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Authors	Burns, David P.;Drummond, Sarah E.;Bolger, Dearbhla;Coiscaud, Amélie;Murphy, Kevin H.;Edge, Deirdre;O'Halloran, Ken D.
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## Article

# N-acetylcysteine Decreases Fibrosis and Increases Force-Generating Capacity of *mdx* Diaphragm

David P. Burns <sup>1,\*</sup>, Sarah E. Drummond <sup>1</sup>, Dearbhla Bolger <sup>2</sup>, Amélie Coiscaud <sup>1</sup>, Kevin H. Murphy <sup>1</sup>, Deirdre Edge <sup>2</sup> and Ken D. O'Halloran <sup>1</sup>

<sup>1</sup> Department of Physiology, School of Medicine, College of Medicine and Health, University College Cork, Cork, Ireland, T12 XF62 ; s.drummond@umail.ucc.ie (S.E.D.); a.coiscaud@stanbiotec.com (A.C.); KevinMurphy12@umail.ucc.ie (K.H.M.); k.ohalloran@ucc.ie (K.D.O.)

<sup>2</sup> Department of Physiology, School of Medicine, Trinity Biomedical Sciences Institute, Trinity College Dublin, the University of Dublin, Dublin, Ireland, D02 R590; bolgerde@tcd.ie (D.B.); ded@kleresca.com (D.E.)

\* Correspondence: david.burns@ucc.ie; Tel.: +353-21-4205486

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**Abstract:** Respiratory muscle weakness occurs due to dystrophin deficiency in Duchenne muscular dystrophy (DMD). The *mdx* mouse model of DMD shows evidence of impaired respiratory muscle performance with attendant inflammation and oxidative stress. We examined the effects of N-acetylcysteine (NAC) supplementation on respiratory system performance in *mdx* mice. Eight-week-old male wild type ( $n = 10$ ) and *mdx* ( $n = 20$ ) mice were studied; a subset of *mdx* ( $n = 10$ ) received 1% NAC in the drinking water for 14 days. We assessed breathing, diaphragm, and external intercostal electromyogram (EMG) activities and inspiratory pressure during ventilatory and non-ventilatory behaviours. Diaphragm muscle structure and function, cytokine concentrations, glutathione status, and mRNA expression were determined. Diaphragm force-generating capacity was impaired in *mdx* compared with wild type. Diaphragm muscle remodelling was observed in *mdx*, characterized by increased muscle fibrosis, immune cell infiltration, and central myonucleation. NAC supplementation rescued *mdx* diaphragm function. Collagen content and immune cell infiltration were decreased in *mdx* + NAC compared with *mdx* diaphragms. The cytokines IL-1 $\beta$ , IL-6 and KC/GRO were increased in *mdx* plasma and diaphragm compared with wild type; NAC decreased systemic IL-1 $\beta$  and KC/GRO concentrations in *mdx* mice. We reveal that NAC treatment improved *mdx* diaphragm force-generating capacity associated with beneficial anti-inflammatory and anti-fibrotic effects. These data support the potential use of NAC as an adjunctive therapy in human dystrophinopathies.

**Keywords:** antioxidant; N-acetylcysteine; DMD; *mdx*; diaphragm; intercostal; interleukin-6; interleukin-1 $\beta$ ; fibrosis

## 1. Introduction

Skeletal muscle weakness is a major feature of Duchenne muscular dystrophy (DMD) due to a deficiency in the protein dystrophin [1]. In healthy muscle, dystrophin has a structural role in supporting muscle fibres during repeated cycles of muscle contraction, acting to limit the mechanical stress applied to muscle fibres [2,3]. In the absence of dystrophin, such as in DMD, skeletal muscle fibres have increased fragility, resulting in fibre damage. Common pathological features of dystrophic muscle include myonecrosis, inflammation, and oxidative stress [4,5]. Due to profound muscle weakness and damage, children with DMD have difficulty with ambulation in their early years, necessitating assistive devices.

Respiratory system morbidity occurs in DMD due to weakness of the respiratory musculature [6], the end effectors of breathing. Respiratory function is compromised in boys with DMD, with reductions in forced vital capacity [7,8] and impaired inspiratory muscle strength [6,9], evident from a young age. Impaired respiratory function is most prominent during sleep, evidenced by an increased risk of sleep disordered breathing in children with DMD [10]. Many patients require ventilatory support during sleep to maintain blood gas homeostasis. Cough and airway clearance manoeuvres are compromised in DMD due to reduced respiratory muscle strength, such that patients are more susceptible to aspiration pneumonia [11]. Therapies aimed at improving respiratory muscle strength in DMD are necessary and attractive, in order to maintain adequate ventilation and prevent respiratory failure, a leading cause of mortality in DMD, notwithstanding recent advances in the respiratory support of DMD boys.

Studies of the respiratory system in the *mdx* mouse, a preclinical model of DMD, have documented profound diaphragm muscle weakness and structural remodelling from a young age as a consequence of dystrophin deficiency [12–16]. Inflammatory markers such as immune cell infiltration and cytokine concentrations are increased in *mdx* diaphragm, as well as the abundance of collagen deposits [17]. Moreover, indices of oxidative stress including lipid peroxidation and superoxide levels are elevated in *mdx* diaphragm compared with control muscle [18]. Inflammation and high levels of reactive oxygen species (ROS) can culminate in skeletal muscle damage leading to poor physiological performance [19]. Oxidative stress is a recognized feature of respiratory disorders including DMD. Targeting oxidative stress within muscle by reducing the bioavailability of ROS or boosting endogenous antioxidant stores are attractive adjunctive therapies, particularly in conditions where redox imbalance presents and contributes to muscle pathology [20,21]. We have previously demonstrated that administration of a superoxide scavenger (Tempol) to *mdx* mice for two weeks restores metabolic enzyme activities and improves diaphragm muscle force-generating capacity [22]. It has been shown by others that Tempol supplementation reduces myonecrosis and inflammation in the diaphragm and biceps brachii muscles of *mdx* mice [23].

N-acetylcysteine (NAC) is a dietary antioxidant and precursor to glutathione, an endogenous antioxidant, safe for use in humans. Interestingly, NAC is a mucolytic agent and is commonly used in patients with cystic fibrosis and chronic obstructive pulmonary disease. Previous studies from our group have demonstrated beneficial effects of NAC supplementation on respiratory muscle function in animal models of respiratory disease [24–26]. Studies utilising NAC as a potential therapeutic for dystrophic disease have yielded promising results. Pinniger et al. (2017) reported improved normalized grip strength and extensor digitorum longus (EDL) force in *mdx* mice supplemented with 2% NAC in the drinking water for 6 weeks [27]. In a separate study, intraperitoneal injections of NAC in 14 day old *mdx* mice for 14 days reduced tumour necrosis factor- $\alpha$  (TNF- $\alpha$ ) and lipid peroxidation levels in *mdx* diaphragm [28]. Terrill et al. (2012) reported that NAC administered in the drinking water (1% NAC for 6 weeks or 4% NAC for one week) prevented exercise-induced myonecrosis in quadriceps muscle of *mdx* mice [29]. Studies by Whitehead et al. (2008) determined that 1% NAC in the drinking water for 6 weeks reduced the concentration of ROS and decreased damage in EDL muscle of *mdx* mice [30]. Collectively, these studies support the use of NAC to target muscle damage mediated by oxidative stress in *mdx* mice, but no studies to date have assessed the efficacy of NAC in ameliorating respiratory system deficits in *mdx* mice.

In the current study, we set out to perform a broad and thorough assessment of the effects of NAC supplementation on respiratory system performance in young (8-week-old), male *mdx* mice. Six-week-old *mdx* mice were treated with 1% NAC in the drinking water for 14 days. We hypothesized that NAC would have beneficial effects on dystrophic respiratory muscle, leading to preserved respiratory system performance.

## 2. Materials and Methods

### 2.1. Ethical Approval

Procedures on live animals were performed under licence in accordance with Irish and European directive 2010/63/EU following ethical approval by University College Cork (AEEC no. 2013/035). Experiments were carried out in accordance with guidelines laid down by University College Cork's Animal Welfare Body, and conform to the principles and regulations described by [31].

### 2.2. Experimental Animals and N-acetylcysteine (NAC) Treatment

Male wild type (WT; C57BL/10ScSnJ;  $n = 10$ ) and *mdx* (C57BL/10ScSn-Dmd<sup>mdx</sup>/J;  $n = 20$ ) mice were purchased from the Jackson Laboratory (Bar Harbor, ME, USA) and were housed at University College Cork's animal facility. Animals were housed conventionally in temperature- and humidity-controlled rooms, operating on a 12 h light:12 h dark cycle with food and water available *ad libitum*. Mice were assigned to three different experimental groups: wild type, *mdx* and *mdx* + NAC. The *mdx* + NAC group received 1% N-acetylcysteine (Sigma-Aldrich, Wicklow, Ireland) in the drinking water for 14 days, beginning at six weeks of age. Drinking water containing NAC was pH matched to control water (pH 8). Mice were studied at eight weeks of age. We performed a thorough assessment of respiratory performance, with measurements of breathing and ventilatory capacity in response to chemoactivation, recordings of thoracic inspiratory pressure and respiratory muscle electromyography (EMG) in vivo and respiratory muscle function tests ex vivo. Diaphragm muscle tissue was collected for structural analysis using histological and immunofluorescence techniques. Diaphragm glutathione status and mRNA expression were examined. Cytokine concentrations were determined in diaphragm and plasma samples.

### 2.3. Whole-Body Plethysmography

Whole-body plethysmography was used to assess respiratory flow in unrestrained, unanaesthetized mice. Mice were introduced into plethysmograph chambers (Model PLY4211; volume 600 mL, Buxco Research Systems, Wilmington, NC, USA) and were allowed an acclimation period typically lasting 40–60 min, with room air passing through each chamber ( $1\text{ L}\cdot\text{min}^{-1}$ ).

Experimental protocol: Following acclimation, during confirmed periods of quiet rest, a 20-min baseline recording was performed in normoxia. This was followed by central and peripheral chemoreceptor stimulation with hypercapnic-hypoxia (10% O<sub>2</sub> & 6% CO<sub>2</sub>) for 5 min, to examine maximum chemoactivated breathing. Respiratory parameters including respiratory frequency ( $f_R$ ), tidal volume ( $V_T$ ) and minute ventilation ( $V_E$ ) were recorded on a breath-by-breath basis for analysis offline.

Data analysis: Baseline normoxic ventilation was determined as an average of the baseline period. For hypercapnic-hypoxia, ventilatory measurements were taken during the final (5th) minute of the challenge.  $V_T$  and  $V_E$  were normalized for body mass (g).

### 2.4. Diaphragm and External Intercostal EMG and Oesophageal Pressure Recordings

Mice were anaesthetized with 5% isoflurane in 60% O<sub>2</sub> (balance N<sub>2</sub>) followed by urethane ( $1.7\text{ g}\cdot\text{kg}^{-1}$  i.p.). Wild type ( $n = 10$ ), *mdx* ( $n = 10$ ) and *mdx* + NAC ( $n = 10$ ) mice were then placed in the supine position and were gradually weaned from the isoflurane. Body temperature was maintained at 37 °C via a rectal probe and thermostatically-controlled heating blanket (Harvard Apparatus, Holliston, MA, USA) for the duration of the experiment. Supplemental anaesthetic was administered as necessary to maintain a surgical plane of anaesthesia. Depth of anaesthesia was determined by assessment of pedal withdrawal reflex to noxious pinch. Peripheral capillary O<sub>2</sub> saturation (SpO<sub>2</sub>) was monitored using a pulse oximeter clip (MouseOx™, Starr Life Sciences Corporation, Oakmount, PA, USA). A thigh of each mouse was shaved and the pulse oximeter was attached. A tracheotomy was performed in the mid-cervical region. A bias flow of supplemental O<sub>2</sub> (FiO<sub>2</sub> = 0.60) was provided

to all animals during baseline conditions. End-tidal carbon dioxide (ETCO<sub>2</sub>) was measured using a MicroCapStar (CWE, Ardmore, PA, USA). A pressure-tip catheter (Mikro-Tip, Millar Inc., Houston, TX, USA) was inserted into the mouth and positioned in the thoracic oesophagus for the assessment of oesophageal pressure, an index of intrapleural sub-atmospheric pressure generated by the respiratory musculature during inspiration [13]. Oesophageal recordings displayed phasic sub-atmospheric pressure swings during inspiration. For the measurement of diaphragm EMG, a concentric needle electrode (26G; Natus Manufacturing Ltd., Galway, Ireland) was inserted into the costal diaphragm for the continuous measurement of EMG activity. In a similar manner, a concentric needle electrode was inserted into the third intercostal space (T3) for the measurement of external intercostal EMG activity. EMG signals were amplified ( $\times 5000$ ), band-pass filtered (50–5000 Hz) and integrated (50 ms time constant; Neurolog system, Digitimer Ltd., Welwyn Garden City, UK). All biological signals were passed through an analogue-to-digital converter (Powerlab r8/30; ADInstruments, Colorado Springs, CO, USA) and were acquired using LabChart 7 (ADInstruments) at a sampling frequency of 4 kHz.

**Experimental protocol:** Following instrumentation, animals were allowed a period of stabilization for at least 10 min, prior to a 10 min baseline recording. Next, the trachea was occluded for 30–40 s until a plateau was observed in the inspiratory pressure recordings. Following this challenge, animals were allowed to recover and were then instrumented for the measurement of ETCO<sub>2</sub> and tracheal airflow. Next, animals were vagotomized by bilateral section of the cervical vagi. Following vagotomy, respiratory parameters were recorded for 10 min under steady-state conditions following vagotomy. Next, animals were challenged with hypercapnic-hypoxia (15% O<sub>2</sub> & 6% CO<sub>2</sub>; 3 min) to examine the effects of chemostimulation on tracheal airflow. Following the experimental protocol, mice were euthanized by decapitation. Blood was collected in 1.5 mL Eppendorf tubes coated with 4% EDTA, which were centrifuged at 1,610 *g* for 10 min at 4 °C. Plasma samples were aliquoted and snap frozen in liquid nitrogen and stored at –80 °C. Tibia length and mass were examined as an index of somatic growth. The following organs were weighed (wet weight): spleen, kidney, lung, right heart ventricle and left heart ventricle and interscapular brown adipose tissue and epididymal white adipose tissue.

**Data analysis:** Integrated inspiratory diaphragm and T3 external intercostal (EIC) EMG activity and peak inspiratory sub-atmospheric oesophageal pressure were analysed. The amplitudes of diaphragm and EIC EMG and oesophageal pressure were averaged under steady-state basal conditions and averaged for the 5 successive maximal sustained efforts (maximal response) of the airway occlusion challenge.  $V_T$  was derived from tracheal airflow measurements following vagotomy during baseline and during hypercapnic-hypoxia. Data for oesophageal pressure and respiratory muscle EMGs in wild type and *mdx* mice were previously published in a separate report [13].

### 2.5. Ex Vivo Diaphragm Muscle Function

The diaphragm muscle (rib and central tendon intact) was immediately excised and placed in a tissue bath at room temperature containing continuously gassed hyperoxic (95% O<sub>2</sub>/5% CO<sub>2</sub>) Krebs solution (in mM: NaCl, 120; KCl, 5; Ca<sup>2+</sup> gluconate, 2.5; MgSO<sub>4</sub>, 1.2; NaH<sub>2</sub>PO<sub>4</sub>, 1.2; NaHCO<sub>3</sub>, 25; and glucose, 11.5) and *d*-tubocurarine (25  $\mu$ M) prior to functional analysis. Diaphragm muscle preparations were suspended vertically between two platinum plate electrodes in a water-jacketed tissue bath at 35 °C containing Krebs solution and were continuously aerated with hyperoxia (95% O<sub>2</sub> & 5% CO<sub>2</sub>). The rib was sutured to an immobile hook and remained in a fixed position for the duration of the experiment. Using non-elastic string, the central tendon was attached to a lever connected to a dual-mode force transducer (Aurora Scientific Inc.; Aurora, ON, Canada). To determine muscle optimum length ( $L_o$ ), the length of the muscle preparations were adjusted using a micro-positioner between intermittent twitch contractions [32,33]. The muscle length which revealed maximal isometric twitch force for a single isometric twitch stimulation (supramaximal stimulation, 1 ms duration) was considered  $L_o$ . Diaphragm preparations were maintained at  $L_o$  for the duration of the protocol.

Experimental protocol: First, a single isometric twitch contraction was measured and peak isometric twitch force ( $P_t$ ), contraction time (CT) and half-relaxation time ( $\frac{1}{2}$  RT) were assessed. To examine the force-frequency relationship, muscle bundles were stimulated sequentially at 10, 20, 40, 60, 80, 100, 120, 140 and 160 Hz (300 ms train duration). Contractions were interspersed by a 1 min interval. Following the isometric protocol, an isotonic contraction was elicited in preparations at 0% load to examine maximum unloaded muscle shortening and velocity of shortening [32,33].

Data analysis: Muscle bundle cross-sectional area (CSA) was determined for the purpose of normalizing muscle force to bundle size. CSA was calculated by dividing muscle mass (weight in grams) by the product of muscle  $L_0$  (cm) and muscle density (assumed to be  $1.06 \text{ g.cm}^{-3}$ ). Muscle force was divided by bundle CSA and expressed as specific force ( $\text{N.cm}^{-2}$ ). CT and  $\frac{1}{2}$  RT were measured as indices of isometric twitch kinetics and expressed in ms. For isotonic contractions, total muscle shortening was considered the total distance shortened during muscle contraction at 0% load. For total muscle shortening ( $S_{\max}$ ), data were expressed in absolute units (cm) and were further normalized to  $L_0$  and expressed in  $L_0^{-1}$ . For the measurement of muscle shortening velocity ( $V_{\max}$ ), the distance shortened during the initial 30 ms of an unloaded contraction was assessed [25,33,34] and determined in absolute units ( $\text{cm.s}^{-1}$ ) and was normalized to  $L_0$  and expressed in  $L_0.s^{-1}$ .

## 2.6. Muscle Immunofluorescence and Histology

### 2.6.1. Tissue Preparation

Sections of hemidiaphragm from wild type ( $n = 6$ ), *mdx* ( $n = 7$ ) and *mdx* + NAC-treated mice ( $n = 4$ ) were mounted on cubes of liver. Diaphragm samples were embedded in optimum cutting temperature embedding medium (OCT; VWR International, Dublin, Ireland) for cryoprotection and then frozen in isopentane (Sigma Aldrich, Wicklow, Ireland) cooled on dry ice. Samples were then stored at  $-80^\circ\text{C}$  for subsequent structural analysis. Serial transverse muscle sections ( $10 \mu\text{m}$ ) were cut using a cryostat (Leica CM3050; Leica Microsystems, Nussloch, Germany) at  $-22^\circ\text{C}$  and mounted across polylysine-coated glass slides (VWR International, Dublin, Ireland) allowing for a distribution of tissue on a given slide.

### 2.6.2. Histological analysis

To examine putative inflammatory cell infiltration of muscle fibres, tissue sections were stained with haematoxylin and eosin (H&E) using an autostainer (Leica ST5010 Autostainer XL, Leica Microsystems, Nussloch, Germany). For collagen staining, picro-sirius red (Leica Biosystems, Wetzlar, Germany) staining was completed. Slides were mounted using DPX mounting medium (Sigma-Aldrich, Wicklow, Ireland), air-dried and visualized on a bright field microscope (Olympus BX51) at  $\times 10$  magnification.

Data analysis: A total of six tissue sections per animal were examined. Muscle histology was scored using ImageJ software. Putative inflammatory cell infiltration (the presence of cells in the extracellular matrix) was scored and expressed as a percentage of the total area of muscle. For slides stained with picro-sirius red, the microscope lighting exposure was standardized during imaging. Images were analysed using a colour balance threshold and the area of collagen was expressed as a percentage of the total area of muscle. Data generated from multiple images was averaged per animal before computing group means.

### 2.6.3. Embryonic Myosin Heavy Chain and Laminin Immunofluorescence

Embryonic myosin ( $\text{MyHC}_{\text{EMB}}$ ) immunofluorescence was carried out as a marker of fibre regeneration/remodelling [35]. Laminin immunofluorescence was used to examine the fibre size (minimum Feret's diameter) distribution of the diaphragm muscle as well as to determine central nucleation of muscle fibres. Two slides were removed from the  $-80^\circ\text{C}$  freezer containing a minimum of four sections per slide, from two distinct regions of muscle tissue for each animal. A hydrophobic barrier was created around the individual tissue samples using a hydrophobic pen (ImmEdge TM Vector Labs, Peterborough, UK). Briefly, tissue samples were immersed in phosphate-buffered saline

(PBS) (0.01 M) containing 1% bovine serum albumin (BSA) for 15 min followed by  $3 \times 5$  min PBS rinses. This was followed by a 30 min wash in PBS containing 5% goat serum (Sigma-Aldrich, Wicklow, Ireland). Before application of the primary embryonic mouse monoclonal antibody (developed by S. Schiaffino and obtained from the Developmental Studies Hybridoma Bank (DSHB) at the University of Iowa, IA, USA), slides were exposed to a mouse IgG blocker, M.O.M. blocking solution (Vector Lab Inc., Burlingame, CA, USA) for 1 h at room temperature. Following  $3 \times 5$  min PBS washes, a cocktail of primary antibodies containing MyHC<sub>EMB</sub> (F1.652; 1:20) and rabbit anti-laminin antibody (Sigma Aldrich, 1:500) diluted in 1% BSA in PBS was applied to the slides and incubated at 4 °C overnight in a humidity chamber. The next day, slides were washed for  $3 \times 5$  min in PBS before the application of the corresponding secondary antibodies, AlexaFluor594-conjugated goat anti-mouse IgG<sub>1</sub> (1:500, Jackson ImmunoResearch Ltd., Ely, UK) and fluorescein isothiocyanate (FITC)-conjugated goat anti-rabbit secondary antibody (1:250, Sigma Aldrich). Slides were incubated for 1 h in the dark at room temperature. Finally, slides were washed for  $3 \times 5$  min rinses in PBS before the application of 4',6-diamidino-2-phenylindole (DAPI; 1:4000, Sigma Aldrich) for 10 min to identify myonuclei, followed by a final  $3 \times 5$  min rinse in PBS. Negative control experiments were run in parallel where the primary antibodies were omitted and the secondary antibodies were applied to ensure that immune-tagging was specific.

**Data analysis:** For muscle fibre distribution, MyHC<sub>EMB</sub> expressing muscle fibres and central nucleation, muscle sections were viewed at  $\times 10$  magnification and images were captured using an Olympus BX51 microscope and an Olympus DP71 camera. Cell Sens<sup>TM</sup> (Olympus) was used to digitally capture the images. Images were taken at random areas across each muscle section with four muscle sections analysed per animal. Individual images of the laminin, DAPI and MyHC<sub>EMB</sub> fibres were captured and merged together using ImageJ software. A  $600 \mu\text{m} \times 600 \mu\text{m}$  grid with inclusion and exclusion borders was placed over the image for analysis [12]. From each muscle section, the number of fibres per area along with minimum Feret's diameter of the individual muscle fibres was analysed using a specialized ImageJ macro [36]. Frequency histograms were constructed to illustrate the distribution of fibre size in muscle sections across groups in addition to the coefficient of variation of minimum Feret's diameter. The number of centrally nucleated and MyHC<sub>EMB</sub> expressing muscle fibres were counted (using the ImageJ multi-point tool) and expressed relative to the total number of myofibres per recorded area. Data generated from multiple images were averaged per animal before computing group means.

## 2.7. Assays

### 2.7.1. Tissue Preparation

Frozen diaphragm muscle samples were removed from storage at  $-80$  °C, weighed and homogenized on ice in ice-cold modified radioimmunoprecipitation assay (1X RIPA) buffer, deionized H<sub>2</sub>O, 200 mM sodium fluoride (NaF), 100 mM phenylmethylsulfonylfluoride (PMSF), 1X protease cocktail inhibitor (Sigma Aldrich, Ireland) and 200 mM sodium orthovanadate (Na<sub>ortho</sub>) at a 5% *w/v* ratio using a general laboratory homogenizer (Omni-Inc., Kennesaw, GA, USA). Homogenates were allowed 20 min of lyse time on ice with intermittent vortexing. Homogenates were centrifuged in a U-320R centrifuge (Boeckel & Co., Hamburg, Germany) at  $15,366 g$  at 4 °C for 20 min to separate insoluble cellular fractions from protein homogenates. The protein-containing supernatant was removed and stored at  $-80$  °C, and the pellet was discarded. The protein concentration of each sample was quantified using a bicinchoninic acid (BCA) assay (Pierce Biotechnology, Fisher Scientific, Dublin, Ireland) as per the manufacturer's instructions, at a dilution of 1:10.

### 2.7.2. Glutathione Reductase and Glutathione Peroxidase Activities, Total Glutathione (tGSH) Concentration and GSSG:GSH Ratio

Glutathione reductase (GR) catalyzes the reduction of glutathione disulfide (GSSG) by nicotinamide adenine dinucleotide phosphate (NADPH) to produce reduced glutathione (GSH). GR

activity was measured in diaphragm muscle homogenates from all three groups: WT ( $n = 7$ ), *mdx* ( $n = 7$ ) and *mdx* + NAC ( $n = 7$ ). Briefly, 50  $\mu$ L (25  $\mu$ g) of sample, 50  $\mu$ L of  $\text{KH}_2\text{PO}_4$  (0.4 M)/EDTA (4 mM; pH 7.5) buffer and 50  $\mu$ L of oxidised GSSG (4 mM) was added in duplicate to wells of a clear 96-well plate. Controls were also plated in duplicate which included: (1) all assay components except oxidised GSSG and (2) all assay components except sample, to account for any non-specific reduction in samples tested. 50  $\mu$ L of NADPH (0.4 mM) was quickly added to each well to initiate the reaction and the absorbance was measured at 340 nm every 15 s for 30 min at 37 °C. The oxidation of NADPH at this wavelength is reflected by a decrease in absorbance over time, proportional to GSSG reduction and thus GR activity. Data are expressed as mU  $\text{mg}^{-1}$  protein. Glutathione peroxidase (GPx) converts GSH to GSSG while also reducing lipid hydroperoxides to alcohols, and hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) to water. GPx activity was measured in diaphragm muscle homogenates from WT ( $n = 7$ ), *mdx* ( $n = 7$ ) and *mdx* + NAC ( $n = 7$ ) groups using a commercially available kit (catalogue number. ABE3639; Source BioScience, Nottingham, UK) as per the manufacturer's instructions. Data are expressed as nmol  $\text{min}^{-1}$   $\text{mg}^{-1}$  protein. Total glutathione (tGSH) comprises GSH + GSSG, which can be used to calculate the GSSG:GSH ratio, a common measure of oxidative stress. tGSH and GSSG concentration were measured in diaphragm muscle homogenates from wild type ( $n = 7$ ), *mdx* ( $n = 7$ ) and *mdx* + NAC ( $n = 7$ ) groups using a commercially available kit (catalogue number. 703002; Cayman Chemical, Ann Arbor, MI, USA) as per the manufacturer's instructions, with data expressed as  $\mu\text{M}$   $\text{mg}^{-1}$  protein and GSSG:GSH.

### 2.7.3. Cytokine Multiplex Assay

A multiplex cytokine assay (K15048G-1; Meso Scale Discovery, Rockville, MD, USA) was used to examine cytokine concentrations in diaphragm muscle (200  $\mu\text{g}$   $\text{well}^{-1}$ ;  $n = 7$  per group) and plasma samples (1:2 dilution;  $n = 8$ –10 per group) from all 3 groups. The assay was performed according to the manufacturer's instructions using an extended incubation time to improve the detection of cytokines in diaphragm muscle homogenates (the plate was incubated overnight at 4 °C) [17]. Following incubation, the plate was read on a Quickplex SQ 120 imager (Meso Scale Discovery).

## 2.8. Gene Expression

### 2.8.1. RNA Extraction and Preparation

Diaphragm muscle samples were removed from storage at  $-80$  °C, weighed and samples ranging from 20–50 mg were homogenized on ice in Tripure Isolation Reagent (Roche Diagnostics, Ltd., Burgess Hill, UK) using a general laboratory homogenizer (Omni-Inc., Kennesaw, GA, USA). Total RNA was isolated from homogenates in accordance with the manufacturer's instructions, with an additional chloroform wash step performed during phase separation. The quantity ( $\text{ng}$   $\mu\text{L}^{-1}$ ) and purity of isolated RNA (260:280 & 260:230 ratios) was assessed using a Nanodrop 1000 (Thermo Scientific, Waltham, MA, USA) and spectrophotometry. The integrity of isolated RNA was assessed by visualization of distinct 18S and 28S ribosomal RNA bands using an agarose gel electrophoresis system (E-gel, Life Technologies, Carlsbad, CA, USA).

### 2.8.2. Reverse Transcription

Diaphragm muscle RNA was reverse transcribed to cDNA using a Transcriptor First Strand cDNA Synthesis Kit (Roche Diagnostics Ltd., Burgess Hill, UK) in accordance with the manufacturer's instructions.

### 2.8.3. qRT-PCR

cDNA was amplified using Realtime ready Catalog or Custom assays (Roche Diagnostics Ltd., Burgess Hill, UK) shown in Table 1 and Fast Start Essential Probe Master (Roche Diagnostics Ltd., Burgess Hill, UK) as per the manufacturer's instructions. Briefly, all reactions were carried out in duplicate on a 96-well plate and consisted of 5  $\mu\text{L}$  cDNA and 15  $\mu\text{L}$  master mix using the Lightcycler



96 (Roche Diagnostics Ltd., Burgess Hill, UK). RNA negatives, reverse transcription negatives, cDNA negatives (no template controls) and plate calibrators were used on every plate. Quantification cycle values (Cq) obtained from experiments for the genes of interest (*Nfkb1*, *Fbxo32*, *Trim63*, *Map1lc3b*, *Gabarapl1*, *Bnip3*, *Cybb*, *Park2*, *Pink*, *Nfe2l2*, *Cat*, *Mstn*, *Myod1*, *Gpx2*, *Myog*, *Sod2*, *Sod1*, *Sirt1*, *Igf1* and *Mef2c*) were normalized to that of a reference gene, *Hprt1*, to account for variations in input amounts of RNA/cDNA and the efficiency of reverse transcription. *Hprt1* was found to be the most stable gene of the candidate reference genes screened, in consideration of genotype and drug treatment (NAC). The relative gene expression was calculated using the  $\Delta\Delta CT$  method (normalized expression of the gene of interest to that of the reference gene), with a change in expression shown as a fold change relative to the control group (wild type).

**Table 1.** Real-time ready catalog and custom assays used for cDNA amplification.

Gene Title	Gene Symbol	Assay ID
Atrogin1	Fbxo32	317844
BNIP3	Bnip3	311465
Catalase	Cat	310718
GABARAPL1	Gabarapl1	317923
GPX2	Gpx2	318630
HPRT	Hprt	307879
IGF1	Igf1	313359
LC3B	Map1lc3b	317920
MEF2C	Mef2c	318629
Myogenin	Myog	313501
Myostatin	Mstn	318626
MyoD	Myod1	313570
MuRF1	Trim63	317843
NFκB	Nfkb1	300085
NOX2	Cybb	317885
Nrf2	Nfe2l2	313377
PARK2	Park2	317264
PINK1	Pink1	331846
Sirtuin1	Sirt1	310480
SOD1	Sod1	310738
SOD2	Sod2	310295

## 2.9. Statistical Analysis

Data are expressed as mean  $\pm$  SD or as box and whisker plots (median, 25–75th percentile and scatter plot). Using Prism, version 6.0 (Graphpad Software Inc., San Diego, CA, USA), data from each group were statistically analysed and assessed for normal distribution and equal variances. Data sets that were of equal variance and normally distributed were compared statistically using one-way analysis of variance (ANOVA) with a Newman–Keuls *post hoc* test. Kruskal–Wallis with Dunn’s *post hoc* test was applied to group data that were not of Gaussian distribution. Diaphragm muscle force–frequency relationship was statistically analysed using repeated measures two-way ANOVA with a Bonferroni *post hoc* test. *P* values are reported.

## 3. Results

### 3.1. Body Mass and Organ Measurements

Table 2 compares body mass, tibia length and mass, organ mass and brown and white adipose mass in wild type, *mdx* and *mdx* mice treated with 1% NAC (*mdx* + NAC). Body mass for *mdx* mice was unaffected by NAC supplementation ( $P > 0.05$ ; Kruskal–Wallis with Dunn’s *post hoc* test). Right

and left ventricle mass was increased for *mdx* mice ( $P < 0.0001$  and  $P < 0.01$ ; one-way ANOVA with Newman-Keuls *post hoc* test and Kruskal–Wallis with Dunn’s *post hoc* test; right and left ventricle, respectively). The ratio of right to left ventricle mass (RV:LV) was increased in *mdx* ( $P < 0.05$ ; one-way ANOVA with Newman–Keuls *post hoc* test); RV:LV was unaffected by NAC. Spleen mass was increased in *mdx* ( $P < 0.0001$ ) compared with wild type and was further increased in *mdx* mice following NAC intervention ( $P < 0.01$ ). Brown and white adipose tissue mass were lower in *mdx* ( $P < 0.01$  and  $P < 0.001$ ; one-way ANOVA with Newman–Keuls *post hoc* test and Kruskal–Wallis with Dunn’s *post hoc* test; brown and white, respectively) compared with wild type. Brown and white adipose tissue mass was unchanged by NAC administration. Tibia length and mass, and kidney and lung mass were equivalent in all three groups (Table 2).

**Table 2.** Body mass, tibia mass and length, organ mass and adipose tissue mass. Definition of abbreviations: NAC, N-acetylcysteine; RV, right heart ventricle; LV, left heart ventricle; BAT, brown adipose tissue; eWAT, epididymal white adipose tissue. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Data are shown as mean  $\pm$  SD and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal–Wallis with Dunn’s *post hoc* test. \* denotes *mdx* different from corresponding wild type group; \*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.001$  and \*\*\*\*  $P < 0.0001$ . § denotes *mdx* + NAC different from corresponding *mdx* group; §  $P < 0.05$  and §§  $P < 0.01$ .

	Wild type (n = 10)	<i>mdx</i> (n = 10)	<i>mdx</i> + NAC (n = 10)	P Value
Body mass (g)	24.8 $\pm$ 1.5	26.0 $\pm$ 0.8	26.7 $\pm$ 1.6	$P = 0.0194$
Tibia length (mm)	16.3 $\pm$ 0.7	16.4 $\pm$ 1.1	16.2 $\pm$ 0.7	$P = 0.9528$
Tibia mass (mg)	40.7 $\pm$ 4.1	39.9 $\pm$ 6.4	38.8 $\pm$ 4.5	$P = 0.7019$
RV (mg)	20.6 $\pm$ 3.6	28.9 $\pm$ 2.8 ****	25.6 $\pm$ 2.2 §	$P < 0.0001$
LV (mg)	70.2 $\pm$ 5.1	82.7 $\pm$ 7.9 **	79.5 $\pm$ 7.8	$P = 0.0048$
RV:LV	0.29 $\pm$ 0.06	0.35 $\pm$ 0.05 *	0.33 $\pm$ 0.05	$P = 0.0411$
Spleen (mg)	58.4 $\pm$ 7.2	99.0 $\pm$ 10.1 ****	119.9 $\pm$ 25.0 §§	$P < 0.0001$
Kidney (mg)	301.7 $\pm$ 26.4	322.9 $\pm$ 64.2	359.8 $\pm$ 31.0	$P = 0.004$
Lung (mg)	136.4 $\pm$ 14.4	150.9 $\pm$ 36.2	159.3 $\pm$ 36.2	$P = 0.4124$
BAT (mg)	64.2 $\pm$ 9.0	43.8 $\pm$ 14.2 **	53.5 $\pm$ 7.5	$P = 0.0048$
eWAT (mg)	407.8 $\pm$ 43.2	216.2 $\pm$ 57.3 ***	217.9 $\pm$ 58.3	$P < 0.0001$

### 3.2. Baseline Ventilation and Ventilatory Responsiveness to Hypercapnic-Hypoxia in Conscious Mice

Respiratory parameters including  $f_R$ ,  $V_T$  and  $V_E$  during baseline (21% O<sub>2</sub>) and hypercapnic-hypoxia (10% O<sub>2</sub> & 6% CO<sub>2</sub>) are shown in Table 3. During baseline conditions,  $f_R$ ,  $V_T$  and  $V_E$  were equivalent between all three groups. During exposure to hypercapnic-hypoxia,  $V_T$  was lower in *mdx* ( $P < 0.05$ ; Kruskal–Wallis with Dunn’s *post hoc* test) compared with wild type,  $f_R$  and  $V_E$  were equivalent between wild type and *mdx*. NAC did not alter  $f_R$ ,  $V_T$  and  $V_E$  during hypercapnic-hypoxia.

**Table 3.** Baseline ventilation in conscious mice. Definition of abbreviations:  $f_R$ , respiratory frequency; NAC, N-acetylcysteine;  $V_E$ , minute ventilation;  $V_T$ , tidal volume. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Data are shown as mean  $\pm$  SD and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal-Wallis with Dunn's *post hoc* test. \* denotes *mdx* different from corresponding wild type group; \*  $P < 0.05$ .

	Wild type (n = 10)	<i>mdx</i> (n = 8)	<i>mdx</i> + NAC (n = 10)	P Value
<b>Normoxia (21% O<sub>2</sub>)</b>				
$f_R$ (brpm)	196 $\pm$ 15	211 $\pm$ 14	205 $\pm$ 16	$P = 0.1136$
$V_T$ (mL g <sup>-1</sup> body mass)	0.007 $\pm$ 0.001	0.006 $\pm$ 0.001	0.006 $\pm$ 0.001	$P = 0.1295$
$V_E$ (mL g <sup>-1</sup> min <sup>-1</sup> )	1.31 $\pm$ 0.15	1.28 $\pm$ 0.25	1.29 $\pm$ 0.10	$P = 0.8891$
<b>Hypercapnic-Hypoxia (10% O<sub>2</sub> &amp; 6% CO<sub>2</sub>)</b>				
$f_R$ (brpm)	366 $\pm$ 28	360 $\pm$ 61	369 $\pm$ 15	$P = 0.8817$
$V_T$ (mL g <sup>-1</sup> body mass)	0.014 $\pm$ 0.002	0.011 $\pm$ 0.002 *	0.012 $\pm$ 0.001	$P = 0.0147$
$V_E$ (mL g <sup>-1</sup> min <sup>-1</sup> )	5.01 $\pm$ 0.98	3.99 $\pm$ 1.24	4.56 $\pm$ 0.47	$P = 0.0903$
Body mass (g)	24.8 $\pm$ 1.5	26.1 $\pm$ 0.8	26.7 $\pm$ 1.6	$P = 0.0222$

### 3.3. Inspiratory Pressure and Diaphragm and Intercostal EMGs during Airway Obstruction in Anaesthetized Mice

Previously, we reported findings from recordings of oesophageal pressure and diaphragm and external intercostal EMG activities for two of the groups, wild type and *mdx*, in the context of a comprehensive assessment of respiratory system deficits and compensations in young *mdx* mice [13]. Here, we compare values for the *mdx* + NAC group to our previously published values. During baseline recordings, oesophageal pressure ( $P = 0.9843$ ; Kruskal-Wallis test; Table 4) and diaphragm ( $P = 0.0907$ ; Kruskal-Wallis test; Table 4) and external intercostal ( $P > 0.05$ ; one-way ANOVA with Newman-Keuls *post hoc* test; Table 4) EMG activities in *mdx* + NAC were equivalent to values for *mdx*. In anaesthetized mice, protracted airway obstruction increased oesophageal sub-atmospheric pressure generation and EMG activity in all groups, measured as the average of five consecutive peak inspiratory efforts. Compared with *mdx* mice, oesophageal pressure ( $P = 0.0954$ ; one-way ANOVA; Table 4) and diaphragm ( $P = 0.1741$ ; Table 4) and external intercostal ( $P > 0.05$ ; Kruskal-Wallis with Dunn's *post hoc* test; Table 4) EMG activities were not different following NAC administration during protracted airway occlusion.

**Table 4.** Oesophageal pressure and diaphragm and T3 external intercostal (EIC) EMG activities during baseline and tracheal occlusion in anaesthetized mice. Definition of abbreviations: NAC, N-acetylcysteine; dia, diaphragm; EIC, external intercostal; NAC, N-acetylcysteine; A.U., arbitrary units. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Sustained tracheal occlusion was reported as the average of five successive peak breaths. Data are shown as mean  $\pm$  SD and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal-Wallis with Dunn's *post hoc* test. \* denotes *mdx* different from corresponding wild type group; \*\*  $P < 0.01$ .

	Wild type (n = 10)	<i>mdx</i> (n = 10)	<i>mdx</i> + NAC (n = 10)	P Value
<b>Baseline</b>				
Inspiratory pressure (cmH <sub>2</sub> O)	-9.0 $\pm$ 3.0	-9.2 $\pm$ 3.5	-9.3 $\pm$ 3.4	$P = 0.9843$
Dia EMG Amplitude (A.U.)	9.6 $\pm$ 6.7	5.5 $\pm$ 5.3	5.5 $\pm$ 3.7	$P = 0.0907$
T3 EIC EMG Amplitude (A.U.)	6.2 $\pm$ 2.7	3.3 $\pm$ 2.0 **	3.2 $\pm$ 1.1	$P = 0.0027$
<b>Obstruction</b>				
Inspiratory pressure (cmH <sub>2</sub> O)	-67.5 $\pm$ 10.2	-59.7 $\pm$ 11.8	-55.4 $\pm$ 13.3	$P = 0.0954$
Dia EMG Amplitude (A.U.)	38.6 $\pm$ 16.2	25.0 $\pm$ 14.6	30.4 $\pm$ 16.8	$P = 0.1741$
T3 EIC EMG Amplitude (A.U.)	43.7 $\pm$ 19.9	14.3 $\pm$ 9.3 **	18.1 $\pm$ 12.5	$P = 0.0011$

### 3.4. Tracheal Airflow in Anaesthetized Mice

Table 5 shows respiratory data in anaesthetized wild type, *mdx* and *mdx* + NAC mice during baseline conditions (60% O<sub>2</sub>) and hypercapnic-hypoxia (F<sub>i</sub>O<sub>2</sub> = 0.15 & F<sub>i</sub>CO<sub>2</sub> = 0.05) in vagotomized mice. All respiratory parameters were equivalent comparing wild type, *mdx* and *mdx* + NAC mice.

**Table 5.** Respiratory measurements in anaesthetized mice in vivo. Definition of abbreviations: ETCO<sub>2</sub>, end-tidal carbon dioxide; *f<sub>R</sub>*, respiratory frequency; NAC, N-acetylcysteine; PIF, peak inspiratory flow; PEF, peak expiratory flow; SpO<sub>2</sub>, peripheral capillary oxygen saturation; V<sub>E</sub>, minute ventilation; V<sub>T</sub>, tidal volume. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Data are shown as mean ± SD and were statistically compared using one-way ANOVA or Kruskal–Wallis.

	Wild type (n = 10)	<i>mdx</i> (n = 7–8)	<i>mdx</i> + NAC (n = 8–10)	P Value
<b>Baseline Post-Vagotomy (60% O<sub>2</sub>)</b>				
<i>f<sub>R</sub></i> (brpm)	65 ± 21	61 ± 12	69 ± 19	P = 0.7421
V <sub>T</sub> (mL g <sup>−1</sup> )	0.028 ± 0.005	0.029 ± 0.003	0.030 ± 0.008	P = 0.8468
V <sub>E</sub> (mL g <sup>−1</sup> min <sup>−1</sup> )	1.8 ± 0.4	1.8 ± 0.3	1.9 ± 0.2	P = 0.5651
PIF (mL g <sup>−1</sup> sec <sup>−1</sup> )	1.10 ± 0.22	1.14 ± 0.12	1.12 ± 0.41	P = 0.9498
PEF (mL g <sup>−1</sup> sec <sup>−1</sup> )	1.08 ± 0.17	1.17 ± 0.27	1.20 ± 0.26	P = 0.5177
ETCO <sub>2</sub> (mmHg)	53.3 ± 3.4	49.6 ± 7.9	52.9 ± 3.3	P = 0.3033
SpO <sub>2</sub> (%)	94.9 ± 2.3	95.1 ± 2.4	93.5 ± 4.1	P = 0.5320
<b>Hypercapnic-Hypoxia (15% O<sub>2</sub> &amp; 6% CO<sub>2</sub>)</b>				
<i>f<sub>R</sub></i> (brpm)	57 ± 17	52 ± 12	62 ± 16	P = 0.4772
V <sub>T</sub> (mL g <sup>−1</sup> )	0.041 ± 0.007	0.046 ± 0.005	0.043 ± 0.009	P = 0.604
V <sub>E</sub> (mL g <sup>−1</sup> min <sup>−1</sup> )	2.3 ± 0.7	2.4 ± 0.6	2.6 ± 0.4	P = 0.6929
PIF (mL g <sup>−1</sup> sec <sup>−1</sup> )	1.43 ± 0.23	1.45 ± 0.14	1.43 ± 0.35	P = 0.611
PEF (mL g <sup>−1</sup> sec <sup>−1</sup> )	1.15 ± 0.19	1.38 ± 0.18	1.29 ± 0.20	P = 0.0655
SpO <sub>2</sub> (%)	59.8 ± 5.9	64.6 ± 9.8	55.7 ± 6.3	P = 0.0865

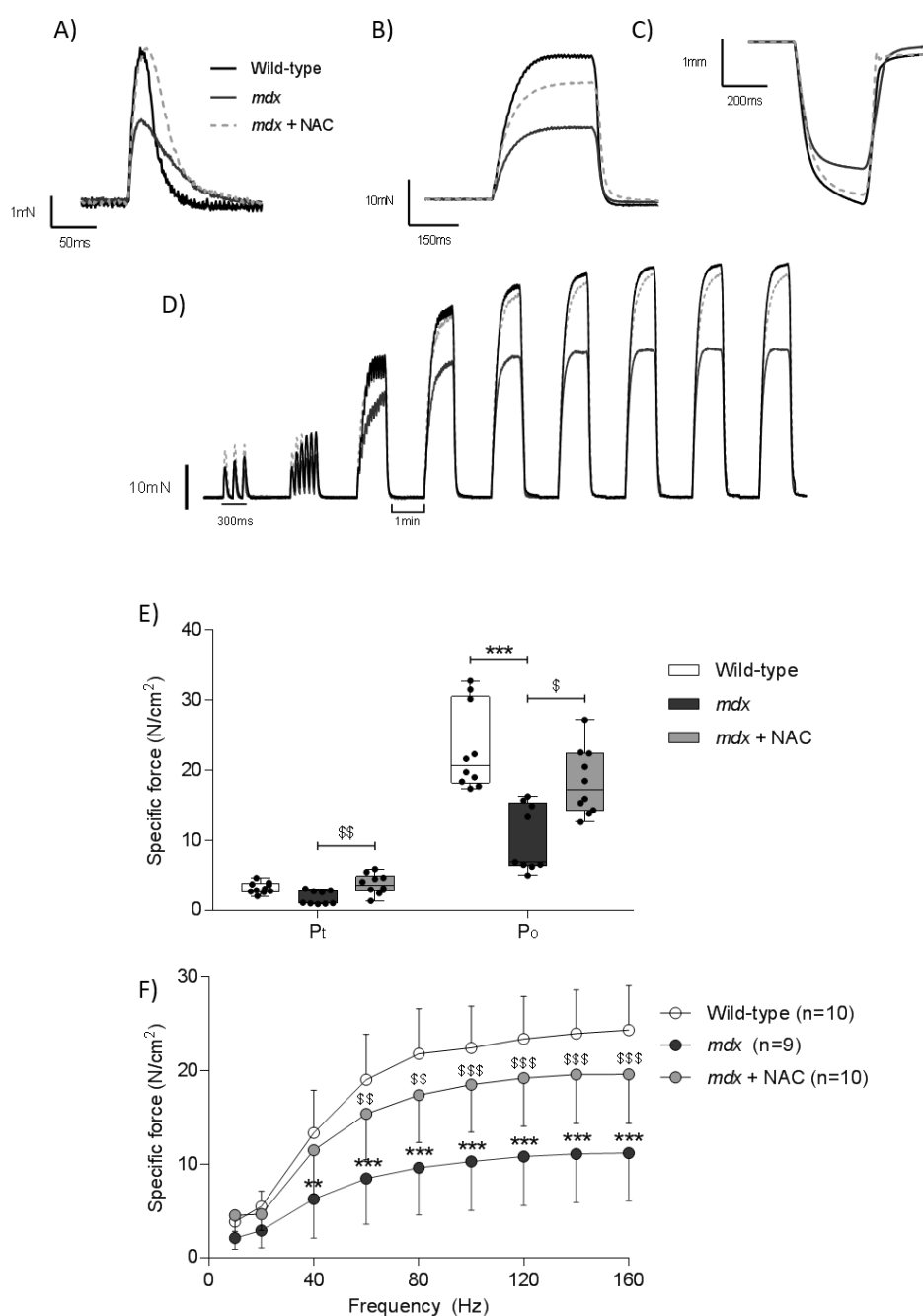
### 3.5. Diaphragm Muscle Contractile Function Ex Vivo

Table 6 shows data for twitch contractile kinetics (CT and ½ RT) and isotonic contractile kinetics (S<sub>max</sub> and V<sub>max</sub>) of diaphragm muscle preparations from all three groups. Diaphragm CT and ½ RT were equivalent in wild type and *mdx* preparations; NAC had no effect on CT and ½ RT for *mdx* diaphragm. Absolute values for S<sub>max</sub> were reduced in *mdx* (P < 0.01; Kruskal–Wallis with Dunn’s *post hoc* test) diaphragm preparations compared with wild type. NAC supplementation in *mdx* mice increased diaphragm S<sub>max</sub> and V<sub>max</sub> (P < 0.05 and P < 0.05, S<sub>max</sub> and V<sub>max</sub> respectively; Kruskal–Wallis with Dunn’s *post hoc* test). Normalized values for S<sub>max</sub> and V<sub>max</sub> were equivalent in all three groups.

**Table 6.** Ex vivo respiratory muscle contractile kinetics. Definition of abbreviations: ½ RT, half-relaxation time; CSA, cross-sectional area; CT, contraction time; L<sub>o</sub>, optimum length; NAC, N-acetylcysteine; P<sub>t</sub>, twitch force; P<sub>o</sub>, tetanic force; S<sub>max</sub>, maximum unloaded shortening; V<sub>max</sub>, maximum shortening velocity. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Data are shown as mean ± SD and were statistically compared using one-way ANOVA with Newman–Keuls *post hoc* test, or Kruskal–Wallis with Dunn’s *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*\* P < 0.01 and \*\*\* P < 0.001. \$ denotes *mdx* + NAC different from corresponding *mdx* group; \$ P < 0.05 and \$\$ P < 0.01.

	Wild type (n = 10)	<i>mdx</i> (n = 8–9)	<i>mdx</i> + NAC (n = 10)	P Value
CT (ms)	17.2 ± 2.6	18.1 ± 4.0	19.2 ± 2.5	P = 0.3816
½ RT (ms)	19.9 ± 6.9	21.7 ± 9.0	22.5 ± 4.2	P = 0.7034
S <sub>max</sub> (cm)	0.39 ± 0.08	0.23 ± 0.06 **	0.36 ± 0.05 \$	P = 0.0014
S <sub>max</sub> (L L <sub>o</sub> <sup>−1</sup> )	0.37 ± 0.06	0.31 ± 0.08	0.35 ± 0.03	P = 0.1016
V <sub>max</sub> (cm s <sup>−1</sup> )	4.1 ± 1.0	2.3 ± 1.2	4.4 ± 1.0 \$	P = 0.0184
V <sub>max</sub> (Lo s <sup>−1</sup> )	4.0 ± 1.0	3.2 ± 1.7	4.2 ± 0.7	P = 0.3223
L <sub>o</sub> (cm)	1.1 ± 0.1	0.8 ± 0.1 ***	1.0 ± 0.1 \$\$	P = 0.0004

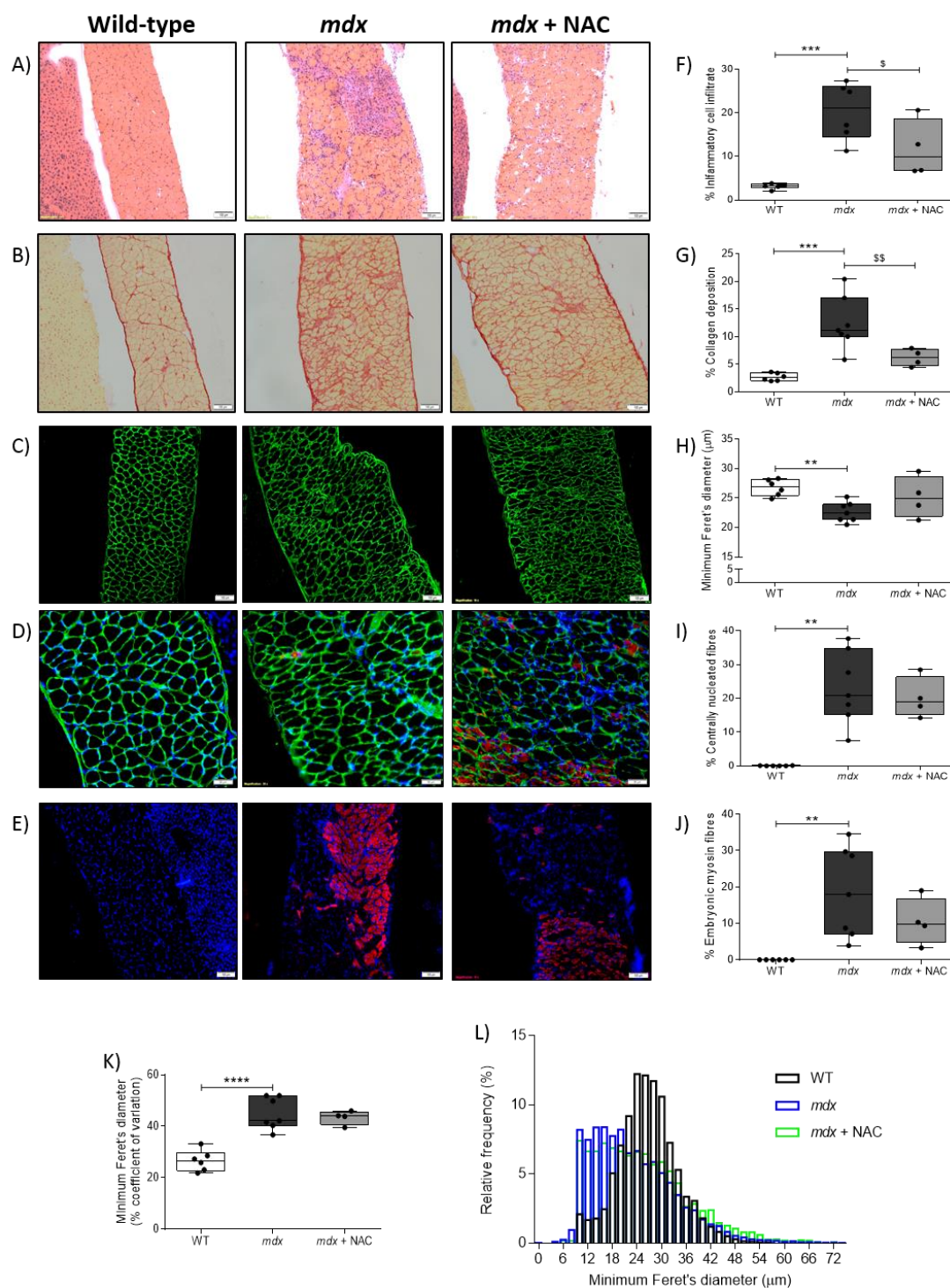
Figure 1 shows representative original traces for diaphragm muscle twitch (A) and tetanic (B) contractions, force-frequency relationship (C) and maximum unloaded shortening (D). NAC treatment increased diaphragm twitch force in *mdx* (Figure 1E,  $P_t$ ;  $P < 0.01$ ; Kruskal–Wallis with Dunn’s *post hoc* test) compared with *mdx*. Diaphragm tetanic force at 100 Hz was lower in *mdx* (Figure 1E,  $P_o$ ;  $P < 0.001$ ; Kruskal–Wallis with Dunn’s *post hoc* test) compared with wild type. NAC treatment increased *mdx* diaphragm tetanic force (Figure 1E,  $P_o$ ;  $P < 0.05$ ; Kruskal–Wallis with Dunn’s *post hoc* test) compared with *mdx*. For the force-frequency relationship, diaphragm specific force was reduced in *mdx* compared with wild type preparations (Figure 1F;  $P < 0.0001$ ; repeated measures two-way ANOVA). *Post hoc* analysis revealed differences between wild type and *mdx* diaphragm across a broad stimulus range (40–160 Hz). NAC supplementation to *mdx* mice improved diaphragm force-generating capacity compared with *mdx* preparations (Figure 1F;  $P < 0.0001$ ; repeated measures two-way ANOVA). *Post hoc* analysis indicated an improvement in *mdx* diaphragm force-generating capacity following NAC supplementation across a broad stimulus range (60–160 Hz).



**Figure 1.** Ex vivo diaphragm muscle contractile force. A–D, original traces of ex vivo diaphragm muscle twitch force (A), tetanic force at 100Hz (B), maximum unloaded shortening (C) and force–frequency relationship (D) for wild type (WT), *mdx* and *mdx* + N-acetylcysteine (NAC) preparations. E and F, group data for ex vivo diaphragm muscle twitch (E;  $P_i$ ) and tetanic (E;  $P_o$ ) force and force–frequency relationship (F) in wild type, *mdx* and *mdx* + NAC preparations. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. For  $P_i$  and  $P_o$ , values are expressed as scatter point box and whisker plots (median, 25–75 percentile and scatter plot) and were statistically compared by Kruskal–Wallis with Dunn’s *post hoc* test. For force–frequency, data are shown as mean  $\pm$  SD and were statistically compared using repeated measures two-way ANOVA with Bonferroni *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*\*  $P < 0.01$ , \*\*\*  $P < 0.001$ . § denotes *mdx* + NAC different from corresponding *mdx* group; §  $P < 0.05$ , §§  $P < 0.01$ , §§§  $P < 0.001$ .

### 3.6. Diaphragm Muscle Histology and Immunofluorescence

Figure 2A–E shows representative images for diaphragm muscle histology and immunohistochemistry for all three groups. The areal density of inflammatory cell infiltration was increased in *mdx* diaphragm compared with wild type (Figure 2F;  $P < 0.001$ ; one-way ANOVA with Newman–Keuls *post hoc* test). There was a reduction in immune cell infiltration in *mdx* diaphragm following NAC treatment compared with *mdx* diaphragm (Figure 2F;  $P < 0.05$ ). The areal density of collagen labelled with Sirius red staining was increased in *mdx* diaphragm compared with wild type (Figure 2G;  $P < 0.001$ ). NAC treatment in *mdx* resulted in a reduction in collagen deposition compared with untreated *mdx* mice (Figure 2G;  $P < 0.01$ ). There was a reduction in muscle fibre size (minimum Feret’s diameter) for *mdx* diaphragm compared with wild type (Figure 2H;  $P < 0.01$ ); NAC treatment had no effect on fibre size. The proportion of diaphragm muscle fibres with centrally located myonuclei was increased in *mdx* compared with wild type (Figure 2I;  $P < 0.01$ ; Kruskal–Wallis with Dunn’s *post hoc* test); NAC had no effect on the proportion of centrally located myonuclei. The proportion of diaphragm muscle fibres expressing embryonic myosin was increased in *mdx* diaphragm compared with wild type (Figure 2J;  $P < 0.01$ ; one-way ANOVA with Newman–Keuls *post hoc* test); NAC treatment had no effect on central nucleation and embryonic myosin expression in *mdx* diaphragm. The coefficient of variation of muscle fibre size was increased in *mdx* diaphragm compared with wild type (Figure 2K;  $P < 0.0001$ ); NAC treatment had no effect on fibre size variability. There was a wider distribution of muscle fibre size (based on minimum Feret’s diameter) in *mdx* diaphragm compared with wild type (Figure 2L). The distribution of diaphragm muscle fibres in *mdx* + NAC was similar to *mdx* mice, with a slightly greater abundance of large fibres.

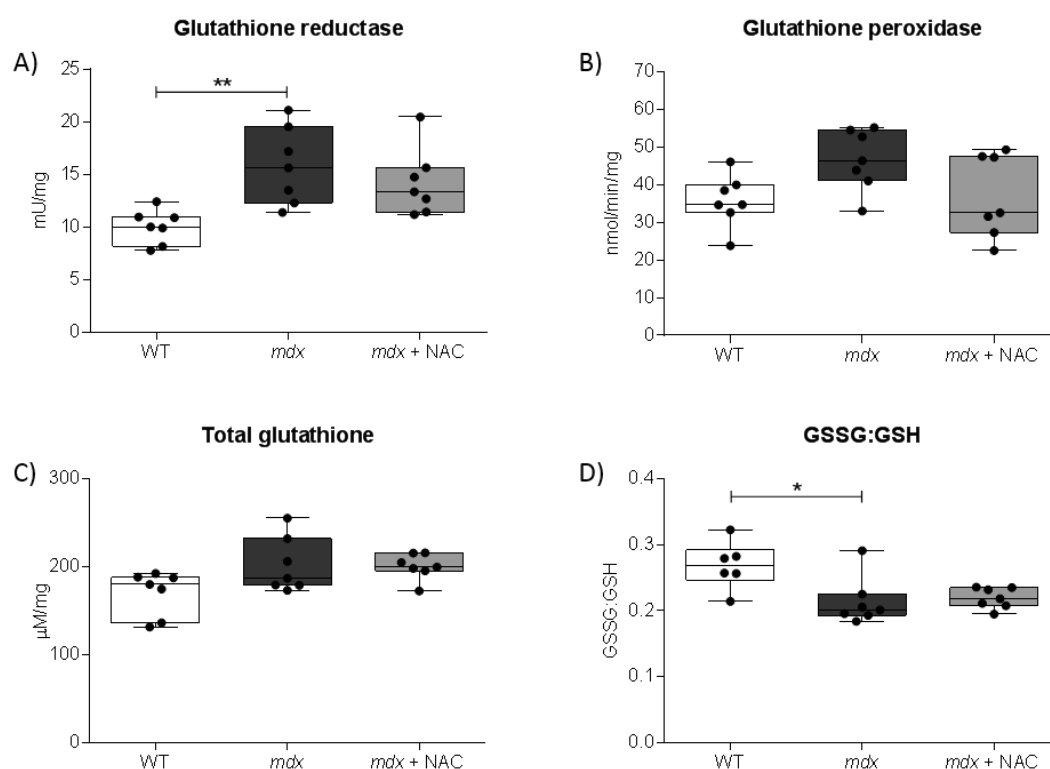


**Figure 2.** Diaphragm muscle histology and immunofluorescence. A and B, representative histological images of transverse sections of diaphragm muscle stained with Haematoxylin and Eosin (A) and Sirius red (B) in wild type (WT), *mdx* and *mdx* + NAC groups. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. C–E, representative images of diaphragm muscle immunofluorescently labelled for laminin (C), laminin, DAPI and embryonic myosin (D) and embryonic myosin and DAPI (E) for WT, *mdx* and *mdx* + NAC groups. Scale bars represent 100  $\mu$ m in A, B, C and E. In D, scale bars represent 50  $\mu$ m. F–J, group data showing the relative area of infiltration of inflammatory cells (F), relative area of collagen deposition (G), muscle fibre size measured by minimum Feret's diameter (H), relative area of centrally nucleated diaphragm muscle fibres (I) and relative area of diaphragm muscle fibres expressing embryonic myosin (J) in diaphragm muscle from WT, *mdx* and *mdx* + NAC groups. K–L, group data for the coefficient of variation of muscle fibre size (K) and the frequency distribution of muscle fibre size (L) for all three groups. Values are expressed

as scatter point box and whisker plots (median, 25–75 percentile and scatter plot) and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal-Wallis with Dunn's *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*\*  $P < 0.01$ , \*\*\*  $P < 0.001$  and \*\*\*\*  $P < 0.0001$ . § denotes *mdx* + NAC different from corresponding *mdx* group; §  $P < 0.05$  and §§  $P < 0.01$ .

### 3.7. Diaphragm Glutathione Status

Glutathione reductase activity was increased in *mdx* diaphragm (Figure 3A;  $P < 0.01$ ; one-way ANOVA with Newman-Keuls *post hoc* test) compared with wild type. GSSG:GSH was decreased in *mdx* diaphragm (Figure 3D;  $P < 0.05$ ; Kruskal-Wallis with Dunn's *post hoc* test) compared with wild type. Diaphragm glutathione peroxidase and total glutathione were equivalent in all three groups. Diaphragm glutathione reductase, glutathione peroxidase, total glutathione and GSSG:GSH were all unaffected by NAC supplementation in *mdx* mice.



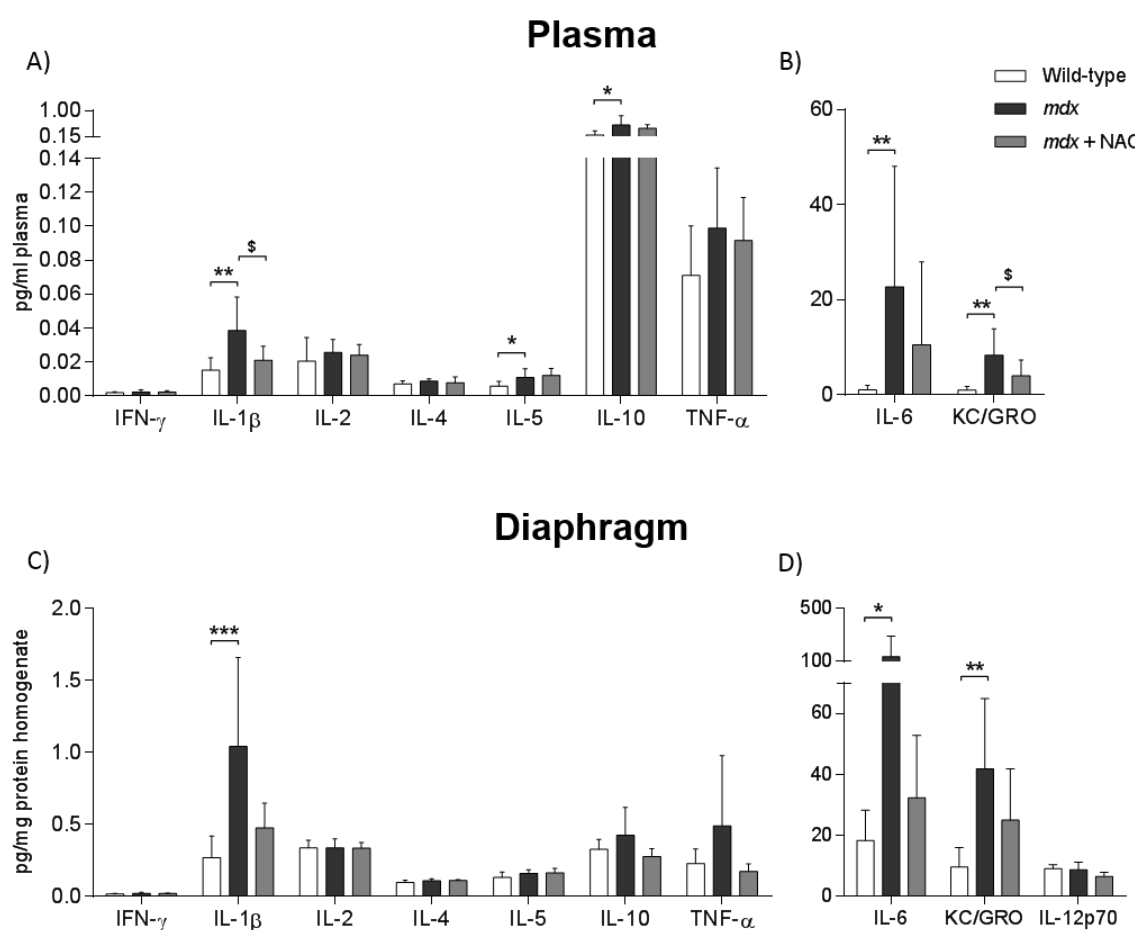
**Figure 3.** Diaphragm muscle glutathione status. A and B, group data for glutathione reductase (A) and glutathione peroxidase (B) activities of diaphragm muscle from wild type (WT), *mdx* and *mdx* + NAC groups. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. C and D, group data for total glutathione (C) and the ratio of oxidised glutathione (GSSG) to glutathione (GSSG:GSH; D). Values are expressed as scatter point box and whisker plots (median, 25–75 percentile and scatter plot) and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal–Wallis with Dunn's *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*  $P < 0.05$  and \*\*  $P < 0.01$ .

### 3.8. Plasma and Diaphragm Cytokine Concentrations

Plasma samples from *mdx* mice had increased concentration of the cytokines, IL-1 $\beta$  (Figure 4A;  $P < 0.01$ ; one-way ANOVA with Newman–Keuls *post hoc* test), IL-5 (Figure 4A;  $P < 0.05$ ); IL-6 (Figure 4B;  $P < 0.01$ ; Kruskal–Wallis with Dunn's *post hoc* test) and IL-10 (Figure 4A;  $P < 0.05$ ; Kruskal–Wallis with Dunn's *post hoc* test). The chemokine KC/GRO was increased in *mdx* plasma (Figure 4B;  $P < 0.01$ ; one-way ANOVA with Newman–Keuls *post hoc* test) compared with wild type. NAC



supplementation decreased the concentration of IL-1 $\beta$  (Figure 4A;  $P < 0.01$ ; one-way ANOVA with Newman–Keuls post hoc test) and KC/GRO (Figure 4B;  $P < 0.05$ ; one-way ANOVA with Newman–Keuls post hoc test) in *mdx* plasma. Plasma IFN- $\gamma$  (Figure 4A), IL-2 (Figure 4A), IL-4 (Figure 4A) and TNF- $\alpha$  (Figure 4A) concentrations were equivalent in all three groups. Increased levels of IL-1 $\beta$  (Figure 4C;  $P < 0.001$ ; Kruskal–Wallis with Dunn’s *post hoc* test), IL-6 (Figure 4D;  $P < 0.05$ ; Kruskal–Wallis with Dunn’s *post hoc* test) and KC/GRO (Figure 4D;  $P < 0.01$ ; one-way ANOVA with Newman–Keuls post hoc test) were observed in *mdx* diaphragm. *Post hoc* analysis revealed no changes in *mdx* diaphragm IL-1 $\beta$ , IL-6 or KC/GRO concentrations following NAC ( $P > 0.05$ ) but notably the concentrations of IL-1 $\beta$  and IL-6 in diaphragms from *mdx* + NAC were >50% reduced compared with *mdx*. Diaphragm IFN- $\gamma$  (Figure 4C), IL-2 (Figure 4C), IL-4 (Figure 4C), IL-5 (Figure 4C), IL-10 (Figure 4C), TNF- $\alpha$  (Figure 4C) and IL-12p70 (Figure 4D) were equivalent in all three groups.



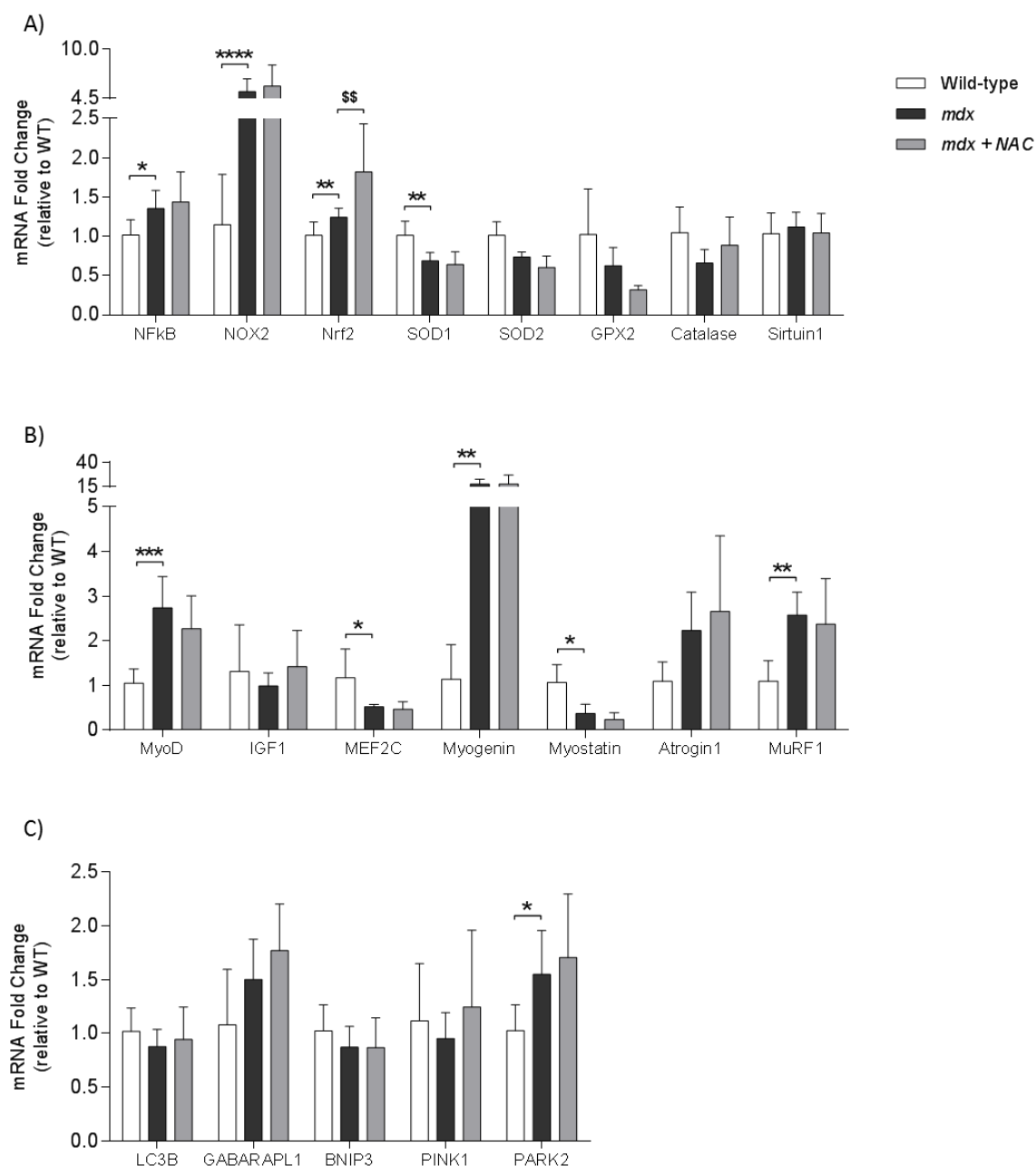
**Figure 4.** Plasma and diaphragm muscle cytokine concentrations. A–D, group data for selective cytokine concentrations in plasma and diaphragm muscle from wild type ( $n = 7$ ), *mdx* ( $n = 7$ –9) and *mdx* + NAC ( $n = 7$ –9) groups. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Plasma cytokines include interferon- $\gamma$  (IFN- $\gamma$ ; A), interleukin-1 $\beta$  (IL-1 $\beta$ ; A), interleukin-2 (IL-2; A), interleukin-4 (IL-4; A), interleukin-5 (IL-5; A), interleukin-6 (IL-6; B), interleukin-10 (IL-10; A), KC/GRO (B), tumour necrosis factor- $\alpha$  (TNF- $\alpha$ ; A). Diaphragm cytokines include IFN- $\gamma$  (C), IL-1 $\beta$  (C), IL-2 (C), IL-4 (C), IL-5 (C), IL-6 (D), IL-10 (P), KC/GRO (D), IL-12 (IL-12p70; D) and TNF- $\alpha$  (C). Data are shown as mean  $\pm$  SD and were statistically compared using one-way ANOVA with Newman–Keuls *post hoc* test, or Kruskal–Wallis with Dunn’s *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*  $P < 0.05$ , \*\*  $P < 0.01$  and \*\*\*  $P < 0.001$ . \$ denotes *mdx* + NAC different from corresponding *mdx* group; \$  $P < 0.05$ .

### 3.9. Diaphragm Muscle mRNA Expression

Figure 5 shows data for mRNA expression of genes related to redox status, inflammation, muscle repair, autophagy and mitophagy in diaphragm muscle from wild type, *mdx* and *mdx* + NAC groups. There was increased expression of the pro-inflammatory transcription factor NF $\kappa$ B in *mdx* diaphragm (Figure 5A;  $P = 0.0184$ ; one-way ANOVA with Newman-Keuls *post hoc* test) compared with wild type; NAC administration had no effect. The expression of the reactive oxygen species generating complex NOX2 was increased in *mdx* diaphragm (Figure 5A;  $P < 0.0001$ ) and was unaffected by NAC administration. Expression of the antioxidant response element, Nrf2, was similar in wild type and *mdx* diaphragm ( $P > 0.05$ ); Nrf2 expression was increased in *mdx* + NAC compared with *mdx* (Figure 5A;  $P < 0.01$ ). Expression of the antioxidant gene SOD1 was decreased in *mdx* diaphragm (Figure 5A;  $P < 0.01$ ) compared with wild type; NAC treatment in *mdx* had no effect. There were no differences between groups for SOD2 and GPX2 in *post hoc* analyses.

The muscle repair genes MyoD and Myogenin had increased mRNA expression in *mdx* diaphragm (Figure 5B;  $P < 0.001$  and  $P = 0.01$ ; MyoD and Myogenin respectively; one-way ANOVA with Newman-Keuls *post hoc* test and Kruskal–Wallis with Dunn's *post hoc* test respectively); NAC treatment in *mdx* had no effect on both genes. mRNA expression of Myostatin, an inhibitor of myogenesis, was reduced in *mdx* diaphragm (Figure 5B;  $P < 0.01$ ; one-way ANOVA with Newman-Keuls *post hoc* test) compared with wild type diaphragm; NAC treatment in *mdx* had no effect. The muscle differentiation transcription factor MEF2C had reduced mRNA expression in *mdx* diaphragm compared with wild type (Figure 5B;  $P < 0.05$ ) and was unaffected by NAC treatment in *mdx*.

The atrophy-related gene MuRF1 had increased mRNA expression in *mdx* diaphragm (Figure 5B;  $P < 0.01$ ) compared with wild type; MuRF1 expression was unaffected by NAC administration. The mitophagy-related gene PARK2, had increased mRNA expression in *mdx* diaphragm (Figure 5C;  $P < 0.05$ ) compared with wild type; PARK2 mRNA expression was unaffected by NAC treatment in *mdx*. There were no differences between groups for GABARAPL1 in *post hoc* analyses. mRNA expression of Catalase (Figure 5A), Sirtuin1 (Figure 5A), IGF1 (Figure 5B), Atrogin1 (Figure 5B), PINK1 (Figure 5C), LC3B (Figure 5C) and BNIP3 (Figure 5C) was equivalent in all three groups.



**Figure 5.** Diaphragm muscle mRNA expression. *Definition of abbreviations:* BNIP-3, BCL3 interacting protein; GPX, glutathione peroxidase; GABARAPL1, GABA type A receptor associated protein Like 1; IGF, insulin-like growth factor; LC3B, light chain 3 beta; MEF2C, myocyte enhancer factor 2C; MyoD, myogenic differentiation; Murf-1, muscle RING-finger protein-1; NAC, N-acetylcysteine; NOX, NADPH oxidase; Nrf-2, nuclear factor (erythroid-derived 2)-like 2; NFκB, nuclear factor kappa-light-chain-enhancer of activated B cells; PARK-2, parkin RBR E3 ubiquitin protein ligase; PINK-1, phosphatase and tensin homolog (PTEN)-induced kinase 1; SOD, superoxide dismutase; WT, wild type. A–C, group data for mRNA expression of NFκB (A), NOX2 (A), Nrf2 (A), SOD1 (A), SOD2 (A), GPX2 (A), catalase (A), sirtuin1 (A), MyoD (B), IGF1 (B), MEF2C (B), myogenin (B), myostatin (B), atrogin1 (B), MuRF1 (B), LC3B (C), GABARAPL1 (C), BNIP3 (C), PINK1 (C) and PARK2 (C) in diaphragm muscle from wild type (WT;  $n = 7$ –8), *mdx* ( $n = 6$ –7) and *mdx* + NAC ( $n = 6$ –7) groups. *Mdx* + NAC group received 1% N-acetylcysteine in the drinking water for 14 days. Data are shown as mean  $\pm$  SD and were statistically compared using one-way ANOVA with Newman-Keuls *post hoc* test, or Kruskal-Wallis with Dunn's *post hoc* test. \* denotes *mdx* different from corresponding WT group; \*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.001$  and \*\*\*\*  $P < 0.0001$ . \$ denotes *mdx* + NAC different from corresponding *mdx* group; \$\$  $P < 0.01$ .

#### 4. Discussion

The major findings of the study are: (i) NAC supplementation improved *mdx* diaphragm functional capacity (ii) NAC supplementation reduced collagen deposition (fibrosis) and immune cell infiltration in *mdx* diaphragm; (iii) NAC treatment reduced IL-1 $\beta$  and KC/GRO concentrations in *mdx* plasma samples; (iv) NAC increased Nrf2 mRNA expression in *mdx* diaphragm; (v) NAC had no adverse effect on ventilation, inspiratory pressure and respiratory muscle EMG or growth measures in *mdx* mice; (vi) NAC did not recover decreased respiratory muscle EMG activity in *mdx* mice during maximum activation (airway obstruction).

Dystrophin deficiency in DMD results in mechanical deficits in striated muscle [5]. Respiratory muscle weakness occurs in DMD, with major implications for respiratory control [37]. Inspiratory muscle strength declines with increasing age in DMD boys [6]. Respiratory disturbances are common during sleep, including hypoventilation [38,39] and apnoeic events [37,40,41]. The *mdx* mouse model of DMD has characteristics similar to DMD, including disruptions to normal breathing [12,42–44] and diaphragm and pharyngeal dilator muscle weakness [12,15,16,32,45]. Adequate respiratory muscle function is essential for ventilation and mechanical manoeuvres including coughing and sneezing, mechanisms which safeguard against airway collapse. Given the clinical implications of poor respiratory system function in DMD, the development of new therapeutic strategies aimed at boosting respiratory muscle performance is essential.

Herein, diaphragm muscle weakness in *mdx* mice was evidenced by reduced tetanic force, consistent with previous reports in young and older *mdx* mice [12–14,16,17]. Specific force was reduced within the frequency range of 40–160 Hz for *mdx* diaphragm compared with wild type. This frequency range corresponds to a broad range of ventilatory and non-ventilatory behaviours [13,46–48]. Structural abnormalities occur in dystrophic muscle as a consequence of muscle fibre damage and degeneration. Our study confirmed structural alterations in *mdx* diaphragm, including a substantial increase in the number of centralized myonuclei (measure of fibre regeneration following damage), increased expression of MyHC<sub>EMB</sub> and increased collagen content (fibrosis). We observed a leftward shift in the frequency distribution of dystrophic muscle fibre size and a concomitant increase in the variability of fibre size. These observations are consistent with and extend our previous findings [12,17] and those of others [49,50], revealing substantial diaphragm muscle pathology in young (8 week old) *mdx* mice.

Systemic and muscle inflammation are prominent features of DMD, secondary to muscle fibre damage. There is elevated expression of chemokines [51,52] and pro-inflammatory cytokines [53,54] in serum and muscle biopsies from DMD boys. In the current study, we report elevated pro-inflammatory (IL-1 $\beta$  and IL-6) and anti-inflammatory (IL-10) cytokine concentrations in plasma samples from *mdx* mice. Immune cells are recruited to areas of damaged muscle, evidenced by increased immune cell infiltration in dystrophic muscle [55]. Here, we report an increase in putative inflammatory cells in *mdx* diaphragm. The concentrations of the pro-inflammatory cytokines, IL-1 $\beta$  and IL-6, and the chemokine KC/GRO are increased in *mdx* diaphragm. Anti-inflammatory drugs have been shown to have beneficial effects on *mdx* skeletal muscle form, but limited improvements in muscle force [55,56]. Encouragingly, however, blockade of pro-inflammatory cytokine signalling has proved beneficial to *mdx* respiratory muscle form and function [17,33,57,58].

Recent studies suggest a close relationship between oxidative stress and inflammation in dystrophic muscle pathology [4,21,59]. Oxidative stress develops as a consequence of increased generation of ROS and/or decreased endogenous antioxidant capacity. Immune cells in muscle can produce ROS [60], which at high levels can lead to impaired muscle force-generating capacity [19]. ROS are primarily produced by the mitochondria and enzyme complexes such as NADPH oxidase (NOX) and xanthine oxidase. Cellular damage occurs during periods of excessive ROS production. Muscle biopsies from DMD boys indicate increased levels of lipid peroxidation, an indirect measure of elevated ROS [61]. Expression of the ROS generating enzyme NOX2 is elevated in DMD muscle [4]. Endogenous antioxidant defence appears to be curtailed in DMD. For example, the activity of superoxide dismutase is decreased in DMD muscle [62]. Studies of the glutathione system in DMD boys suggests glutathione deficiency [63,64]. In DMD muscle biopsies, decreased antioxidant activity

confers reduced antioxidant defence increasing the risk of oxidative damage. Similarly, diaphragm muscle weakness in *mdx* mice is associated with altered redox balance [21,65]. ROS production is increased in *mdx* muscle [66,67]. NOX is elevated in *mdx* diaphragm and limb muscle [68–70]. Levels of 4-hydroxynonenal are elevated in *mdx* diaphragm, indicating increased lipid peroxidation [28]. Respiratory and limb muscle samples from *mdx* mice have decreased citrate synthase activity, suggesting reduced mitochondrial content [22,27]. Protein thiol oxidation and carbonyl content are increased in *mdx* diaphragm [71], further supporting redox modulation of dystrophic respiratory muscle.

Our examination of diaphragm mRNA expression indicated an increase in mRNA expression related to muscle atrophy (MuRF1), regeneration (MyoD, myogenin), inflammation (NFκB), oxidative stress (NOX2) and mitophagy (PARK2) in *mdx* diaphragm, and a reduction in antioxidant (SOD1)-related mRNA expression compared with wild type. The transcription factor Nrf2 is considered a regulator of antioxidant proteins in response to oxidative injury or inflammation. Nrf2 mRNA was increased in *mdx* diaphragm, consistent with studies of elevated Nrf2 in DMD muscle [4]. An upregulation of Nrf2 signalling in dystrophic muscle suggests a defensive antioxidant response.

We investigated the effects of NAC supplementation in *mdx* mice on ventilation in conscious mice, and inspiratory pressure and respiratory (diaphragm and external intercostal) EMG activity in anaesthetized mice. In addition, we determined diaphragm muscle structure and force-generating capacity, diaphragm glutathione, diaphragm and plasma cytokine concentrations, and diaphragm mRNA expression. NAC is a free radical scavenger and a precursor of reduced glutathione (GSH), boosting synthesis of the endogenous antioxidant. We administered 1% NAC in the drinking water to *mdx* mice for 14 consecutive days beginning at 6 weeks of age. We have previously demonstrated the beneficial effects of NAC supplementation in rodent models of hypoxia-induced respiratory muscle weakness, where NAC improves respiratory muscle force-generating capacity and boosts endogenous cell survival signalling [24–26,72]. In addition to its effects on the glutathione system and its important role as a ROS scavenger, NAC is also a mucolytic agent and has been considered as a therapeutic agent in respiratory conditions such as chronic obstructive pulmonary disease [73]. Previously, NAC was shown to attenuate diaphragm muscle fatigue in humans [74].

In the current study, NAC supplementation increased diaphragm specific force in *mdx* mice across the frequency range of 60–160 Hz. Structural improvements following NAC included a reduction in diaphragm collagen content and a decrease in the infiltration of putative inflammatory cells. Total absolute muscle shortening and maximal shortening velocity were both improved in *mdx* diaphragm following NAC supplementation due to improvements in *mdx* muscle optimal length. This was likely mediated, at least in part, by the reduction in diaphragm collagen content following NAC, decreasing tissue stiffness. Recent studies of NAC administration in *mdx* mice demonstrated improved grip strength, increased EDL force output *ex vivo*, and reduced muscle inflammation and oxidative stress [27]. Whitehead et al. (2008) also reported increased EDL force and reduced muscle damage and inflammation following NAC intervention [30]. Furthermore, NAC reduced myosin protein thiol oxidation in *mdx* limb muscle [27]. Interestingly, NAC has been shown to prevent myofibre damage in *mdx* mice following exercise [29,75]. Anti-fibrotic actions of NAC have also been observed in animal models of disease. For example, NAC decreased TGFβ signalling in fibroblasts [76]; cardiac fibrosis was decreased by NAC in a mouse model of heart failure [77]; NAC decreased fibrosis in soleus muscle in a murine model of peripheral arterial insufficiency [78]; and renal fibrosis was decreased by NAC in a model of dilated cardiomyopathy [79].

We examined the glutathione system in the diaphragm of wild type, *mdx* and *mdx* + NAC mice. Glutathione is an endogenous antioxidant that exists in both a reduced (GSH) and oxidized (GSSG) state. Glutathione peroxidase (GPx) catalyses the oxidation reaction of GSH to GSSG, whereas glutathione reductase catalyses the reduction of GSSG to GSH. Glutathione reductase activity was increased in *mdx* diaphragm, along with a reduction in the GSSG:GSH ratio. These data suggest an early antioxidant defence, perhaps in response to a modest oxidative stress (buffered by glutathione reductase) in diaphragm from young *mdx* mice.

Given the close relationship between inflammation and oxidative stress, we examined immune cell infiltration in the diaphragm and the concentration of cytokines in diaphragm and plasma samples. Supplementation with NAC reduced the relative area of infiltrating inflammatory cells in *mdx* diaphragm. Cytokine concentrations were determined in plasma and diaphragm muscle samples following NAC administration. The pro-inflammatory cytokine IL-1 $\beta$  and the chemokine KC/GRO were reduced in *mdx* plasma samples following NAC. In addition, there was a sizeable reduction in plasma IL-6 levels ( $11 \pm 18$  versus  $23 \pm 25$  pg.mL<sup>-1</sup> plasma; *mdx* + NAC versus *mdx*). For diaphragm, sizeable reductions in the concentration of IL-1 $\beta$  ( $0.47 \pm 0.17$  versus  $1.04 \pm 0.62$  pg.mg<sup>-1</sup> protein homogenate; *mdx* + NAC versus *mdx*), IL-6 ( $32 \pm 21$  versus  $137 \pm 155$  pg.mg<sup>-1</sup> protein homogenate) and KC/GRO ( $25 \pm 17$  versus  $42 \pm 23$  pg.mg<sup>-1</sup> protein homogenate) were observed following NAC. These data are in agreement with previous studies of NAC intervention in *mdx* mice. Pinniger et al. (2017) demonstrated that NAC supplementation reduced neutrophils and myeloperoxidase activity in *mdx* gastrocnemius muscle [27]. De Senzi Moraes Pinto et al. (2013) reported that NAC supplementation in young *mdx* mice reduced TNF- $\alpha$  protein expression [28]. Given the emerging benefits of reduced pro-inflammatory cytokine signalling in *mdx* respiratory muscle, these anti-inflammatory effects of NAC likely contributed to improved respiratory muscle performance in *mdx* mice. Our study suggests that the potential anti-inflammatory effects of NAC in dystrophic muscle warrant further attention.

Breathing was assessed using whole-body plethysmography in conscious mice during baseline (21% O<sub>2</sub>) and chemoactivated breathing with hypercapnic-hypoxia (F<sub>I</sub>O<sub>2</sub> = 0.10 and F<sub>I</sub>CO<sub>2</sub> = 0.06). These assessments confirmed the capacity for *mdx* mice to enhance ventilation as previously described [12,13]. Tidal volume generation during hypercapnic-hypoxia was lower in *mdx* compared with wild type and was not rescued by NAC. In addition, we examined respiratory airflow and volume measures in anaesthetized mice following bilateral section of the vagi and with superimposed chemo-activation (F<sub>I</sub>O<sub>2</sub> = 0.15 and F<sub>I</sub>CO<sub>2</sub> = 0.06) to maximally activate ventilation. V<sub>T</sub> and inspiratory and expiratory flows were equivalent or greater in *mdx* mice compared with wild type revealing preserved ventilatory capacity in young *mdx* mice despite dystrophic disease. Direct measures of f<sub>R</sub>, V<sub>T</sub> and V<sub>E</sub> and peak inspiratory and expiratory flows were equivalent between *mdx* and *mdx* + NAC groups, revealing no adverse effect of NAC on respiratory airflow and volume in *mdx* mice.

Our recent study reported lower diaphragm EMG activity in anaesthetized *mdx* mice during maximum activation [13]. Despite structural remodelling, profound weakness and lower EMG activity in *mdx* diaphragm, inspiratory pressure-generating capacity is remarkably preserved across ventilatory (chemoactivated breathing and augmented breaths) and non-ventilatory behaviours (sustained tracheal occlusion) [13], revealing a role for accessory and auxiliary muscles in support of peak inspiratory pressure generation in dystrophic disease [80]. We compared oesophageal pressure and respiratory muscle EMG (diaphragm and external intercostal) measurements in *mdx* mice following supplementation with 1% NAC to our previously published results. Values for oesophageal pressure during baseline ( $-9.3 \pm 3.4$  versus  $-9.2 \pm 3.5$  cmH<sub>2</sub>O; *mdx* + NAC versus *mdx*) and maximal inspiratory efforts ( $-55.4 \pm 13.3$  versus  $59.7 \pm 11.8$  cmH<sub>2</sub>O; *mdx* + NAC versus *mdx*) were equivalent between *mdx* and *mdx* + NAC mice. Furthermore, NAC supplementation had no effect on diaphragm and external intercostal EMG activity in *mdx* mice during baseline conditions (diaphragm:  $6 \pm 4$  versus  $6 \pm 5$  A.U.; EIC:  $3 \pm 1$  versus  $3 \pm 2$  A.U.; *mdx* + NAC versus *mdx*) and peak inspiratory efforts during airway obstruction (diaphragm:  $30 \pm 17$  versus  $25 \pm 15$  A.U.; EIC:  $18 \pm 13$  versus  $14 \pm 9$  A.U.; *mdx* + NAC versus *mdx*). Decreased maximal respiratory EMG activities in *mdx* mice is likely reflective of neuromuscular junction impairment [81,82], which appears not to have been affected by NAC administration, but this requires further investigation.

Measures of somatic growth such as tibia length and mass were equivalent across all three groups. In terms of organ measures, *mdx* mice had heavier right and left heart ventricles and spleens than age-matched wild type mice. This is consistent with our previous observations [17,42]. Brown and white adipose tissues were decreased in *mdx* compared with wild type; NAC supplementation resulted in a modest increase in brown adipose tissue mass. Collectively, these are interesting

findings in the context of previous research addressing potential side effects of NAC particularly in respect of reductions in body mass [27,75,83]. Left ventricle mass was unaffected by NAC supplementation in *mdx* mice, while right ventricle mass was decreased. This could potentially be a beneficial effect of NAC acting to reduce cardiac muscle hypertrophy or collagen deposition in the right ventricle. Overall, NAC supplementation had no major effects on body mass, indices of somatic growth, organ mass and brown and white adipose tissue mass, demonstrating no adverse effects of our relatively short intervention. Changes in right ventricle and spleen mass were observed and it will be important to discern in future work if these changes are adaptive or maladaptive outcomes in dystrophic tissue in response to NAC. Future work should widen the scope of assessment of the effects of NAC supplementation on physiological systems and incorporate assessments of NAC in other animal models of muscular dystrophy. It will be important to establish the efficacy of NAC supplementation in models of advanced dystrophic disease affecting ventilatory performance.

## 5. Conclusions

In summary, *mdx* mice have diaphragm muscle weakness and myofibre remodelling, immune cell infiltration and collagen deposition. Pro-inflammatory cytokines are increased in diaphragm muscle and plasma samples from *mdx* mice. Supplementation of *mdx* mice with NAC resulted in functional improvements of the diaphragm muscle, immune cell infiltrates and collagen deposits were reduced in the diaphragm, and systemic cytokine concentrations were reduced and diaphragm Nrf2 mRNA expression was increased. There were also sizeable reductions in the concentrations of IL-1 $\beta$  and IL-6 in *mdx* diaphragms following NAC supplementation. These findings are important and relevant to the development of adjunctive therapies with an ambition to alleviate muscle pathology in human dystrophinopathies.

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## References

1. Yiu, E.M.; Kornberg, A.J. Duchenne muscular dystrophy. *Neurol India* **2008**, *56*, 236–247.
2. Petrof, B.J.; Shrager, J.B.; Stedman, H.H.; Kelly, A.M.; Sweeney, H.L. Dystrophin protects the sarcolemma from stresses developed during muscle contraction. *Proc Natl Acad Sci U S A* **1993**, *90*, 3710–3714.
3. Ervasti, J.M.; Campbell, K.P. Membrane organization of the dystrophin-glycoprotein complex. *Cell* **1991**, *66*, 1121–1131.
4. Petrillo, S.; Pelosi, L.; Piemonte, F.; Travaglini, L.; Forcina, L.; Catteruccia, M.; Petrini, S.; Verardo, M.; D'Amico, A.; Musarò, A., et al. Oxidative stress in Duchenne muscular dystrophy: focus on the NRF2 redox pathway. *Hum Mol Genet* **2017**, *26*, 2781–2790, doi:10.1093/hmg/ddx173.
5. Deconinck, N.; Dan, B. Pathophysiology of duchenne muscular dystrophy: current hypotheses. *Pediatr Neurol* **2007**, *36*, 1–7, doi:10.1016/j.pediatrneurol.2006.09.016.
6. Khirani, S.; Ramirez, A.; Aubertin, G.; Boulé, M.; Chemouny, C.; Forin, V.; Fauroux, B. Respiratory muscle decline in Duchenne muscular dystrophy. *Pediatr Pulmonol* **2014**, *49*, 473–481, doi:10.1002/ppul.22847.
7. Mayer, O.H.; Finkel, R.S.; Rummey, C.; Benton, M.J.; Glanzman, A.M.; Flickinger, J.; Lindström, B.M.; Meier, T. Characterization of pulmonary function in Duchenne Muscular Dystrophy. *Pediatr Pulmonol* **2015**, *50*, 487–494, doi:10.1002/ppul.23172.
8. Meier, T.; Rummey, C.; Leinonen, M.; Spagnolo, P.; Mayer, O.H.; Buyse, G.M.; Group, D.S. Characterization of pulmonary function in 10–18 year old patients with Duchenne muscular dystrophy. *Neuromuscul Disord* **2017**, *27*, 307–314, doi:10.1016/j.nmd.2016.12.014.

9. De Bruin, P.F.; Ueki, J.; Bush, A.; Khan, Y.; Watson, A.; Pride, N.B. Diaphragm thickness and inspiratory strength in patients with Duchenne muscular dystrophy. *Thorax* **1997**, *52*, 472–475.
10. Barbé, F.; Quera-Salva, M.A.; McCann, C.; Gajdos, P.; Raphael, J.C.; de Lattre, J.; Agustí, A.G. Sleep-related respiratory disturbances in patients with Duchenne muscular dystrophy. *Eur Respir J* **1994**, *7*, 1403–1408.
11. Toussaint, M.; Davidson, Z.; Bouvoie, V.; Evenepoel, N.; Haan, J.; Soudon, P. Dysphagia in Duchenne muscular dystrophy: practical recommendations to guide management. *Disabil Rehabil* **2016**, *38*, 2052–2062, doi:10.3109/09638288.2015.1111434.
12. Burns, D.P.; Roy, A.; Lucking, E.F.; McDonald, F.B.; Gray, S.; Wilson, R.J.; Edge, D.; O'Halloran, K.D. Sensorimotor control of breathing in the mdx mouse model of Duchenne muscular dystrophy. *J Physiol* **2017**, *595*, 6653–6672, doi:10.1113/JP274792.
13. Burns, D.P.; Murphy, K.H.; Lucking, E.F.; O'Halloran, K.D. Inspiratory pressure-generating capacity is preserved during ventilatory and non-ventilatory behaviours in young dystrophic mdx mice despite profound diaphragm muscle weakness. *J Physiol* **2019**, *597*, 831–848, doi:10.1113/JP277443.
14. Coirault, C.; Pignol, B.; Cooper, R.N.; Butler-Browne, G.; Chabrier, P.E.; Lecarpentier, Y. Severe muscle dysfunction precedes collagen tissue proliferation in mdx mouse diaphragm. *J Appl Physiol (1985)* **2003**, *94*, 1744–1750, doi:10.1152/jappphysiol.00989.2002.
15. Coirault, C.; Lambert, F.; Marchand-Adam, S.; Attal, P.; Chemla, D.; Lecarpentier, Y. Myosin molecular motor dysfunction in dystrophic mouse diaphragm. *Am J Physiol* **1999**, *277*, C1170–1176.
16. Stedman, H.H.; Sweeney, H.L.; Shrager, J.B.; Maguire, H.C.; Panettieri, R.A.; Petrof, B.; Narusawa, M.; Leferovich, J.M.; Sladky, J.T.; Kelly, A.M. The mdx mouse diaphragm reproduces the degenerative changes of Duchenne muscular dystrophy. *Nature* **1991**, *352*, 536–539, doi:10.1038/352536a0.
17. Burns, D.P.; Canavan, L.; Rowland, J.; O'Flaherty, R.; Brannock, M.; Drummond, S.E.; O'Malley, D.; Edge, D.; O'Halloran, K.D. Recovery of respiratory function in mdx mice co-treated with neutralizing interleukin-6 receptor antibodies and urocortin-2. *J Physiol* **2018**, *596*, 5175–5197, doi:10.1113/JP276954.
18. Moraes, L.H.; de Burgos, R.R.; Macedo, A.B.; de Almeida Hermes, T.; de Faria, F.M.; Minatel, E. Reduction of Oxidative Damage and Inflammatory Response in the Diaphragm Muscle of mdx Mice Using Iron Chelator Deferoxamine. *Biol Trace Elem Res* **2015**, *167*, 115–120, doi:10.1007/s12011-015-0290-y.
19. Reid, M.B. Invited Review: redox modulation of skeletal muscle contraction: what we know and what we don't. *J Appl Physiol* **2001**, *90*, 724–731.
20. O'Halloran, K.D.; Lewis, P. Respiratory muscle dysfunction in animal models of hypoxic disease: antioxidant therapy goes from strength to strength. *Hypoxia (Auckl)* **2017**, *5*, 75–84, doi:10.2147/HP.S141283.
21. Lawler, J.M. Exacerbation of pathology by oxidative stress in respiratory and locomotor muscles with Duchenne muscular dystrophy. *J Physiol* **2011**, *589*, 2161–2170, doi:10.1113/jphysiol.2011.207456.
22. Burns, D.P.; Ali, I.; Rieux, C.; Healy, J.; Jasione, G.; O'Halloran, K.D. Tempol Supplementation Restores Diaphragm Force and Metabolic Enzyme Activities in mdx Mice. *Antioxidants*, **2017**, *6*, 101, doi:10.3390/antiox6040101.
23. Hermes, T.A.; Mâncio, R.D.; Macedo, A.B.; Mizobuti, D.S.; Rocha, G.L.D.; Cagnon, V.H.A.; Minatel, E. Tempol treatment shows phenotype improvement in mdx mice. *PLoS One* **2019**, *14*, e0215590, doi:10.1371/journal.pone.0215590.
24. O'Leary, A.J.; Drummond, S.E.; Edge, D.; O'Halloran, K.D. Diaphragm Muscle Weakness Following Acute Sustained Hypoxic Stress in the Mouse Is Prevented by Pretreatment with N-Acetyl Cysteine. *Oxid Med Cell Longev* **2018**, *2018*, 4805493, doi:10.1155/2018/4805493.
25. Lewis, P.; Sheehan, D.; Soares, R.; Coelho, A.V.; O'Halloran, K.D. Redox Remodeling Is Pivotal in Murine Diaphragm Muscle Adaptation to Chronic Sustained Hypoxia. *Am J Respir Cell Mol Biol* **2016**, *55*, 12–23, doi:10.1165/rcmb.2015-0272OC.
26. Shortt, C.M.; Fredsted, A.; Chow, H.B.; Williams, R.; Skelly, J.R.; Edge, D.; Bradford, A.; O'Halloran, K.D. Reactive oxygen species mediated diaphragm fatigue in a rat model of chronic intermittent hypoxia. *Exp Physiol* **2014**, *99*, 688–700, doi:10.1113/expphysiol.2013.076828.
27. Pinniger, G.J.; Terrill, J.R.; Assan, E.B.; Grounds, M.D.; Arthur, P.G. Pre-clinical evaluation of N-acetylcysteine reveals side effects in the mdx mouse model of Duchenne muscular dystrophy. *J Physiol* **2017**, *595*, 7093–7107, doi:10.1113/JP274229.
28. de Senzi Moraes Pinto, R.; Ferretti, R.; Moraes, L.H.; Neto, H.S.; Marques, M.J.; Minatel, E. N-acetylcysteine treatment reduces TNF- $\alpha$  levels and myonecrosis in diaphragm muscle of mdx mice. *Clin Nutr* **2013**, *32*, 472–475, doi:10.1016/j.clnu.2012.06.001.
29. Terrill, J.R.; Radley-Crabb, H.G.; Grounds, M.D.; Arthur, P.G. N-Acetylcysteine treatment of dystrophic mdx mice results in protein thiol modifications and inhibition of exercise induced myofibre necrosis. *Neuromuscul Disord* **2012**, *22*, 427–434, doi:10.1016/j.nmd.2011.11.007.



30. Whitehead, N.P.; Pham, C.; Gervasio, O.L.; Allen, D.G. N-Acetylcysteine ameliorates skeletal muscle pathophysiology in mdx mice. *J Physiol* **2008**, *586*, 2003–2014, doi:10.1113/jphysiol.2007.148338.
31. Grundy, D. Principles and standards for reporting animal experiments in The Journal of Physiology and Experimental Physiology. *J Physiol* **2015**, *593*, 2547–2549, doi:10.1113/JP270818.
32. Burns, D.P.; O'Halloran, K.D. Evidence of hypoxic tolerance in weak upper airway muscle from young mdx mice. *Respir Physiol Neurobiol* **2016**, *226*, 68–75, doi:10.1016/j.resp.2015.12.001.
33. Burns, D.P.; Rowland, J.; Canavan, L.; Murphy, K.H.; Brannock, M.; O'Malley, D.; O'Halloran, K.D.; Edge, D. Restoration of pharyngeal dilator muscle force in dystrophin-deficient (mdx) mice following co-treatment with neutralizing interleukin-6 receptor antibodies and urocortin 2. *Exp Physiol* **2017**, *102*, 1177–1193, doi:10.1113/EP086232.
34. Lewis, P.; Sheehan, D.; Soares, R.; Varela Coelho, A.; O'Halloran, K.D. Chronic sustained hypoxia-induced redox remodeling causes contractile dysfunction in mouse sternohyoid muscle. *Front Physiol* **2015**, *6*, 122, doi:10.3389/fphys.2015.00122.
35. Schiaffino, S.; Rossi, A.C.; Smerdu, V.; Leinwand, L.A.; Reggiani, C. Developmental myosins: expression patterns and functional significance. *Skelet Muscle* **2015**, *5*, 22, doi:10.1186/s13395-015-0046-6.
36. Noirez, P.; Torres, S.; Cebrian, J.; Agbulut, O.; Peltzer, J.; Butler-Browne, G.; Daegelen, D.; Martelly, I.; Keller, A.; Ferry, A. TGF-beta1 favors the development of fast type identity during soleus muscle regeneration. *J Muscle Res Cell Motil* **2006**, *27*, 1–8, doi:10.1007/s10974-005-9014-9.
37. LoMauro, A.; D'Angelo, M.G.; Aliverti, A. Sleep Disordered Breathing in Duchenne Muscular Dystrophy. *Curr Neurol Neurosci Rep* **2017**, *17*, 44, doi:10.1007/s11910-017-0750-1.
38. Smith, P.E.; Edwards, R.H.; Calverley, P.M. Ventilation and breathing pattern during sleep in Duchenne muscular dystrophy. *Chest* **1989**, *96*, 1346–1351.
39. Kirk, V.G.; Flemons, W.W.; Adams, C.; Rimmer, K.P.; Montgomery, M.D. Sleep-disordered breathing in Duchenne muscular dystrophy: a preliminary study of the role of portable monitoring. *Pediatr Pulmonol* **2000**, *29*, 135–140.
40. Khan, Y.; Heckmatt, J.Z. Obstructive apnoeas in Duchenne muscular dystrophy. *Thorax* **1994**, *49*, 157–161.
41. Suresh, S.; Wales, P.; Dakin, C.; Harris, M.A.; Cooper, D.G. Sleep-related breathing disorder in Duchenne muscular dystrophy: disease spectrum in the paediatric population. *J Paediatr Child Health* **2005**, *41*, 500–503, doi:10.1111/j.1440-1754.2005.00691.x.
42. Burns, D.P.; Edge, D.; O'Malley, D.; O'Halloran, K.D. Respiratory Control in the mdx Mouse Model of Duchenne Muscular Dystrophy. *Adv Exp Med Biol* **2015**, *860*, 239–244, doi:10.1007/978-3-319-18440-1\_27.
43. Mosqueira, M.; Baby, S.M.; Lahiri, S.; Khurana, T.S. Ventilatory chemosensory drive is blunted in the mdx mouse model of Duchenne Muscular Dystrophy (DMD). *PLoS One* **2013**, *8*, e69567, doi:10.1371/journal.pone.0069567.
44. Gayraud, J.; Matecki, S.; Hnia, K.; Mornet, D.; Prefaut, C.; Mercier, J.; Michel, A.; Ramonatxo, M. Ventilation during air breathing and in response to hypercapnia in 5 and 16 month-old mdx and C57 mice. *J Muscle Res Cell Motil* **2007**, *28*, 29–37, doi:10.1007/s10974-007-9101-1.
45. Attal, P.; Lambert, F.; Marchand-Adam, S.; Bobin, S.; Pourny, J.C.; Chemla, D.; Lecarpentier, Y.; Coirault, C. Severe mechanical dysfunction in pharyngeal muscle from adult mdx mice. *Am J Respir Crit Care Med* **2000**, *162*, 278–281.
46. Sieck, G.C.; Ferreira, L.F.; Reid, M.B.; Mantilla, C.B. Mechanical properties of respiratory muscles. *Compr Physiol* **2013**, *3*, 1553–1567, doi:10.1002/cphy.c130003.
47. Mantilla, C.B.; Seven, Y.B.; Sieck, G.C. Convergence of pattern generator outputs on a common mechanism of diaphragm motor unit recruitment. *Prog Brain Res* **2014**, *209*, 309–329, doi:10.1016/B978-0-444-63274-6.00016-3.
48. O'Halloran, K.D.; Burns, D.P. Breathing with neuromuscular disease: Does compensatory plasticity in the motor drive to breathe offer a potential therapeutic target in muscular dystrophy? *Respir Physiol Neurobiol* **2019**, *265*, 49–54, doi:10.1016/j.resp.2018.06.009.
49. Barros Maranhão, J.; de Oliveira Moreira, D.; Maurício, A.F.; de Carvalho, S.C.; Ferretti, R.; Pereira, J.A.; Santo Neto, H.; Marques, M.J. Changes in calsequestrin, TNF- $\alpha$ , TGF- $\beta$  and MyoD levels during the progression of skeletal muscle dystrophy in mdx mice: a comparative analysis of the quadriceps, diaphragm and intrinsic laryngeal muscles. *Int J Exp Pathol* **2015**, *96*, 285–293, doi:10.1111/iep.12142.
50. Gutpell, K.M.; Hrinivich, W.T.; Hoffman, L.M. Skeletal muscle fibrosis in the mdx/utrn $\pm$  mouse validates its suitability as a murine model of Duchenne muscular dystrophy. *PLoS One* **2015**, *10*, e0117306, doi:10.1371/journal.pone.0117306.
51. Pescatori, M.; Broccolini, A.; Minetti, C.; Bertini, E.; Bruno, C.; D'amico, A.; Bernardini, C.; Mirabella, M.; Silvestri, G.; Giglio, V., et al. Gene expression profiling in the early phases of DMD: a constant molecular

- signature characterizes DMD muscle from early postnatal life throughout disease progression. *FASEB J* **2007**, *21*, 1210-1226, doi:10.1096/fj.06-7285com.
52. De Paepe, B.; Creus, K.K.; Martin, J.J.; De Bleecker, J.L. Upregulation of chemokines and their receptors in Duchenne muscular dystrophy: potential for attenuation of myofiber necrosis. *Muscle Nerve* **2012**, *46*, 917-925, doi:10.1002/mus.23481.
  53. Messina, S.; Vita, G.L.; Aguenouz, M.; Sframeli, M.; Romeo, S.; Rodolico, C.; Vita, G. Activation of NF-kappaB pathway in Duchenne muscular dystrophy: relation to age. *Acta Myol* **2011**, *30*, 16-23.
  54. Cruz-Guzmán, O.e.R.; Rodríguez-Cruz, M.; Escobar Cedillo, R.E. Systemic Inflammation in Duchenne Muscular Dystrophy: Association with Muscle Function and Nutritional Status. *Biomed Res Int* **2015**, *2015*, 891972, doi:10.1155/2015/891972.
  55. Eghtesad, S.; Jhunjhunwala, S.; Little, S.R.; Clemens, P.R. Rapamycin ameliorates dystrophic phenotype in mdx mouse skeletal muscle. *Mol Med* **2011**, *17*, 917-924, doi:10.2119/molmed.2010.00256.
  56. Serra, F.; Quarta, M.; Canato, M.; Toniolo, L.; De Arcangelis, V.; Trotta, A.; Spath, L.; Monaco, L.; Reggiani, C.; Naro, F. Inflammation in muscular dystrophy and the beneficial effects of non-steroidal anti-inflammatory drugs. *Muscle Nerve* **2012**, *46*, 773-784, doi:10.1002/mus.23432.
  57. Pelosi, L.; Berardinelli, M.G.; De Pasquale, L.; Nicoletti, C.; D'Amico, A.; Carvello, F.; Moneta, G.M.; Catizone, A.; Bertini, E.; De Benedetti, F., et al. Functional and Morphological Improvement of Dystrophic Muscle by Interleukin 6 Receptor Blockade. *EBioMedicine* **2015**, *2*, 285-293, doi:10.1016/j.ebiom.2015.02.014.
  58. Manning, J.; Buckley, M.M.; O'Halloran, K.D.; O'Malley, D. Combined xIL-6R and urocortin-2 treatment restores mdx diaphragm muscle force. *Muscle Nerve* **2017**, *56*, E134-140, doi:10.1002/mus.25644.
  59. Pelosi, L.; Forcina, L.; Nicoletti, C.; Scicchitano, B.M.; Musarò, A. Increased Circulating Levels of Interleukin-6 Induce Perturbation in Redox-Regulated Signaling Cascades in Muscle of Dystrophic Mice. *Oxid Med Cell Longev* **2017**, *2017*, 1987218, doi:10.1155/2017/1987218.
  60. Tidball, J.G. Inflammatory processes in muscle injury and repair. *Am J Physiol Regul Integr Comp Physiol* **2005**, *288*, R345-353, doi:10.1152/ajpregu.00454.2004.
  61. Mechler, F.; Imre, S.; Dioszeghy, P. Lipid peroxidation and superoxide dismutase activity in muscle and erythrocytes in Duchenne muscular dystrophy. *J Neurol Sci* **1984**, *63*, 279-283.
  62. Burr, I.M.; Asayama, K.; Fenichel, G.M. Superoxide dismutases, glutathione peroxidase, and catalase in neuromuscular disease. *Muscle Nerve* **1987**, *10*, 150-154, doi:10.1002/mus.880100208.
  63. Khouzami, L.; Bourin, M.C.; Christov, C.; Damy, T.; Escoubet, B.; Caramelle, P.; Perier, M.; Wahbi, K.; Meune, C.; Pavoiné, C., et al. Delayed cardiomyopathy in dystrophin deficient mdx mice relies on intrinsic glutathione resource. *Am J Pathol* **2010**, *177*, 1356-1364, doi:10.2353/ajpath.2010.090479.
  64. Renjini, R.; Gayathri, N.; Nalini, A.; Srinivas Bharath, M.M. Oxidative damage in muscular dystrophy correlates with the severity of the pathology: role of glutathione metabolism. *Neurochem Res* **2012**, *37*, 885-898, doi:10.1007/s11064-011-0683-z.
  65. Kim, J.H.; Kwak, H.B.; Thompson, L.V.; Lawler, J.M. Contribution of oxidative stress to pathology in diaphragm and limb muscles with Duchenne muscular dystrophy. *J Muscle Res Cell Motil* **2013**, *34*, 1-13, doi:10.1007/s10974-012-9330-9.
  66. Dudley, R.W.; Danelou, G.; Govindaraju, K.; Lands, L.; Eidelman, D.E.; Petrof, B.J. Sarcolemmal damage in dystrophin deficiency is modulated by synergistic interactions between mechanical and oxidative/nitrosative stresses. *Am J Pathol* **2006**, *168*, 1276-1287, doi:10.2353/ajpath.2006.050683.
  67. Whitehead, N.P.; Yeung, E.W.; Allen, D.G. Muscle damage in mdx (dystrophic) mice: role of calcium and reactive oxygen species. *Clin Exp Pharmacol Physiol* **2006**, *33*, 657-662, doi:10.1111/j.1440-1681.2006.04394.x.
  68. Whitehead, N.P.; Yeung, E.W.; Froehner, S.C.; Allen, D.G. Skeletal muscle NADPH oxidase is increased and triggers stretch-induced damage in the mdx mouse. *PLoS One* **2010**, *5*, e15354, doi:10.1371/journal.pone.0015354.
  69. Loehr, J.A.; Wang, S.; Cully, T.R.; Pal, R.; Larina, I.V.; Larin, K.V.; Rodney, G.G. NADPH oxidase mediates microtubule alterations and diaphragm dysfunction in dystrophic mice. *eLife* **2018**, *7*, e31732, doi:10.7554/eLife.31732.
  70. Whitehead, N.P.; Kim, M.J.; Bible, K.L.; Adams, M.E.; Froehner, S.C. A new therapeutic effect of simvastatin revealed by functional improvement in muscular dystrophy. *Proc Natl Acad Sci U S A* **2015**, *112*, 12864-12869, doi:10.1073/pnas.1509536112.
  71. El-Shafey, A.F.; Armstrong, A.E.; Terrill, J.R.; Grounds, M.D.; Arthur, P.G. Screening for increased protein thiol oxidation in oxidatively stressed muscle tissue. *Free Radic Res* **2011**, *45*, 991-999, doi:10.3109/10715762.2011.590136.
  72. Dunleavy, M.; Bradford, A.; O'Halloran, K.D. Oxidative stress impairs upper airway muscle endurance in an animal model of sleep-disordered breathing. *Adv Exp Med Biol* **2008**, *605*, 458-462, doi:10.1007/978-0-387-73693-8\_80.

73. Sadowska, A.M.; Verbraecken, J.; Darquennes, K.; De Backer, W.A. Role of N-acetylcysteine in the management of COPD. *Int J Chron Obstruct Pulmon Dis* **2006**, *1*, 425-434.
74. Travalline, J.M.; Sudarshan, S.; Roy, B.G.; Cordova, F.; Leyenson, V.; Criner, G.J. Effect of N-acetylcysteine on human diaphragm strength and fatigability. *Am J Respir Crit Care Med* **1997**, *156*, 1567-1571, doi:10.1164/ajrccm.156.5.96-09133.
75. O'Halloran, K.D.; Murphy, K.H.; Burns, D.P. Antioxidant therapy for muscular dystrophy: caveat lector! *J Physiol* **2018**, *596*, 737-738, doi:10.1113/JP275598.
76. Kopp, J.; Seyhan, H.; Müller, B.; Lanczak, J.; Pausch, E.; Gressner, A.M.; Dooley, S.; Horch, R.E. N-acetyl-L-cysteine abrogates fibrogenic properties of fibroblasts isolated from Dupuytren's disease by blunting TGF-beta signalling. *J Cell Mol Med* **2006**, *10*, 157-165.
77. Giam, B.; Chu, P.Y.; Kuruppu, S.; Smith, A.I.; Horlock, D.; Kiriazis, H.; Du, X.J.; Kaye, D.M.; Rajapakse, N.W. N-acetylcysteine attenuates the development of cardiac fibrosis and remodeling in a mouse model of heart failure. *Physiol Rep* **2016**, *4*, e12757, doi:10.14814/phy2.12757.
78. Roseguini, B.T.; Silva, L.M.; Polotow, T.G.; Barros, M.P.; Souccar, C.; Han, S.W. Effects of N-acetylcysteine on skeletal muscle structure and function in a mouse model of peripheral arterial insufficiency. *J Vasc Surg* **2015**, *61*, 777-786, doi:10.1016/j.jvs.2013.10.098.
79. Giam, B.; Kuruppu, S.; Chu, P.Y.; Smith, A.I.; Marques, F.Z.; Fiedler, A.; Horlock, D.; Kiriazis, H.; Du, X.J.; Kaye, D.M., et al. N-Acetylcysteine Attenuates the Development of Renal Fibrosis in Transgenic Mice with Dilated Cardiomyopathy. *Sci Rep* **2017**, *7*, 17718, doi:10.1038/s41598-017-17927-5.
80. Burns, D.P.; Lucking, E.F.; O'Halloran, K.D. Reply from David P. Burns, Eric F. Lucking and Ken D. O'Halloran: Auxiliary compensation for diaphragm dysfunction in dystrophic disease. *J Physiol* **2019**, *597*, 4103-4105, doi:10.1113/JP278371.
81. Pratt, S.J.; Valencia, A.P.; Le, G.K.; Shah, S.B.; Lovering, R.M. Pre- and postsynaptic changes in the neuromuscular junction in dystrophic mice. *Front Physiol* **2015**, *6*, 252, doi:10.3389/fphys.2015.00252.
82. van der Pijl, E.M.; van Putten, M.; Niks, E.H.; Verschuuren, J.J.; Aartsma-Rus, A.; Plomp, J.J. Characterization of neuromuscular synapse function abnormalities in multiple Duchenne muscular dystrophy mouse models. *Eur J Neurosci* **2016**, *43*, 1623-1635, doi:10.1111/ejn.13249.
83. Head, S.I. Antioxidant therapy in a mouse model of Duchenne muscular dystrophy: some promising results but with a weighty caveat. *J Physiol* **2017**, *595*, 7015, doi:10.1113/JP275232.



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