

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](#).

Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

n/a	Confirmed
☐	☒ The exact sample size (<i>n</i>) 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
☐	☒ The statistical test(s) used AND whether they are one- or two-sided <i>Only common tests should be described solely by name; describe more complex techniques in the Methods section.</i>
☒	☐ A description of all covariates tested
☒	☐ A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
☐	☒ 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)
☐	☒ 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 <i>Give P values as exact values whenever suitable.</i>
☐	☒ For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
☒	☐ For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
☐	☒ Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i>), 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 [availability of computer code](#)

Data collection	Sequence data were processed using the following programs to obtain genotype data used in the analysis: AdapterRemoval v2.2.2, BWA v0.7.15, Samtools v1.9, Picard markduplicate v2.21.8, pileupCaller (https://github.com/stschiff/sequenceTools), mapDamage v2.1.0, ANGSD v0.920, Schmutzi v1.5.6. These programs are publicly available.
Data analysis	Population genetic data analysis in this study was performed using the following publicly available programs: HaploGrep v2.4, Y_leaf, smartpca v13050, ADMIXTURE v1.3.0, PLINK v1.90, qp3Pop v412, qpDstat v712, KIN(https://github.com/DivyaranPopli/Kinship_Inference), anclBD(https://github.com/hringbauer/anclBD), hapROH(https://haproh.readthedocs.io/en/latest/intro.html), GLIMPSE2. Non-default parameters used in our analysis are described in the Methods section. The base map in Figure 1a was generated by ArcGIS pro v3.0. Calibration of AMS 14C dating results was done by either IntCal20 or OxCal v4.4.4.

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 [data availability statement](#). 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](#)

Raw sequence data (BAM format) are available at the Genome Sequence Archive in the National Genomics Data Center (GSA) under accession HRA007922. The mitochondrial DNA sequences have been deposited in GenBase in National Genomic Data Center, Beijing Institute of Genomics, Chinese Academy of Sciences/China National Center for Bioinformation, under accession number C_AA111067.1 to C_AA111103.1 that is publicly accessible at <https://ngdc.cncb.ac.cn/genbase>.

Research involving human participants, their data, or biological material

Policy information about studies with [human participants or human data](#). See also policy information about [sex, gender \(identity/presentation\), and sexual orientation](#) and [race, ethnicity and racism](#).

Reporting on sex and gender	The genetic sex of our ancient samples was determined by analyzing the ratio of X and Y chromosome coverages relative to autosomes.
Reporting on race, ethnicity, or other socially relevant groupings	The concept of race or ethnicity was not applicable to our study, as we focused on ancient DNA samples where such social constructs do not apply. Instead, we classified our samples based on archaeological context and cultural associations with the Dawenkou culture. The categorization was done based on the site location, burial practices, and associated artifacts typical of the Dawenkou culture, as identified through archaeological methods.
Population characteristics	Our study examined individuals from the Baligang archaeological site. The samples are dated between 6,100 and 2,500 cal BP, based on radiocarbon dating of the burial remains. The demographic characteristics such as age at death were inferred from osteological analysis. Detailed genetic information was obtained through sequencing ancient DNA, revealing insights into the genetic structure and diversity of this ancient population.
Recruitment	Participants were not recruited in the traditional sense, as this study involved ancient human remains. The samples were obtained from well-documented archaeological excavations conducted by the school of archaeology and museology at Peking University. We ensured that all samples were ethically sourced and that the excavation and analysis followed proper archaeological and bioethical guidelines to minimize contamination and ensure respectful handling of human remains.
Ethics oversight	The study protocol was approved by the ethics committee of the school of archaeology and museology at Peking University. Additionally, all necessary permits for the excavation and analysis of the ancient human remains were obtained from relevant local authorities.

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

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

☐ Life sciences ☐ Behavioural & social sciences ☒ Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see [nature.com/documents/nr-reporting-summary-flat.pdf](https://www.nature.com/documents/nr-reporting-summary-flat.pdf)

Ecological, evolutionary & environmental sciences study design

All studies must disclose on these points even when the disclosure is negative.

Study description	This study includes whole genome sequencing of 58 ancient individuals from Henan province, China, dating between 6,100 and 2,500 cal BP, associated with both North China's Yellow River cultures and South China's Yangtze River cultures.
Research sample	Research samples are composed of 58 ancient genomes from the Baligang archaeological site. The samples are dated between 6,100 and 2,500 cal BP, associated with both North China's Yellow River cultures and South China's Yangtze River cultures.
Sampling strategy	No sample-size selection was performed prior to the study. To produce ancient genomes reported in this study, we screened the accessible skeletal elements from the relevant geographic regions and time periods, and produced in-depth sequencing data for those with sufficient endogenous DNA preservation and without substantial contamination.
Data collection	Sequencing of the libraries was performed on an Illumina Novaseq platform with paired-end 2x150 cycles, HiSeq2500 platform with paired-end 2x100 cycles and CYGNUS S100 platform with 1 x 100 cycles.

Timing and spatial scale	Laboratory works and sequencing was conducted over the period from February 2015 to December 2023. Samples were taken from China, the location of which are provided in Fig 1.
Data exclusions	We excluded samples only if the samples do not meet the quality criteria, either by having low level of endogenous human DNA prohibiting genome-scale sequencing or by showing high level of contamination estimates. For population genetic analysis that requires exclusion of genetic relatives, we excluded closely related individuals (1st degree relatives) by removing one with lower coverage from each pair.
Reproducibility	We took multiple individuals from each archaeological site, if available, to support the representativeness of their genetic profiles. For each sample, we estimated contamination level to support the authenticity of data.
Randomization	Ancient genomes were first analyzed by each individual, and then were allocated into the analysis group based on their archaeological context, absolute date (14C dating), and their individual genetic profile. Randomization is not applicable.
Blinding	There was no experimental treatment of samples involved in this study that requires blinding. Data analysis was performed based on the analysis groups that were defined by archaeological information (archaeological context and date) and genetic homogeneity.

Did the study involve field work? ☐ Yes ☒ No

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, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

n/a	Involved in the study
☒	☐ Antibodies
☒	☐ Eukaryotic cell lines
☐	☒ Palaeontology and archaeology
☒	☐ Animals and other organisms
☒	☐ Clinical data
☒	☐ Dual use research of concern
☒	☐ Plants

Methods

n/a	Involved in the study
☒	☐ ChIP-seq
☒	☐ Flow cytometry
☒	☐ MRI-based neuroimaging

Palaeontology and Archaeology

Specimen provenance	The specimens analyzed in this study were obtained from the Baligang archaeological site, China. Permits for the excavation and export of these samples were obtained from the school of archaeology and museology at Peking University.
Specimen deposition	The specimens analyzed in this study have been deposited at the ancient DNA laboratory at Peking University. These specimens are available for further research upon request, under the guidelines and supervision of the school of archaeology and museology at Peking University, to permit free access by other researchers.
Dating methods	New radiocarbon dates were provided for this study. The samples were pretreated, and measurements were taken at the Radiocarbon Dating Laboratory of Peking University. The calibration program used was OxCal v4.4, and quality assurance protocols were strictly followed as per the laboratory standards. No new dates are provided beyond those obtained during this pretreatment and measurement process.
☒ Tick this box to confirm that the raw and calibrated dates are available in the paper or in Supplementary Information.	
Ethics oversight	The study protocol was approved by the ethics committee of the school of archaeology and museology at Peking University. Additionally, all necessary permits for the excavation and analysis of the ancient human remains were obtained from relevant local authorities.

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

Plants

Seed stocks	N/A
Novel plant genotypes	N/A
Authentication	N/A