Title

*Cmah*-dystrophin deficient *mdx* mice display an accelerated cardiac phenotype that is improved following peptide-PMO exon skipping treatment

Authors

Corinne A Betts¹, Graham McClorey¹, Richard Healicon¹, Suzan M Hammond¹, Raquel Manzano¹, Sofia Muses², Vicky Ball¹, Caroline Godfrey¹, Thomas M Merritt¹, Tirsa van Westering¹, Liz O’Donovan⁴, Kim E Wells², Michael J Gait⁴, Dominic J Wells², Damian Tyler¹, Matthew J Wood¹*

¹Department of Physiology, Anatomy and Genetics, University of Oxford, South Parks Road, Oxford, OX1 3QX, United Kingdom.

²Department of Comparative Biomedical Sciences, Royal Veterinary College, Royal College Street, London, NW1 0TU, United Kingdom.

³Clinical Biomanufacturing Facility, Nuffield Department of Clinical Medicine, University of Oxford, Old Road, Oxford, OX3 7JT, United Kingdom

⁴Medical Research Council, Laboratory of Molecular Biology, Francis Crick Avenue, Cambridge CB2 0QH, UK

*To whom correspondence should be addressed: Professor Matthew J Wood, Department of Physiology, Anatomy and Genetics, University of Oxford, South Parks Road, Oxford, OX1 3QX, United Kingdom, +44(0) 1865 272 419, matthew.wood@dpag.ox.ac.uk
Abstract

Duchenne muscular dystrophy (DMD) is caused by loss of dystrophin protein, leading to progressive muscle weakness and premature death due to respiratory and/or cardiac complications. Cardiac involvement is characterised by progressive dilated cardiomyopathy, decreased fractional shortening and metabolic dysfunction involving reduced metabolism of fatty acids - the major cardiac metabolic substrate. Several mouse models have been developed to study molecular and pathological consequences of dystrophin deficiency, but do not recapitulate all aspects of human disease pathology and exhibit a mild cardiac phenotype. Here we demonstrate that Cmah-deficient mdx mice (Cmah-/-;mdx) have an accelerated cardiac phenotype compared to the established mdx model. Cmah-/-;mdx mice display earlier functional deterioration, specifically a reduction in right ventricle ejection fraction and stroke volume at 12 weeks of age, and decreased left ventricle diastolic volume with subsequent reduced stroke volume compared to mdx mice by 24 weeks. They further show earlier elevation of cardiac damage markers for fibrosis (Ctgf), oxidative damage (Nox4) and haemodynamic load (Nppa). Cardiac metabolic substrate requirement was assessed using hyperpolarised-MRS indicating increased in vivo glycolytic flux in Cmah-/-;mdx mice. Early upregulation of mitochondrial genes (Ucp3 and Cpt1), and down-regulation of key glycolytic genes (Pdk1, Pdk4, Ppara), also denote disturbed cardiac metabolism and shift towards glucose utilisation in Cmah-/-;mdx mice. Moreover, we show long-term treatment with peptide-conjugated exon skipping antisense oligonucleotides (20 week regimen), resulted in 20% cardiac dystrophin protein restoration and significantly improved right ventricle cardiac function. Therefore Cmah-/-;mdx mice represent an appropriate model for evaluating cardiac benefit of novel DMD therapeutics.

Introduction

Duchenne muscular dystrophy (DMD) is caused by the loss of dystrophin protein due to genetic defects in the DMD gene(1). This results in rapidly progressing muscle wasting with subsequent early
loss of ambulation(2) and death in the third or fourth decade of life. As the standard of care has
improved for patients, predominantly through the use of corticosteroids to delay disease
progression(3) and the use of assisted ventilation to preserve respiratory function(4); men are living
longer and as such, secondary pathologies such as cardiomyopathy have become important factors to
consider for disease management. Cardiomyopathy in DMD patients first presents with left ventricle
(LV) dilation and hypertrophy accompanied by decreased fractional shortening and electrocardiogram
(ECG) abnormalities(5-8). As the disease progresses, patients exhibit dilated cardiomyopathy, and by
20 years of age most DMD patients suffer from cardiac complications(5). Cardiomyopathy is a major
contributor of death amongst DMD patients(9) and therefore multiple approaches have been
implemented to treat the cardiac phenotype, including corticosteroids, beta-blockers, ACE inhibitors,
and antimineralocorticoid diuretics(10).

The heart is unique in its requirement for metabolic substrate particularly compared to skeletal
muscle. The heart predominantly relies on free fatty acids for approximately 70% of its energy
requirements and for the remainder on glucose reserves(11). When the heart undergoes a shift in
metabolic substrate requirement, for instance an increase in glucose utilisation, this is generally
deemed pathological, as is the case in heart failure patients. Disruption of lipid handling in skeletal
muscle of DMD patients is well characterised and includes the accumulation of cholesterol,
sphingomyelin and triglycerides, as well as high levels of monosaturated fatty acid in damaged areas
of muscle(12). As cardiac biopsies are not feasible, assessment of cardiac metabolic alterations is
more difficult in patients. However, there is one study that utilised radioiodinated branched fatty acid
$^{123}$I-BMIPP for in vivo measurement of myocardial metabolism in DMD patients(13). Momose et al.
showed a reduction in myocardial fatty acid metabolism in a substantial proportion of patients.

To understand the molecular and pathological effects of the loss of dystrophin protein, a number of
mouse models of DMD are available, including the mdx mouse which has a premature termination
codon in exon 23 that results in loss of the full-length dystrophin isoform(14). The cardiac phenotype
in this model is well characterised, typified by right ventricle (RV) dysfunction preceding LV
involvement; although it should be noted that generally LV function deteriorates prior to RV in DMD patients(15, 16). Multiple studies also report altered proteomic profile of key metabolic proteins, in particular mitochondrial proteins associated with the electron transport chain and ATP synthesis(17, 18). Further to this, alterations in metabolic flux in isolated \( mdx \) hearts was observed as early as 12 weeks of age(19), which is prior to obvious histological and functional changes(15, 20). Specifically, \( mdx \) mouse hearts displayed a marked shift from long chain fatty acid to carbohydrate oxidation for energy requirements, a decrease in utilisation of the pyruvate carboxylation pathway, and enhanced glycolysis.

More recently, another DMD mouse model, the \( Cmah-/\);\( mdx \) mouse, has been generated(21). In this model, the \( Cmah \) gene (cytidine monophosphate-sialic acid hydroxylase), which is also absent in humans, is deleted on the \( mdx \) background, resulting in a more severe phenotype representative of the human disease. This included advanced inflammation and fibrosis, reduced specific force in muscle, increased susceptibility to contraction/exercise induced damage, and increased mortality rates. The characterisation of this model did not report on cardiac disease progression, and given that other measures of disease were worsened, we hypothesised that this model may be more appropriate to assess cardiac defects than the \( mdx \) model. We therefore sought to assess cardiac function by magnetic resonance imaging (MRI) and cardiac metabolic flux \textit{in vivo} using hyperpolarised magnetic resonance spectroscopy (MRS) technology. Additionally, we sought to assess the benefit of dystrophin restoration in this model utilising a splice-switching approach. This model contains a premature termination codon at exon 23, therefore we target the removal of exon 23 so as to restore near full-length dystrophin protein. To achieve this we used cell-penetrating peptide conjugated morpholinos, specifically Pip6A-PMO, that have been used in previous studies in \( mdx \) mice to achieve robust dystrophin expression in skeletal muscle and in the heart (22, 23).

**Results**
Marked cardiac functional changes and damage in mdx and Cmah-/-;mdx mice from 12 weeks of age.

Mouse weights and cardiac function (cine-MRI) in C57LB10, mdx, and Cmah-/-;mdx male mice were recorded at 12 and 24 weeks of age. At 12 weeks there was no significant difference in body weight between genotypes; however at 24 weeks, body weight was significantly raised in Cmah-/-;mdx mice compared to control and mdx mice, with mdx mice also significantly heavier than controls (Table 1). Multiple parameters including systolic and diastolic volumes, stroke volume and cardiac output were normalised to body weight. Figure 1a shows representative stills for each cohort at diastole and systole for 12 and 24 week old mice. As previously observed, there were no significant functional changes in the mdx cohort compared to C57BL10 mice at 12 weeks of age, with the exception of LV cardiac output (LV CO; Table 1) as previously reported(15, 22). Interestingly, Cmah-/-;mdx mice exhibited reduced RV ejection fraction (RV EF; Fig 1b), lower RV stroke volume (RV SV; Fig 1b) with resultant lower cardiac output (RV CO; Table 1) compared to C57BL10 at 12 weeks. At 24 weeks of age, the mdx mice revealed a smaller average LV mass and changes in RV function including reduced RV EF, lowered RV end systolic volume (RV ESV) and RV SV, leading to reduced RV CO (Table 1). In addition to marked RV changes, Cmah-/-;mdx mice also exhibited reduced LV end diastolic volume (LV EDV), LV SV and LV CO (Fig 1b and Table 1). The heart rate of Cmah-/-;mdx mice was also significantly elevated. Further to functional abnormalities, Cmah-/-;mdx mice also display elevated gene expression of the fibrosis marker Connective tissue growth factor (Ctgf; Fig 1c) at 12 and 24 weeks of age. Markers of cardiac damage/haemodynamic load (oxidative stress (NADPH oxidase 4; Nox4) and natriuretic peptide precursor A; Nppa(24, 25)), were also significantly upregulated in Cmah-/-;mdx mice at these time points. It is noteworthy to add that in mdx mice the expression of Ctgf and Nppa was only significantly elevated at 24 weeks of age. Nox4 expression was elevated in mdx mice from 12 weeks of age but was significantly lower than the Cmah-/-;mdx cohort. This data is complimented by histology showing marked fibrosis at the LV and RV walls of Cmah-/-;mdx mouse hearts at 12 weeks of age compared to mdx and C57BL10 cohorts (Supp Fig 1).
In vivo metabolic perturbations in Cmah-/-;mdx mice at 24 weeks

Hyperpolarized magnetic resonance spectroscopy was used to determine metabolic substrate allocation by measuring $^{13}$C labelled pyruvate flux through pyruvate dehydrogenase (PDH) and the citric acid cycle(26). This was measured in the hearts of 12 and 24 week old mdx and Cmah-/-;mdx male mice compared to C57LB10 controls. The rate of bicarbonate production at 24 weeks was significantly raised in Cmah-/-;mdx mice compared to the C57BL10 (Fig 2a), indicating increased pyruvate flux through PDH. Although lactate levels were unchanged by genotype, age had a highly significant effect, with lactate production declining from 12 to 24 weeks of age across all genotypes. No statistically significant differences in alanine production were observed between any of the test groups at either time point (Supp Fig. 2).

The expression levels of key metabolic genes involved in glucose and fatty acid metabolism were also altered. Pyruvate dehydrogenase lipoamide kinase isozymes 1 and 4 (Pdk1 and Pdk4), were significantly downregulated in the Cmah-/-;mdx mouse cohort at 24 weeks of age (Fig 2b; also downregulated in mdx for Pdk4). These kinases act to prevent the conversion of pyruvate from glucose, and thus a decrease in gene expression would suggest a shift toward glucose metabolism(27). Mitochondrial uncoupling protein 3 (Ucp3) is involved in the transfer of anions across the mitochondrial membrane in order to protect against oxidative stress(28). It is markedly upregulated at 12 weeks in Cmah-/-;mdx mice, suggesting a spike in oxidative stress. Carnitine palmitoyltransferase 1 (Cpt1) is a mitochondrial enzyme involved in fatty acid metabolism. Its expression was unexpectedly upregulated in Cmah-/-;mdx mice at 12 weeks, however could be a compensatory mechanism attempting to catabolise fatty acids. Another important membrane protein, cluster of differentiation 36 (CD36), allows the importation of fatty acids, and is significantly down-regulated in both the mdx and Cmah-/-;mdx mice cohorts at 24 weeks. Peroxisome proliferator-activated receptor alpha (Ppara) is a key transcription factor which regulates lipid metabolism, and is also significantly downregulated at 24 weeks of age. The downregulation of both CD36 and Ppara support the
hypothesis for a reduction in fatty acid metabolism. Indeed, circulating levels of total cholesterol, high-density lipoproteins (HDL) and low-density lipoproteins (LDL) were all elevated in Cmah-/-;mdx mice compared to C57BL10 and mdx cohorts, suggesting that these lipid reserves are not being utilised (Fig 3 and Supp Table 1). Free fatty acid levels were significantly raised in mdx mice at 12 weeks of age, and in both mdx and Cmah-/-;mdx mice at 24 weeks.

Chronic PPMO treatment moderately improves right ventricle function in Cmah-/-;mdx

To determine the treatment benefit of an exon-skipping approach to restore dystrophin in dystrophic hearts, 12 week old Cmah-/-;mdx male mice (n=7) were administered 10 x 10mg/kg Pip6A-PMO (Pip-PMO) fortnightly over a 20 week period. Approximately 18% dystrophin protein restoration was detected in heart by western blot, concurrent with moderate exon 23 skipping (Fig 4a, Supp Fig 3). Immunohistochemistry also confirmed correct localisation of dystrophin to the sarcolemma and widespread dystrophin positive fibres throughout the heart (Supp Fig 4).

Cardiac function in C57LB10, mdx, Cmah-/-;mdx and Cmah-/-;mdx treated mice was recorded using cine-MRI at 8 months of age. There was no significant difference in body weight between genotypes (Table 2), however the heart rate in Cmah-/-;mdx cohort was significantly raised compared to all other cohorts, suggesting compensatory activation of the sympathetic pathway (Fig 4b). In addition, the RV ESV was markedly larger than the control cohort, resulting in decreased RV EF. However, Pip-PMO treatment restored heart rate and RV ESV to control levels. Interestingly, gene expression analysis of key metabolic factors at 36 weeks, indicate no significant difference between Cmah-/-;mdx and Cmah-/-;mdx treated cohorts. However, Nppa (trending) and Nox4 were significantly down-regulated in treated Cmah-/-;mdx cohorts indicating a reduction in damage and oxidative species (Fig 4c). However, it should be noted that histological data shows marked fibrosis in the hearts of treated Cmah-/-;mdx, mdx and Cmah-/-;mdx mouse hearts at 8 months of age (Supp Fig 5).
Chronic PPMO treatment induces robust dystrophin expression and improves and protects muscle function

Whilst the primary focus of this study was to evaluate cardiac function in this model, we also sought to determine if Pip-PMO mediated benefit to muscle function could be conferred by assessing resistance to eccentric contraction-induced damage in treated and untreated littermate controls (Fig 5a). Untreated Cmah-/-;mdx mice exhibited a significant final drop in force of 33% compared with initial baseline force (p<0.001), although this was not as pronounced as levels observed for mdx mice of ~60% observed in previous studies (29). Administration of Pip-PMO prevented this force drop with maximal force production maintained throughout the eccentric contraction protocol, similar to wild-type C57BL10 mice (not shown). Specific isometric force was also significantly improved (p<0.001) following Pip-PMO treatment with a 71% improvement towards wild-type levels compared to untreated Cmah-/-;mdx controls (Fig 5b). Robust dystrophin protein restoration was detected in tibialis anterior (TA), diaphragm and heart, concurrent with high levels of exon 23 skipping which correlated to relative levels of correction in the different tissues (Fig 4a). Immunohistochemistry also confirmed correct localisation of dystrophin to the sarcolemma and widespread dystrophin positive fibres throughout quadriceps and diaphragm (Supp Fig 4). Muscle hypertrophy was also assessed with physiological cross-sectional area of the TA significantly reduced in Pip-PMO treated animals, indicative of reduced muscle hypertrophy (Supp Table 2).

Discussion

The predominant aim of this study was to evaluate cardiac dysfunction and metabolic profile in Cmah-/-;mdx and mdx mice to determine to what extent these models recapitulate the cardiac dystrophic pathology in DMD. Furthermore we sought to determine if Pip-PMO mediated exon skipping could confer functional benefit in the Cmah-/-;mdx model.
We show significant changes in functional cardiac parameters in the Cmah-/-;mdx model compared to C57BL10. We report alterations in multiple RV parameters as early as 12 weeks of age, and prominent LV involvement by 24 weeks, which is markedly earlier then the mdx model(15, 23). RV involvement does not generally precede LV dysfunction in patients, though it should be noted that DMD patients undergo rapid respiratory decline, and therefore ventilatory intervention is administered early. It is believed this improves pulmonary hypertension thus compensating for RV dysfunction(30, 31). Indeed the decline in RV function in DMD patients does correlate with a reduction in respiratory function(32). This could account for marked disparity observed between DMD patients and the Cmah-/-;mdx and mdx models which do not receive ventilatory assistance. However most importantly, Cmah-/-;mdx mice do exhibit a much earlier and more pronounced cardiac phenotype compared to mdx. These functional events also correlate with the elevation of multiple markers of pathology and damage. We are also the first to describe functional cardiac metabolic alterations in the Cmah-/-;mdx mouse, which again correlate with the down regulation of key fatty acid-handling genes, and glucose pathway mediators. These data support previous reports in mdx mice(19) that show the dystrophic heart has a preference for glucose utilization, which is an important pathological indicator. Although we did observe significant changes in metabolic flux in Cmah-/-;mdx mice, we did not detect alterations in mdx hearts at 12 weeks of age as previously reported(19). It is important to note that our measurements were performed in vivo, but Khairallah et al experiments were performed ex vivo. It is likely this discrepancy is due to systemic compensatory mechanisms attempting to reinstate a homeostatic state, and it is these events which are masking the underlying pathology of the mdx heart. However, most importantly we were able to detect significant changes in the Cmah-/-;mdx heart, indicating this is a more appropriate model for studying early metabolic perturbations. In short, the Cmah-/-;mdx model exhibits an earlier cardiac phenotype as determined by deterioration in heart function, cardiac metabolic perturbations and progressed pathology from 12 weeks of age, prior to alterations in mdx heart. It should be noted that by 24 weeks of age, the mdx heart also deteriorates, and many functional and pathological changes are evident.
Similarly to previous studies in the *mdx* mouse model (29), we observed marked dystrophin restoration in heart of *Cmah-/-;mdx* mice following administration of Pip-PMO targeting removal of exon 23 from the *Dmd* transcript. This restoration led to moderate improvement in some RV cardiac parameters and heart rate, but no improvement in metabolic gene expression levels. As such, PDH flux levels were not reported for the treated samples. These modest changes may be due to several factors; it may be that the treatment was started too late and we missed the therapeutic window, that there is greater intraspecies variation within each cohort in older mice (evidenced by the higher SEM for some *Cmah-/-;mdx* cardiac parameters at 36 weeks), or that the interspecies variation between the *Cmah-/-;mdx* and *mdx* cohorts is diminished (the cardiac *mdx* phenotype deteriorates by 24 weeks exhibiting a similar phenotype to *Cmah-/-;mdx* mice). We also observed marked dystrophin restoration in skeletal muscle, and these levels were sufficient to induce a significant improvement in muscle physiology measures, with approximately 70% correction towards wild-type levels for muscle specific force and complete protection against muscle force drop following eccentric contraction protocol. Interestingly, for the *Cmah-/-;mdx* vehicle group a number of mice did not exhibit as great a force drop following the eccentric protocol as observed previously for *mdx*, which suggests that they may not be as good a model as the *mdx* for assessing treatments targeted to skeletal muscle.

Assessment of metabolic pathways in the *mdx* model have utilized *ex vivo* approaches that demonstrated a clear shift in metabolic substrates towards glycolysis from as early as 12 weeks. In contrast, utilising *in vivo* hyperpolarised MRS, we observed only a small shift towards glycolytic flux in *Cmah-/-;mdx*, but not in *mdx*, even by 24 weeks of age. This suggests that despite a cardiac functional deficit in these mouse models, alterations in metabolic function are less evident and *in vivo* hyper-polarised MRS may not represent an appropriate measure for treatment benefit in these models.

Previous studies in *mdx* mice demonstrate onset of cardiomyopathy from 6 months. Our data presented here suggest that cardiac dysfunction in the *Cmah/mdx* model is not only established earlier (by 12 weeks) but also displays greater severity, and as such may be a more appropriate model to
assess cardiac function. Ultimately, this research shows that the Cmah-/-;mdx mouse is an appropriate model to use if an early dystrophic heart is required for study.

Materials and Methods

All experiments were performed at the University of Oxford, except for the muscle physiology experiments which were conducted at the Royal Veterinary College London. All experiments were performed under the authorisation of the UK Home Office. Mice were housed in IVCs, with access to food and water ad libitum. All animals used were male.

[1-13C]pyruvate hyperpolarized magnetic resonance spectroscopy (MRS) and MRS analysis The hyperpolarized MRS protocol was performed as previously described(26). In brief, mice were anaesthetized using 4% isoflurane for induction and then maintained at 1.5% isoflurane for the duration of the experiment. An intravenous line was established in the tail vein for the administration of hyperpolarized solutions. Mice were weighed and then placed supine into a custom-made cradle and ECG leads were inserted in the forepaws. The cradle was then inserted into the horizontal bore of a 7 T MRI scanner connected to a direct-drive console (Varian Inc). After preliminary axial scans to localise the heart, 0.15 ml of [1-13C]pyruvate, hyperpolarized as described24, was injected over 10 seconds and cardiac gated 13C MR spectra acquired every second for 60 seconds (10 mm radius 13C RF coil). MRS spectra were analysed using the AMARES algorithm in jMRUI software25, employing the maximum metabolite signal normalized to the maximum pyruvate signal to correct for variation in sample polarization and injection rate.

Cine MRI protocol and data analysis Mice underwent Cine MRI as previously described(22). In brief, mice were anaesthetized using 4% isoflurane for induction and then maintained at 1.5%
isoflurane for the duration of the experiment. Mice were weighed and placed supine in a custom-made cradle. ECG leads were inserted into the forearms and a respiratory lead taped across the chest of the mouse. The cradle was inserted into the vertical-bore of an 11.7 T MR system (Magnex Scientific) containing a 40mm birdcage RF coil (Rapid Biomedical). Images were taken using a Bruker console running Paravision 2.1.1 (Bruker Medical). Preliminary axial scans ensured correct positioning of the animal within the coil. The left and right ventricles were imaged by taking a contiguous stack of cine images in 1mm increments and images were analysed using ImageJ software (NIH Image, Bethesda). The epicardial and endocardial borders at end-diastole and end-systole were outlined using the ImageJ free-hand tool. These measurements were used to calculate multiple cardiac structure and function parameters.

Mice were sacrificed by CO₂ inhalation followed by cervical dislocation. Blood, heart, TA and diaphragm tissue samples were collected and snap frozen and stored at -80°C.

**RNA extraction, cDNA synthesis and qPCR** Total RNA was extracted using TRIzol reagent (Invitrogen) as described in the manufacturer’s instructions. 1 ug of RNA was reverse transcribed using a High Capacity cDNA Synthesis kit (ThermoFisher). Diluted cDNA was run using gene specific primers sets (IDT; see Supp Table 3 for Assay IDs) and TaqMan probe set (Integrated DNA Technologies, Leuven, Belgium) on the StepOne Plus Real-Time PCR system (Applied Biosystems). Samples were quantified using the Pfaffl method by attaining the CT values for each reaction and then normalising relative to the house-keeping gene, Ywhaz.

Levels of *Dmd* exon 23 skipping were determined by multiplex qPCR of FAM-labelled primers spanning Exon 20–21 (Assay Mm.PT.47.9564450, Integrated DNA Technologies, Leuven, Belgium) and HEX-labelled primers spanning Exon 23–24 (Mm.PT.47.7668824, Integrated DNA Technologies, Leuven, Belgium). The percentage of *Dmd* transcripts containing exon 23 was determined by normalizing exon 23–24 amplification levels to exon 20–21 levels.
Plasma and liver biochemistry Blood was collected from the jugular vein of mice immediately after sacrifice in plasma tubes (Sarstedt), and sent to the MRC Harwell Mary Lyon Centre (Oxfordshire, UK) for biochemical analysis.

Masson trichrome staining 8 μm transverse sections of heart were cut and mounted on slides. Slides were sent to the Dunn School of Pathology, University of Oxford, for staining. Images were captured with a DM IRB Leica upright microscope (Zeiss monochrome camera) and AxioVision Rel. 4.8 software.

P-PMO synthesis, preparation and administration Pip6a was synthesized by standard solid phase Fmoc chemistry and purified by HPLC, as previously described(22, 23). The PMO sequence (5'-GGCCAAACCTCGGCTTACCTGAAAT-3') was purchased from Gene Tools LLC. Pip6a was conjugated to PMO through an amide linkage at the 3’ end of the PMO, followed by purification by HPLC. The final product was analysed by MALDI-TOF MS and HPLC.

For chronic PPMO treatment, 12-week-old male Cmah-/-mdx mice were administered 10 intravenous tail-vein injections of Pip6a-PMO (12.5 mg/kg) at 2-week intervals. Littermate mice were used as untreated controls.

Muscle physiology Two weeks after the last Pip6a-PMO injection, muscle function was assessed using the right TA muscle. Mice were surgically prepared and analysed as previously described(29). Briefly, optimal muscle length (Lo) was determined by increasing muscle length until the maximal twitch force was achieved. For force–frequency relationship, TA muscles were stimulated at different frequencies, delivered 1min apart (1, 10, 30, 40, 50, 80, 100, 120, 150 and 180 Hz). Muscle fibre physiological cross-sectional area (CSA in cm²) was determined as previously described (29)and
specific isometric force (N/cm²) was calculated by dividing the absolute force (N) at each stimulation frequency by TA muscle physiological cross-sectional area.

Following the force/frequency testing the muscle was tested with an eccentric contraction protocol. The TA muscle was stimulated at 120 Hz for 500 ms before lengthening the muscle by 10% of the Lo at a velocity of 0.5 Lo s⁻¹ for a further 200 ms. Between each contraction a 2-min rest period was permitted to avoid muscle fatigue. A total of 10 eccentric contractions were performed on each mouse. After each eccentric contraction, the maximum isometric force was measured and expressed as a percentage of the initial maximum isometric force achieved at the start of the protocol, prior to the first eccentric contraction. Statistical analysis for the force–frequency and eccentric contraction studies was measured by a repeated measure two-way ANOVA followed by a Tukey’s post-hoc comparison. Statistical significance was defined as a value of \( P < 0.05 \).

**Dystrophin protein extraction and western blot analysis**

For dystrophin protein quantification, transverse cryosections (8 µm thick) were lysed in buffer containing 75 mmol/l Tris–HCl (pH 6.5), 10% sodium dodecyl sulphate, 5% 2-mercaptoethanol, centrifuged at 13 000 rpm for 10 min and supernatant was collected and heated at 100°C for 3 min. Proteins were resolved on a 3–8% Tris–Acetate gel (Invitrogen) and transferred to PVDF membranes for 100 minutes at 30V (Millipore). Subsequently, membranes were probed with monoclonal anti-dystrophin (1:200, NCL-DYS1, Novocastra) and anti-vinculin (loading control, 1:100 000, hVIN-1, Sigma) antibodies and detected with secondary antibody IRDye 800CW goat anti-mouse (LiCOR). Fluorescence was recorded using the Odyssey imaging system. To quantify dystrophin expression, the ratio between dystrophin and vinculin signals was plotted and referred to the ratio of C57BL10 protein standard dilutions on each gel, set as 100%.
Dystrophin Immunohistochemistry analysis

8 μm transverse sections of quadriceps, diaphragm and heart tissues were cut and mounted on slides. Unfixed sections were blocked for 2 hours in 20% foetal bovine serum 20% goat serum solution and double stained for 2 hours with antibodies for dystrophin (Abcam, ab15277) and laminin α2 (Sigma, L0663). Primary antibodies were detected with Alexa 594 and 488 respectively and mounted in Dako Fluorescent Mounting Medium. Images were captured with a DM IRB Leica upright microscope (Zeiss monochrome camera) and AxioVision Rel. 4.8 software.

Statistical analysis All reported values are mean ± standard error of the mean (SEM). When 2 groups were compared, a Student’s t-test was utilised. When multiple groups were analysed, Two-way ANOVA and One-way ANOVA with Tukey post-hoc method was employed. Graphs were drawn using GraphPad Prism 5 software (GraphPad Software Inc, USA). Grubbs test was performed to exclude outliers.

Acknowledgements

We would like to thank the Laboratory of Dr PT Martin from the Research Institute at Nationwide Children's Hospital (Columbus) for providing the Cmah-/-;mdx model. We would also like to thank the Dun School of Pathology for their help with the histology. CAB and TvW were funded by the British Heart Foundation (PG/14/2/30595) and Muscular Dystrophy UK (AVRZFL00), and GM and RM by Association Française contre les Myopathies (AFM). RH, VB and DT were funded by British Heart Foundation (Fellowships FS/10/002/28078 & FS/14/17/30634, Programme Grant RG/11/9/28921), the OXFORD-BHF Centre for Research Excellence and the NIHR Oxford Biomedical Research Centre. SMH was funded by a MRC DPFS Grant.
Conflicts of Interest

No conflicts of interest.
References

1 Koenig, M., Beggs, A.H., Moyer, M., Scherpf, S., Heindrich, K., Betteken, T., Meng, G., Muller, C.R., Lindlof, M., Kaariainen, H. et al. (1989) The molecular basis for Duchenne versus Becker muscular dystrophy: correlation of severity with type of deletion. Am. J. Hum. Genet., 45, 498-506.

2 Vry, J., Gramsch, K., Rodger, S., Thompson, R., Steffensen, B.F., Rahbek, J., Doerken, S., Tassoni, A., Beytia, M.L., Guergueltcheva, V. et al. (2016) European Cross-Sectional Survey of Current Care Practices for Duchenne Muscular Dystrophy Reveals Regional and Age-Dependent Differences. J. Neuromuscul. Dis., 3, 517-527.

3 Matthews, E., Brassington, R., Kuntzer, T., Jichi, F. and Manzur, A.Y. (2016) Corticosteroids for the treatment of Duchenne muscular dystrophy. Cochrane Database Syst. Rev., in press., CD003725.

4 Eagle, M., Baudouin, S.V., Chandler, C., Giddings, D.R., Bullock, R. and Bushby, K. (2002) Survival in Duchenne muscular dystrophy: improvements in life expectancy since 1967 and the impact of home nocturnal ventilation. Neuromuscul. Disord., 12, 926-929.

5 Nigro, G., Comi, L.I., Politano, L. and Bain, R.J. (1990) The incidence and evolution of cardiomyopathy in Duchenne muscular dystrophy. Int. J. Cardiol., 26, 271-277.

6 Bushby, K., Muntoni, F., Urtizberea, A., Hughes, R. and Griggs, R. (2004) Report on the 124th ENMC International Workshop. Treatment of Duchenne muscular dystrophy; defining the gold standards of management in the use of corticosteroids. 2-4 April 2004, Naarden, The Netherlands. Neuromuscul. Disord., 14, 526-534.

7 Sultan, A. and Fayaz, M. (2008) Prevalence of cardiomyopathy in Duchenne and Becker’s muscular dystrophy. J. Ayub. Med. Coll. Abbottabad., 20, 7-13.

8 Kirchmann, C., Kececioglu, D., Kornthember, R. and Dittrich, S. (2005) Echocardiographic and electrocardiographic findings of cardiomyopathy in Duchenne and Becker-Kiener muscular dystrophies. Pediatric cardiology, 26, 66-72.
9 Connuck, D.M., Sleeper, L.A., Colan, S.D., Cox, G.F., Towbin, J.A., Lowe, A.M., Wilkinson, J.D., Orav, E.J., Cuniberti, L., Salbert, B.A. et al. (2008) Characteristics and outcomes of cardiomyopathy in children with Duchenne or Becker muscular dystrophy: a comparative study from the Pediatric Cardiomyopathy Registry. *Am. Heart J.*, **155**, 998-1005.

10 D’Amario, D., Amodeo, A., Adorisio, R., Tiziano, F.D., Leone, A.M., Perri, G., Bruno, P., Massetti, M., Ferlini, A., Pane, M. *et al.* (2017) A current approach to heart failure in Duchenne muscular dystrophy. *Heart*, in press.

11 Kerr, M., Dodd, M.S. and Heather, L.C. (2017) The ‘Goldilocks zone’ of fatty acid metabolism; to ensure that the relationship with cardiac function is just right. *Clin. Sci. (Lond.)*, **131**, 2079-2094.

12 Saini-Chohan, H.K., Mitchell, R.W., Vaz, F.M., Zelinski, T. and Hatch, G.M. (2012) Delineating the role of alterations in lipid metabolism to the pathogenesis of inherited skeletal and cardiac muscle disorders: Thematic Review Series: Genetics of Human Lipid Diseases. *J. Lipid Res.*, **53**, 4-27.

13 Momose, M., Iguchi, N., Imamura, K., Usui, H., Ueda, T., Miyamoto, K. and Inaba, S. (2001) Depressed myocardial fatty acid metabolism in patients with muscular dystrophy. *Neuromuscul. Disord.*, **11**, 464-469.

14 Sicinski, P., Geng, Y., Ryder-Cook, A.S., Barnard, E.A., Darlison, M.G. and Barnard, P.J. (1989) The molecular basis of muscular dystrophy in the mdx mouse: a point mutation. *Science*, **244**, 1578-1580.

15 Stuckey, D.J., Carr, C.A., Camelliti, P., Tyler, D.J., Davies, K.E. and Clarke, K. (2012) In vivo MRI Characterization of Progressive Cardiac Dysfunction in the mdx Mouse Model of Muscular Dystrophy. *PLoS One*, **7**, e28569.

16 Meyers, T.A. and Townsend, D. (2015) Early right ventricular fibrosis and reduction in biventricular cardiac reserve in the dystrophin-deficient mdx heart. *American journal of physiology. Heart and circulatory physiology*, **308**, H303-315.
17 Holland, A., Dowling, P., Zweyer, M., Swandulla, D., Henry, M., Clynés, M. and Ohlendieck, K. (2013) Proteomic profiling of cardiomyopathic tissue from the aged mdx model of Duchenne muscular dystrophy reveals a drastic decrease in laminin, nidogen and annexin. *Proteomics*, in press.

18 Gulston, M.K., Rubtsov, D.V., Atherton, H.J., Clarke, K., Davies, K.E., Lilley, K.S. and Griffin, J.L. (2008) A combined metabolomic and proteomic investigation of the effects of a failure to express dystrophin in the mouse heart. *J. Proteome Res.*, 7, 2069-2077.

19 Khairallah, M., Khairallah, R., Young, M.E., Dyck, J.R., Petrof, B.J. and Des Rosiers, C. (2007) Metabolic and signaling alterations in dystrophin-deficient hearts precede overt cardiomyopathy. *J. Mol. Cell. Cardiol.*, 43, 119-129.

20 Stuckey, D.J., Carr, C.A., Tyler, D.J. and Clarke, K. (2008) Cine-MRI versus two-dimensional echocardiography to measure in vivo left ventricular function in rat heart. *NMR Biomed.*, 21, 765-772.

21 Chandrasekharan, K., Yoon, J.H., Xu, Y., deVries, S., Camboni, M., Janssen, P.M., Varki, A. and Martin, P.T. (2010) A human-specific deletion in mouse Cmah increases disease severity in the mdx model of Duchenne muscular dystrophy. *Sci. Transl. Med.*, 2, 42ra54.

22 Betts, C.A., Saleh, A.F., Carr, C.A., Hammond, S.M., Coenen-Stass, A.M., Godfrey, C., McClory, G., Varela, M.A., Roberts, T.C., Clarke, K. et al. (2015) Prevention of exercised induced cardiomyopathy following Pip-PMO treatment in dystrophic mdx mice. *Sci. Rep.*, 5, 8986.

23 Betts, C., Saleh, A.F., Arzumanov, A.A., Hammond, S.M., Godfrey, C., Coursindel, T., Gait, M.J. and Wood, M.J. (2012) Pip6-PMO, A New Generation of Peptide-oligonucleotide Conjugates With Improved Cardiac Exon Skipping Activity for DMD Treatment. *Mol. Ther. Nucleic Acids*, 1, e38.

24 Potter, L.R., Yoder, A.R., Flora, D.R., Antos, L.K. and Dickey, D.M. (2009) Natriuretic peptides: their structures, receptors, physiologic functions and therapeutic applications. *Handb. Exp. Pharmacol.*, in press., 341-366.

25 Zhang, M., Perino, A., Ghigo, A., Hirsch, E. and Shah, A.M. (2013) NADPH oxidases in heart failure: poachers or gamekeepers? *Antioxid. Redox Signal*, 18, 1024-1041.
26 Dodd, M.S., Ball, V., Bray, R., Ashrafian, H., Watkins, H., Clarke, K. and Tyler, D.J. (2013) In vivo mouse cardiac hyperpolarized magnetic resonance spectroscopy. *J. Cardiovasc. Magn. Reson.*, 15, 19.

27 Sun, W., Liu, Q., Leng, J., Zheng, Y. and Li, J. (2015) The role of Pyruvate Dehydrogenase Complex in cardiovascular diseases. *Life Sci.*, 121, 97-103.

28 Lopez-Bernardo, E., Anedda, A., Sanchez-Perez, P., Acosta-Iborra, B. and Cadenas, S. (2015) 4-Hydroxynonenal induces Nrf2-mediated UCP3 upregulation in mouse cardiomyocytes. *Free Radic Biol. Med.*, 88, 427-438.

29 Godfrey, C., Muses, S., McClorey, G., Wells, K.E., Coursindel, T., Terry, R.L., Betts, C., Hammond, S., O'Donovan, L., Hildyard, J. *et al.* (2015) How much dystrophin is enough: the physiological consequences of different levels of dystrophin in the mdx mouse. *Human molecular genetics*, 24, 4225-4237.

30 LoMauro, A., D'Angelo, M.G. and Aliverti, A. (2015) Assessment and management of respiratory function in patients with Duchenne muscular dystrophy: current and emerging options. *Therapeutics and clinical risk management*, 11, 1475-1488.

31 Yotsukura, M., Miyagawa, M., Tsuya, T., Ishihara, T. and Ishikawa, K. (1988) Pulmonary hypertension in progressive muscular dystrophy of the Duchenne type. *Jpn. Circ. J.*, 52, 321-326.

32 Mehmood, M., Ambach, S.A., Taylor, M.D., Jefferies, J.L., Raman, S.V., Taylor, R.J., Sawani, H., Mathew, J., Mazur, W., Hor, K.N. *et al.* (2016) Relationship of Right Ventricular Size and Function with Respiratory Status in Duchenne Muscular Dystrophy. *Pediatric cardiology*, in press.
Figures

Figure 1. Cardiac function and pathology of mdx and Cmah-/--;mdx hearts compared to C57BL10 at 12 and 24 weeks of age. a) Representative cine-MRI images showing left and right ventricles for heart during diastole and systole. b) Cardiac function parameters, right ventricle ejection fraction (RV EF; %), right ventricle stroke volume (RV SV), left ventricle end diastolic volume (LV EDV), and left ventricle stroke volume (LV SV). b) Quantitative real time (qRT)-PCR for the expression of fibrotic and injury markers namely Ctgf, Nppa and Nox4 in heart tissue normalised to 12 week C57BL10. Data displayed as mean ± SEM. For cardiac function parameters N = 5 to 7, and qRT-PCR data n = 3 to 6. Significance calculated using two-way ANOVA, Tukey post-hoc test (***=P<0.001, **=P<0.01*=P<0.05).
Figure 2. Metabolic profile of C57BL10, *mdx* and *Cmah-/-;mdx* hearts at 12 and 24 weeks of age- hyperpolarised MRS and gene expression analysis. a) Bicarbonate and lactate production normalized to maximum pyruvate signal. Data displayed as mean ± SEM. N = 5 to 7. b) Quantitative real time (qRT)-PCR for the expression of *Pdk1*, *Pdk4*, *Ucp3*, *Cpt1*, *CD36* and *Ppara* in heart tissue normalised to 12 week C57BL10. Data displayed as mean ± SEM. N = 3 to 6. Statistical significance was determined using two-way ANOVA, Tukey post-hoc test (****P<0.0001, ***P<0.001, **P<0.01*=P<0.05).
Figure 3. Biochemical data from plasma samples of C57BL10, mdx and Cmah-/-;mdx mice at 12 and 24 weeks. Graphs showing total cholesterol, free fatty acid, high-density lipoproteins (HDL) and low-density lipoproteins (LDL) measurements. Data displayed as mean ± SEM. N = 4 to 6. Statistical significance was determined using two-way ANOVA, Tukey post-hoc test (**P<0.001, ***P<0.001, **P<0.01, *P<0.05).
Figure 4. PPMO treatment restores dystrophin and improves right ventricle function and pathology in *Cmah-/-;mdx* hearts at 8 months of age. a) Quantification of exon 23-skipped transcripts (RT-qPCR) and dystrophin protein restoration in tibialis anterior (TA), diaphragm and heart following Pip6A-PMO treatment (n=4). b) Cardiac function parameters were measured by cine-MRI. Graphs indicating right ventricle end systolic volume (RV ESV), right ventricle ejection fraction (RV EF), and heart rate (HR) shown. c) RT-qPCR for the expression of injury and metabolic markers namely *Nox4, Nppa, Ucp3, Pdk1 and Pdk4* in heart tissue normalised to C57BL10. Data displayed as mean ± SEM. For cardiac function parameters significance calculated using one-way ANOVA, Tukey post-hoc test. For RT-qPCR data significance was determined using Student’s *t*-test (***=P<0.001, **=P<0.01*=P<0.05).
Figure 5. PPMO treatment improves muscle function in untreated and treated $Cmah^{-/-};mdx$ mice. a) Specific force – frequency relationship for untreated $Cmah^{-/-};mdx$ (n=6), chronically Pip6a-PMO treated $Cmah^{-/-};mdx$ (n=8), and C57BL10 mice (n=12). Treatment with Pip6a-PMO significantly increased specific force in the $Cmah^{-/-};mdx$ mice (p<0.001). b) Force drop associated with eccentric exercise in untreated $Cmah^{-/-};mdx$ and chronically Pip6a-PMO treated $Cmah^{-/-};mdx$. C57BL10 mice not included as their plot overlays the result for the chronically Pip6a-PMO treated $Cmah^{-/-};mdx$ mice. The $Cmah^{-/-};mdx$ mice show a significant force drop with eccentric exercise which is prevented by Pip6a-PMO treatment (p<0.001). Pip6a-PMO treated $Cmah^{-/-};mdx$ and C57BL10 mice are not significantly different.
Table 1. Cine MRI measurements from C57BL10, *mdx* and *Cmah-/-;mdx* mice at 12 and 24 weeks of age. Data displayed as mean ± SEM. N = 5 to 7. Significance calculated using two-way ANOVA, Tukey post-hoc test (****P<0.0001, ***P<0.001, **P<0.01, *P<0.05). BW, body weight; HR, heart rate; LV, left ventricle; RV, right ventricle; EF, ejection fraction; ESV, end-systolic volume; EDV, end-diastolic volume; SV, stroke volume; CO, cardiac output.

| Parameter      | 12 Weeks C57BL10 | 12 Weeks *mdx* | 12 Weeks *Cmah-/-;mdx* | 24 Weeks C57BL10 | 24 Weeks *mdx* | 24 Weeks *Cmah-/-;mdx* |
|----------------|------------------|----------------|------------------------|------------------|----------------|------------------------|
| BW (g)         | 27.1 ± 0.5       | 28.3 ± 0.4     | 27.3 ± 0.5             | 32.3 ± 0.5       | 34.1 ± 0.5*     | 36.2 ± 0.5****†        |
| HR (bpm)       | 482 ± 10         | 452 ± 10       | 466 ± 8                | 453 ± 20         | 445 ± 10        | 505 ± 10**†             |
| Average LV mass/BW | 3.95 ± 0.07     | 3.62 ± 0.09    | 3.57 ± 0.09*           | 4.14 ± 0.15      | 3.44 ± 0.06**** | 3.5 ± 0.2***           |
| LV EF (%)      | 61 ± 4           | 58 ± 4         | 61 ± 3                 | 60 ± 3           | 59 ± 3          | 55 ± 4                 |
| LV EDV/BW      | 2.18 ± 0.09      | 2.00 ± 0.05    | 1.9 ± 0.1              | 2.1 ± 0.1        | 1.87 ± 0.04     | 1.60 ± 0.08**           |
| LV ESV/BW      | 0.9 ± 0.1        | 0.85 ± 0.08    | 0.76 ± 0.08            | 0.9 ± 0.1        | 0.78 ± 0.06     | 0.72 ± 0.09             |
| LV SV/BW       | 1.31 ± 0.06      | 1.15 ± 0.07    | 1.15 ± 0.07            | 1.23 ± 0.04      | 1.10 ± 0.05     | 0.87 ± 0.04****†        |
| LV CO/BW       | 0.63 ± 0.04      | 0.52 ± 0.04*   | 0.53 ± 0.03*           | 0.56 ± 0.03      | 0.49 ± 0.02     | 0.4 ± 0.2*              |
| RV EF (%)      | 71 ± 3           | 68 ± 4         | 60 ± 2*                | 71 ± 2           | 55 ± 2***       | 59 ± 4*                 |
| RV EDV/BW      | 1.86 ± 0.05      | 1.72 ± 0.04    | 1.75 ± 0.09            | 1.80 ± 0.08      | 1.87 ± 0.05     | 1.61 ± 0.03†            |
| RV ESV/BW      | 0.53 ± 0.05      | 0.55 ± 0.07    | 0.71 ± 0.07            | 0.54 ± 0.05      | 0.84 ± 0.04**   | 0.67 ± 0.08             |
| RV SV/BW       | 1.32 ± 0.07      | 1.17 ± 0.07    | 1.04 ± 0.04**          | 1.27 ± 0.05      | 1.04 ± 0.04*    | 0.94 ± 0.05**           |
| RV CO/BW       | 0.64 ± 0.05      | 0.53 ± 0.04    | 0.48 ± 0.02**          | 0.57 ± 0.03      | 0.46 ± 0.02*    | 0.48 ± 0.03             |
Table 2. Cine MRI measurements from C57BL10, mdx, Cmah-/mdx and Cmah-/-mdx treated mice at 8 months of age. Data displayed as mean ± SEM. N = 6 to 9. Significance calculated using one-way ANOVA, Tukey post-hoc test (***P<0.001, **P<0.01*P<0.05). BW, body weight; HR, heart rate; LV, left ventricle; RV, EF, ejection fraction; right ventricle; ESV, end-systolic volume; EDV, end-diastolic volume; SV, stroke volume; CO, cardiac output.

| Parameter                  | C57BL10  | mdx     | Cmah-/mdx | Cmah-/-mdx Treated |
|----------------------------|----------|---------|-----------|--------------------|
| BW (g)                     | 35.00    | 38.00   | 38.33     | 38.25              |
| HR (bpm)                   | 442.62 ± 13.7 | 442.41 ± 15.2 | 523.1 ± 13.4 **// | 454.27 ± 15.8 ++   |
| Average LV mass/BW         | 3.48 ± 0.15 | 2.93 ± 0.09 # | 3.61 ± 0.21 / | 3.51 ± 0.09 /     |
| LV EF (%)                  | 63 ± 2.8 | 60 ± 3.8 | 54 ± 4.4 | 55 ± 4.3           |
| LV EDV/BW                  | 1.73 ± 0.11 | 1.37 ± 0.08 | 1.69 ± 0.18 | 1.69 ± 0.05       |
| LV ESV/BW                  | 0.65 ± 0.07 | 0.56 ± 0.06 | 0.83 ± 0.19 | 0.78 ± 0.09       |
| LV SV/BW                   | 1.09 ± 0.07 | 0.82 ± 0.06 * | 0.86 ± 0.06 | 0.91 ± 0.05       |
| LV CO/BW                   | 0.48 ± 0.04 | 0.36 ± * | 0.45 ± 0.03 | 0.41 ± 0.02       |
| RV EF (%)                  | 72 ± 1.2 | 57 ± 3 ** | 52 ± 2.8 *** | 60 ± 2.3 *         |
| RV EDV/BW                  | 1.51 ± 0.09 | 1.41 ± 0.07 | 1.67 ± 0.01 | 1.47 ± 0.03       |
| RV ESV/BW                  | 0.43 ± 0.03 | 0.60 ± 0.04 | 0.81 ± 0.08 *** | 0.58 ± 0.03 +     |
| RV SV/BW                   | 1.08 ± 0.07 | 0.81 ± 0.06 * | 0.87 ± 0.06 | 0.89 ± 0.04       |
| RV CO/BW                   | 0.48 | 0.36 * | 0.45 / | 0.40 |
**Abbreviations**

ALT- alanine transaminase

AO- antisense oligonucleotide

AST- aspartate transaminase

BW- body weight

CK- creatine kinase

Cmah- cytidine monophosphate-sialic acid hydroxylase

CO- cardiac output

ECG- electrocardiogram

EDV- end-diastolic volume

EF- ejection fraction

ESV- end-systolic volume

FFA- free fatty acid

HDL- high-density lipoproteins

HR- heart rate

LDH- lactate dehydrogenase

LDL- low-density lipoproteins

LV- left ventricle
MRIs- magnetic resonance imaging

MRS- magnetic resonance spectroscopy

PDH- pyruvate dehydrogenase

RV- right ventricle

SV- stroke volume

TA- tibialis anterior