibited similar levels of glucose oxidation capacity across all disease stages (Fig. 4E and F). In contrast to the decline in fat oxidation and dependence in WT mouse skeletal muscle fibres with age, we observed sustained dependence on fat oxidation in isolated EDL muscle fibre bundles from SOD1<sup>G93A</sup> mice by the mid-stage of disease (Fig. 4G and H). Moreover, the capacity for fat oxidation to sustain mitochondrial respiration after inhibition of pyruvate and glutamine entry into mitochondria was also increased in EDL muscle fibre bundles from SOD1<sup>G93A</sup> mice at the mid-stage of disease when compared to WT controls, which exhibited a decline in fat oxidation capacity with age (Fig. 4I and J).

To determine the capacity of glucose and fatty acid oxidation pathways to sustain mitochondrial respiration in the absence of substrate competition, and to study substrate utilization in the presence of increased energy demand, we measured the capacity of mitochondria in EDL fibre...
bundles to oxidize pyruvate or palmitate in the presence of the mitochondrial uncoupler, carbonyl cyanide-\(p\)-trifluoromethoxyphenylhydrazone (Fig. 5A and B). The basal OCR of EDL muscles from WT and SOD1\(^{G93A}\) mice was similar (Fig. 5C). In the presence of pyruvate, we observed a significant elevation in maximal peak OCR in EDL muscle fibre bundles from SOD1\(^{G93A}\) mice at the mid-stage stage of disease (Fig. 5A). In the presence of palmitate, we observe no differences in maximal OCR in EDL muscle fibre bundles between SOD1\(^{G93A}\) and WT controls (Fig. 5B). Compared to WT mice, total oxygen consumption in the EDL muscle fibres from SOD1\(^{G93A}\) mice was significantly higher at the mid-stage of disease when pyruvate was provided as the external energy substrate (Fig. 5D). However, oxygen consumption between EDL muscle fibres from SOD1\(^{G93A}\) mice and WT mice was comparable when palmitate-bovine serum albumin was provided as the external energy substrate (Fig. 5E). Thus, despite exhibiting a functional preference towards fatty acid oxidation, glycolytic EDL muscle fibres from SOD1\(^{G93A}\) mice are capable of utilizing glucose metabolism pathways to sustain mitochondrial function when there are no competing fatty acid substrates.

**ALS patient-derived myotubes have increased dependence on fat oxidation**

We next aimed to determine whether the preferential use of lipids by mitochondria of SOD1\(^{G93A}\) mice was also present in human ALS subjects. We generated primary myotubes from skeletal muscle from ALS patients and age-matched healthy controls and conducted real-time assessment of substrate utilization. Demographics of our study population are detailed in Table 1. Sex, age and clinical demographics of ALS patients are detailed in Supplementary Table 1. When compared to myotubes derived from healthy controls, myotubes from patients with ALS had similar levels of glucose oxidation.
dependency and capacity (Fig. 6A). ALS patient-derived myotubes also had similar levels of fat oxidation capacity compared to myotubes derived from healthy controls. However, they exhibited an increased dependence on fat oxidation (Fig. 6B).

The transition between glucose and fatty acid oxidation can ultimately impact the flux of metabolic substrates through their respective pathways. We therefore sought to determine whether the increase in fat oxidation dependence in ALS patient-derived myotubes was associated with altered glycolytic function. We found no difference in glycolysis, glycolytic capacity or glycolytic reserve between myotubes derived from control individuals and myotubes derived from patients with ALS (Fig. 6C).
Figure 4 A functional shift in mitochondrial fuel preference from glucose to fat occurs in glycolytic EDL muscle of SOD1<sup>G93A</sup> mice. (A) ORO staining and (B) quantification of intramuscular lipid in the glycolytic EDL muscle of SOD1<sup>G93A</sup> and age-matched WT mice (n=5/group), scale bar = 50 μm. (C) Data trace of mitochondrial OCR (% baseline) contributed to by glucose oxidation dependence. (D) Quantification of glucose oxidation dependence in isolated EDL muscle fibre bundles from SOD1<sup>G93A</sup> mice and WT controls. (E) Data trace of mitochondrial OCR (% baseline) contributed to by glucose oxidation capacity. (F) Quantification of glucose oxidation capacity in isolated EDL muscle fibre bundles from SOD1<sup>G93A</sup> mice and WT controls. (G) Data trace of mitochondrial OCR (% baseline) contributed to by fat oxidation dependence. (H) Quantification of fat oxidation dependence in SOD1<sup>G93A</sup> mice and WT control mice. (I) Data trace of mitochondrial OCR (% baseline) contributed to by fat oxidation capacity. (J) Quantification of fat oxidation capacity in SOD1<sup>G93A</sup> mice and WT controls. White circles represent WT mice; teal circles represent SOD1<sup>G93A</sup> transgenic mice. All data presented as mean ± SD for n=3–13/group. *P < 0.05, two-way ANOVA with Bonferroni’s post hoc test. MS = mid-stage; OS = onset; PS = presymptomatic.
We next assessed the relationships between substrate oxidation and glycolytic function in myotubes and the metabolic and clinical characteristics of our ALS cohort. The metabolic measures were resting energy expenditure and the MI, which we have previously used to define hypermetabolism in ALS patients (Supplementary Table 2) (Steyn et al., 2018a). We found that glucose oxidation capacity and glycolytic reserve were negatively correlated with resting energy expenditure, and that fat oxidation dependency and capacity, as well as glycolysis, were greater in myotubes derived from ALS patients with higher resting energy expenditure.
We observed no relationship between substrate oxidation or glycolytic function in ALS patient-derived myotubes relative to the severity of disability as determined by the ALS functional rating scale-revised at the time of metabolic assessment (Supplementary Table 2). However, glucose oxidation capacity was greater, and fat oxidation dependence was lower in patients with a more rapidly progressing disease [indicated by a faster decline in ALS functional rating scale-revised scores (deltaFRS); Supplementary Table 2 and Fig. 7]. Collectively, these data indicate that substrate utilization in myotubes is not related to hypermetabolism but is linked to the resting energy expenditure of ALS patients. Moreover, our data also suggest that substrate utilization in myotubes is linked to the resting energy expenditure of ALS patients.

Figure 6 Myotubes derived from patients with ALS have increased dependence on fat oxidation. (A) Data traces and quantification of mitochondrial OCR (% baseline) contributed to by glucose oxidation dependency and capacity in ALS and control myotubes. (B) Data traces and quantification of mitochondrial OCR (% baseline) contributed to by fat oxidation dependency and capacity in ALS and control myotubes. (C) Data trace of extracellular acidification rate, and quantification of glycolysis, glycolytic capacity and glycolytic reserve in ALS and control myotubes. All data presented as mean ± SD for n = 11 control and n = 18 ALS individuals. *P < 0.05, Unpaired t-test. Individuals with a C9orf72 expansion are noted in grey, and an individual with a SOD1 mutation is noted in blue.
associated with the rate of functional decline in patients with ALS.

Discussion

The primary aim of this study was to explore the potential mechanisms underpinning altered whole-body metabolic balance in ALS (Ahmed et al., 2018; Steyn et al., 2018a; Vandoorne et al., 2018). We show that an increase in whole-body energy expenditure in symptomatic SOD1<sup>G93A</sup> mice is associated with a decline in both fat mass and fat free mass. We also find that glycolytic muscle from symptomatic SOD1<sup>G93A</sup> mice exhibit an increased dependence on fatty acids as an energy substrate, as well as an increased capacity to utilize fatty acids. In myotubes derived from patients with ALS, we show a similar association between dependence on fatty acid oxidation and resting energy expenditure. Further there was an association between muscle cellular metabolism and rate of disease progression.

Compared to WT littermates, SOD1<sup>G93A</sup> mice exhibited an increase in total whole-body oxygen consumption (a proxy for energy expenditure) alongside a concomitant decline in body mass, fat mass and fat free mass over the course of disease. These observations are in line with the notion that increased energy expenditure in ALS contributes to weight loss (Dupuis et al., 2011). However, given that hypermetabolism has not been shown to be associated with weight loss in human ALS (Steyn et al., 2018a), these divergent observations between ALS mice and patients suggest that multiple factors are likely to contribute to weight loss in patients with ALS. These include reduced capacity to meet energy requirements (Ngo et al., 2017) [including loss of appetite resulting in loss of fat mass (Ngo et al., 2019; Mezoian et al., 2020)], as well as neurogenic wasting (Al-Sarraji et al., 2014).

A failure to accumulate fat mass was an early feature of disease in SOD1<sup>G93A</sup> mice. When compared to WT mice, where the lipolytic rate of WAT explants decreased with age, the rate of lipolysis of WAT explants from SOD1<sup>G93A</sup> mice was sustained over the course of disease. The sustained rate of lipolysis by disease mid-stage corresponded to greater energy needs, as indicated by the higher total energy expenditure of SOD1<sup>G93A</sup> mice at this time. That circulating levels of fatty acids (NEFA) were not increased in SOD1<sup>G93A</sup> mice, even though WAT explants were primed for lipolysis, would suggest that lipids that entered circulation were rapidly utilized. It is well established that lipolysis and fatty acid mobilization are upregulated in response to increased muscle energy requirements, for example, during exercise (Goodpaster and Sparks, 2017). Thus, increased mobilization of lipids in ALS mice presumably occurs as a means to sustain metabolic requirements in peripheral skeletal muscle, as has been suggested previously (Dupuis et al., 2004; Fergani et al., 2007). Expanding on previous reports of increased expression of genes associated with fatty acid oxidation in skeletal muscle of SOD1<sup>G86R</sup> mice (Palamiuc et al., 2015), we have now generated the first evidence to show that there is a greater functional dependence and capacity of glycolytic skeletal muscle from symptomatic SOD1<sup>G93A</sup> mice to utilize fatty acids as a fuel substrate. This is in contrast to what is observed in WT mice, where fat oxidation declined from 3 weeks of age through to 20–25 weeks of age. This decline in fat oxidation is in line with, and likely to be secondary to our observations of decreased lipolytic rate of WT epididymal WAT explants and stable levels of plasma NEFA, indicating that the metabolic demand of skeletal muscle in WT mice is possibly lower than that of SOD1<sup>G93A</sup> mice. Although the increased dependence and capacity for fatty acid oxidation in SOD1<sup>G93A</sup> mice could be an adaptive response to muscle metabolic demand, it might also occur due to reduced muscle glucose uptake and glucose intolerance (Pradat et al., 2010; Desseille et al., 2017). Yet, we found that glycolytic muscle fibres were able to utilize pyruvate, a product of glucose oxidation, following inhibition of fatty acid and glutamine pathways. While providing evidence to suggest that glucose oxidation mechanisms remain intact in glycolytic SOD1<sup>G93A</sup> muscle, our data lend further support to the notion that increased utilization of fatty acids could inhibit the use of glucose in ALS.
and glucose oxidation capacity is paradoxical, given that a negative correlation between resting energy expenditure to increased energy demand through this pathway. The maximum, and therefore, there is reduced capacity to respond lytic reserve. High levels of glycolysis indicate that the correlation between resting energy expenditure and glycolysis metabolism (Turner et al., 2014). As such, an increase in the need for fatty acid oxidation in myotubes obtained from patients with higher resting energy expenditure may function to increase ATP availability. Moreover, the correlation observed between resting energy expenditure and fatty acid oxidation dependence aligns with physiological processes whereby continued dependence on fatty acid metabolism requires greater levels of oxygen consumption when compared to glucose metabolism (Turner et al., 2014). Interestingly, myotubes derived from ALS patients with higher resting energy expenditure also had higher glycolysis, but lower glucose oxidation capacity. The finding of increased glycolysis is in agreement with previous studies demonstrating higher activity of phosphofructokinase, the rate limiting enzyme of glycolysis, in skeletal muscles from individuals with higher resting energy expenditure (Zurlo et al., 1994).

Higher levels of glycolysis likely explains the negative correlation between resting energy expenditure and glycolytic reserve. High levels of glycolysis indicate that the glycolytic activity of a cell is close to the theoretical maximum, and therefore, there is reduced capacity to respond to increased energy demand through this pathway. The negative correlation between resting energy expenditure and glucose oxidation capacity is paradoxical, given that glucose metabolism provides more ATP when compared to oxygen consumption when compared to glucose metabolism (Turner et al., 2014). As such, an increase in the need for fatty acid oxidation in myotubes obtained from patients with higher resting energy expenditure may function to increase ATP availability. Moreover, the correlation observed between resting energy expenditure and fatty acid oxidation dependence aligns with physiological processes whereby continued dependence on fatty acid metabolism requires greater levels of oxygen consumption when compared to glucose metabolism (Turner et al., 2014).

We found that myotubes with higher fatty acid oxidation dependence were derived from patients with slower clinical decline, indicating that substrate availability and/ or use could be a factor that determines disease progression in ALS. Previous studies linking higher levels of serum lipids and fat mass with longer survival (Marin et al., 2011; Lindauer et al., 2013; Huang et al., 2015) provide some evidence to suggest that lipids are beneficial in ALS. This is further supported by recent clinical observations demonstrating that high fat dietary supplementation is able to slow disease progression and extend survival in patients with more rapid progressing disease (Ludolph et al., 2020).

The cause for the change in fuel substrate preference towards fatty acids in skeletal muscle remains unknown. An interesting observation in this study was the functional change in fuel oxidation preference at the symptomatic stage of disease in SOD1G93A mice—this finding is in contrast to work published by Palamiuc et al. who identified gene expression changes suggestive of a similar preference for fat oxidation in asymptomatic SOD1G86R mice (Palamiuc et al., 2015). The differences observed between the two SOD1 mouse models are curious and could be due to the more aggressive disease course in SOD1G86R mice. It is plausible that the timing in the change in preference for fatty acid oxidation is associated with alterations in global metabolic homeostasis (which are...
apparent in asymptomatic SOD1\textsuperscript{G86R} mice) that results in cellular bioenergetic deficit (Ngo and Steyn, 2015; Loeffler et al., 2016; Vandoorne et al., 2018), and this in turn might subsequently lead to the utilization of fatty acids as an initial adaptive response to yield more ATP than glucose to sustain cellular function and survival. As disease progresses, persistent fatty acid oxidation would likely become maladaptive, with increased production of reactive oxygen species and lipotoxicity exacerbating the degenerative process (Aon et al., 2014; Tracey et al., 2018). If this were the case, could modulation of glucose–fatty acid flux offer therapeutic benefit in ALS? It has previously been shown that dichloroacetate, a synthetic halogenated organic acid that promotes glucose oxidation and pyruvate oxidation, improves function and survival in SOD1 mouse models of ALS (Miquel et al., 2012; Palamiuc et al., 2015). Moreover, the direct targeting of glycolysis or glycolysis associated pathways in skeletal muscle might also show promise. This approach is supported by observations that promotion of glycolytic function in motor neurons improves locomotor function in a transactive response DNA binding protein 43 (TDP-43) fly model of ALS (Manzo et al., 2019), and that pre-supplementation of chromosome 9 open reading frame 72 (C9orf72) induced astrocytes with inosine improves survival of motor neurons by increasing glycolytic flux (Allen et al., 2019).

The biochemical mechanisms that underpin the increased dependence for fatty acid oxidation in skeletal muscle also remain to be determined. It is well known that substrate competition between glucose and fatty acids at the mitochondrial outer membrane is dictated by the activities of the pyruvate dehydrogenase complex and the carnitine palmitoyl transferase system/complexes, and that the interaction between these two complexes is influenced by nutrient availability (Sugden and Holness, 1994). In ALS, an increase in energy expenditure that outstrips nutrient supply might result in negative energy balance, and a switch in substrate preference from glucose to fatty acids through altered pyruvate dehydrogenase and carnitine palmitoyl transferase activity. Future, in-depth and longitudinal assessment of the complex interactions between glucose and fatty acid metabolism pathways is needed to decipher the mechanisms that modulate the adaptation of ALS skeletal muscle towards higher fatty acid oxidation.

Although we have used both mouse and human-derived models to investigate aspects of muscle metabolism in ALS, there are some limitations to our study. First, the SOD1\textsuperscript{G93A} mouse, albeit an accepted pre-clinical model of ALS, is representative of only a small proportion of ALS patients. However, we found similar alterations in substrate oxidation in the myotubes derived from some ALS patients, indicating that muscle metabolic changes are possibly present widely in the disease. Second, we matched measures in SOD1\textsuperscript{G93A} mice with measures in WT control mice rather than measures in mice overexpressing wild-type human SOD1 (WTSOD1). Similar to mutant human SOD1\textsuperscript{G93A}, the overexpression of WTSOD1 in motor neuron-like cells is associated with altered glycolytic function (Richardson et al., 2013; Valbuena et al., 2016), although, to our knowledge, the impact of WTSOD1 on fatty acid metabolism remains to be investigated. Thus, detailed metabolic characterization of WTSOD1 mice is needed if we are to generate further insight into the degree to which mutant human SOD1 alters glucose–fatty acid flux when compared to WTSOD1. Third, while we conducted assessment of energy expenditure in ALS patients using indirect calorimetry (Haugen et al., 2007), and assessed substrate oxidation in real-time using Seahorse technology, our use of primary human myotubes is a caveat. Previously, it has been shown that human primary myotubes mirror the metabolic phenotypes of their donors (Ukropcova et al., 2005). Regardless, as primary myotubes are grown from cells, which have been isolated from muscle biopsies that were removed from their physiological milieu, future studies assessing substrate oxidation in muscle in vivo are needed.

In summary, we demonstrate that an increase in fatty acid oxidation dependence and capacity occurs in glycolytic muscle of SOD1\textsuperscript{G93A} mice during the symptomatic stages of disease when they exhibit increased energy expenditure. In myotubes derived from ALS patients, a similar increase in fatty acid oxidation dependence occurs. While this change in fatty acid oxidation appears to be associated with the progression of disease, it is not linked to hypermetabolism in human ALS. Given the heterogeneity of disease in ALS, there remains an important need for further studies that delineate mechanisms of metabolic imbalance and the link between substrate utilization and energy expenditure throughout the course of the disease. A comprehensive understanding of specific metabolic changes at an individual patient level will be essential for the development of treatments that aim to target metabolic pathways in ALS.

**Supplementary material**

Supplementary material is available at *Brain Communications* online.

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Competing interests

The authors have no conflicts of interest to declares.

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