nature portfolio

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Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our Editorial Policies and the Editorial Policy Checklist.

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FOI 6	an statistical analyses, commit that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed
\boxtimes	\square The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
\boxtimes	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.
\boxtimes	A description of all covariates tested
\boxtimes	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
\boxtimes	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
\boxtimes	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted Give <i>P</i> values as exact values whenever suitable.
\boxtimes	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
\boxtimes	\square Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated
'	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection

qPCR data collected with QuantStudio 6 and 7 Flex Real-Time PCR System Software. Western Blot data collected with Bio-Rad Chemi-Doc & Image-Lab Software. LRS data collected using Oxford Nanopore MinKNOW software.

Data analysis

RNA-seq data were analyzed following the GATK Best Practices recommendations. Blots were analyzed using the Image-Lab Software. Tools used for data analysis, such as FLAIR for LRS data, are mentioned in the method section of the paper. Custom codes are a negligible part of

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.

Data

Policy information about availability of data

All manuscripts must include a <u>data availability statement</u>. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our policy

The sequencing data of this study were stored in NCBI Bio Project (PRJNA809363).

Field-spe	cific re	porting
Please select the or	ne below that is	the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
∑ Life sciences	В	ehavioural & social sciences
For a reference copy of t	he document with a	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>
Life scien	ices stu	ıdy design
All studies must dis	close on these	points even when the disclosure is negative.
Sample size	Single patient st	cudy with matching healthy control.
Data exclusions	N/A	
Replication	qPCR & WB rep	licated. LRS not replicated due to great amount of data generated and high cost of replication.
Randomization	N/A	
Blinding	N/A	
We require informatic system or method list Materials & exp n/a Involved in th Antibodies Eukaryotic Palaeontold Animals and Human reset Clinical data Dual use re	on from authors a ed is relevant to perimental so e study cell lines ogy and archaeol d other organism earch participant	n/a Involved in the study ChIP-seq Flow cytometry MRI-based neuroimaging ss
Antibodies		
Antibodies used	MLIP (Invitrogen, PA5-72759); LMNA (NEB, 2026S); GAPDH (NEB, 5174S); Anti-Rabbit (Jackson ImmunoResearch, 111-035-114)	
Validation	All primary antibodies are validated for use against human proteins. All antibodies, except for MLIP, are validated for use with Western blots. MLIP antibody is validated for immunohistochemistry.	
Human resea	arch parti	cipants
Policy information about studies involving human research participants		
Population characteristics Single patient with late		Single patient with late-onset myopathy without a genetic cause.
Recruitment		Recruited for RNA-seq following failed attempt to identify genetic cause with gene panels.

McGill University Health Centre Research Ethics Board

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Ethics oversight