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 <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

		CS

FOI	all St	atistical analyses, commit that the following items are present in the figure regend, table regend, main text, or Methods section.
n/a	Cor	nfirmed
	\boxtimes	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
\boxtimes		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.
X		A description of all covariates tested
X		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. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted Give P 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	Estimates of effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated
		Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about <u>availability of computer code</u>

Data collection

Provide a description of all commercial, open source and custom code used to collect the data in this study, specifying the version used OR state that no software was used.

Data analysis

Provide a description of all commercial, open source and custom code used to analyse the data in this study, specifying the version used OR state that no software was used.

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 generated datasets are available in the European Nucleotide Archive under the accession number PRJEB49648

Research inv	volving hu	man participants, their data, or biological material				
,		vith human participants or human data. See also policy information about sex, gender (identity/presentation), thnicity and racism.				
Reporting on sex and gender		The genetic sex was determined for the human samples.				
Reporting on race, ethnicity, or other socially relevant groupings		The human samples came from an archaeological context and were analyzed with contemporaneous samples.				
Population characteristics		N/A				
Recruitment		The samples that could be sampled were analyzed.				
Ethics oversight		N/A				
Note that full informa	ation on the appro	oval of the study protocol must also be provided in the manuscript.				
Field-spe	ecific re	porting				
Please select the o	ne below that is	s 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				
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Life scier	nces stu	udy design				
All studies must dis	sclose on these	points even when the disclosure is negative.				
Sample size	2					
Data exclusions	N/A					
Replication	N/A					
Randomization	ndomization N/A					
Blinding	N/A					
Reporting for specific materials, systems and methods We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material,						
,		your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.				
Materials & ex	perimental s	vstems Methods				
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Antibodies		ChIP-seq				
Eukaryotic cell lines		Flow cytometry				
Palaeontology and archaeology Animals and other organisms		———				
Clinical dat						
Dual use research of concern						
Plants						

Palaeontology and Archaeology

Specimen provenance Warburg, Germany

Specimen deposition The human skeletons from Warburg are kept at the LWL in Münster.

Dating methods The samples were directly dated using the radiocarbon method.

💢 Tick this box to confirm that the raw and calibrated dates are available in the paper or in Supplementary Information.

Ethics oversight The State Monuments Office has approved the sampling and analysis.

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

Plants

Seed stocks

Report on the source of all seed stocks or other plant material used. If applicable, state the seed stock centre and catalogue number. If plant specimens were collected from the field, describe the collection location, date and sampling procedures.

Novel plant genotypes

Describe the methods by which all novel plant genotypes were produced. This includes those generated by transgenic approaches, gene editing, chemical/radiation-based mutagenesis and hybridization. For transgenic lines, describe the transformation method, the number of independent lines analyzed and the generation upon which experiments were performed. For gene-edited lines, describe the editor used, the endogenous sequence targeted for editing, the targeting guide RNA sequence (if applicable) and how the editor was applied.

Authentication

Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosiacism, off-target gene editing) were examined.