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Biological Sciences; Ecology

Climatic niche shifts drove rapid expansion of Paleolithic Modern Humans

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16 **Abstract**

17 The routes by which Palearctic modern humans expanded their geographic ranges across Eurasia to colonize the Americas
18 during the Late Pleistocene have been intensively analyzed. Whether this geographic expansion occurred as a result of tracking
19 a specific set of favourable climatic conditions, or via concurrent colonization of novel climates, remains unclear. Analyses of
20 the ecological niche linking archeological and paleoclimatic data revealed that Palearctic hunter-gatherers significantly altered
21 their climatic niche during the last phase of the Late Pleistocene, colonizing novel climatic as well as geographic space
22 between 46ka-26ka. In contrast, from 26ka-11ka, the climatic niche was more stable, even as humans dispersed to different
23 geographic regions. This dispersal was facilitated by a persistent climatically suitable ‘corridor’ linking Western Europe to Far
24 East Asia beginning 32ka, via a mid-latitude belt in South Siberia. Other areas with suitable climates over long periods
25 included Kamchatka and regions of Beringia which are currently submerged. Niche dynamics were controlled by changes in
26 seasonal water availability and Upper Paleolithic technological innovations associated with transitions between cultural
27 periods.

28
29 *Keywords:* Late Pleistocene; modern humans; climatic niche; climatic refugia

30
31 **Significance Statement**

32 Humans have successfully colonized every continent except Antarctica and live in nearly every ecosystem, from the wet
33 tropics to the high arctic. However, at some point in our history, our survival was dependent upon a much more constrained set
34 of climatic conditions. We explore how changes in the environmental conditions in which early humans were able to persist, as
35 well as changes in cultural periods, facilitated human expansion through Eurasia during the Late Pleistocene. Using a unique
36 framework that integrates paleo-information sources with macroecology we show that humans expanded their ability to survive
37 in more diverse climatic conditions until ~26ka while afterwards tracked specific types of climates, probably due to the adverse
38 conditions around the Last Glacial Maximum.

Introduction

Anatomically Modern Humans (AMHs), hereafter modern humans, are believed to have originated in Africa approximately 200ka (ka= thousand years ago). From there, they moved outward through Eurasia (~60ka) (but see 1), crossing to the Americas through Beringia by at least 15ka, and finally populating South America (2-4). The spread and colonization of modern humans throughout Europe and Asia occurred during a period of intense climate change, and these changing conditions may have driven humans to colonize new regions via specific dispersal routes (5-7).

Whereas the timing and routes of these dispersal events across geographical space have been intensively analysed and heavily disputed (1, 8-10) less is known about the niche dynamics of hunter-gatherers populations during this dispersal. A fundamental question is whether modern humans colonised novel climatic conditions as they expanded their geographic distribution, or tracked a specific set of climates in which they were able to persist. In comparison, the responses of other iconic glacial-era species to the climatic changes of this period are relatively well investigated. Macroecological approaches have successfully reconstructed the Late Pleistocene biogeography of numerous animal and plant species by quantifying species' climatic niches (11), revealing changes in niche size (12, 13), locations of refugia (14, 15) or extinction probability (16-18). These studies suggest that climate change responses were species-specific: some species tracked favorable climatic conditions as they shifted in space, while others remained in the same geographic regions by adapting to new climatic conditions or expanded their distributions into novel regions and climates (16, 17). However, similar studies for modern humans are still in their infancy (but see 19-21).

Here, we present evidence that modern humans shifted and expanded their niche into new climatic conditions, in response to the magnitude of climate change and transitions between cultural periods from 46ka until 11ka. Moreover, we provide evidence for a continuous corridor across South Siberia that maintained relatively stable climatic conditions suitable for dispersal by modern humans between Europe and central Asia. We also demonstrate a unique quantitative framework that integrates a suite of paleo-information sources with macroecological and community ecological tools for identifying the responses of modern humans to past climatic changes.

We quantified changes in the climatic niche, or the set of climatic conditions occupied by modern human from Europe and central and north Asia, between 46ka and 11ka. We used 3,993 Eurasian Paleolithic radiocarbon-dated occurrences and 25 paleoclimatic simulations at intervals of 1,000/ 2,000-years, to calculate changes in three niche parameters: i) niche overlap,

defined as the amount of climate space continuously occupied by modern humans in two consecutive time intervals (22); ii) niche breadth, or the range of climatic conditions inhabited at each interval, and iii) niche marginality, measured as the distance between the average climatic conditions inhabited by modern humans within each interval and the average climatic conditions of the study area across all 25 climatic intervals (23-25) (Fig.1). In addition, we estimated the geographical distribution of modern humans' suitable climatic conditions in each time interval using Climatic Envelope Models (CEMs) (26-28). We used the resulting maps of climatic suitability (as in 29) to quantify changes in the distribution of modern humans and to identify areas with consistently high climatic suitability throughout the study period (14). Finally, we implemented Generalized Additive Models (GAMs) to separate the effects of climate change versus technological/cultural transitions as potential drivers of the observed changes in niche parameters.

We provide answers to three fundamental questions about the historical biogeography of modern human expansion across Eurasia: i) to what extent did this expansion occur via colonization of novel climatic conditions versus dispersal to novel regions following a specific set of climate conditions; ii) how important are changes in climate versus cultural periods in explaining niche dynamics; and iii) what was the most plausible route, and over which time periods, connecting modern humans from Europe and central and northeast Asia?

Results

Climatic niche dynamics

The climatic niche of modern humans changed extensively from 46ka to 11ka, as indicated by varying levels of niche overlap between intervals across this period ($D = 0.271$; with 0 being no overlap and 1 complete overlap; low overlap is 0-0.3, medium overlap is 0.3-0.7 and high overlap is 0.7-1; these categories follow 22, 30). Changes in the climatic niche happened gradually over this 35,000 year time period ($0.3 < D < 0.7$, Fig. 2A), punctuated by brief periods of rapid niche change. The episodes of rapid change were concentrated in the first half of the study period, as indicated by large differences in niche overlap for some consecutive periods occurring before 26ka. Following 26ka, the climatic niche of modern humans showed consistently higher niche overlap.

Analysis of niche breadth indicate that these changes reflect expansion of the climatic niche, with niche breadth consistently increasing from 40ky until reaching its largest extent at 22ka, with some intermittent periods of small contractions. Two time intervals exhibited very rapid growth, as niche breadth expanded by 483% between 40ka and 38ka, and by 83%

between 30ka and 26ka (Fig. 2B). Following the maximum niche breadth at 22ka, the niche contracted gradually until the end of the Pleistocene at 11ka (Fig. 2B).

In contrast to niche breadth, niche marginality, or the distance between the average climatic conditions occupied by modern humans in each time interval and the average conditions from 46ka to 11ka of the study area, roughly declined throughout the study period (Fig. 2B). As modern humans expanded their niche into new climatic conditions, the centroid of their climatic niche approached the mean climatic conditions of the geographical study area across the 25 climatic simulation intervals. Modern humans' climatic niche became progressively less marginal relative to the available climate space (i.e. people occupied a larger fraction of the climatic zones available across Eurasia) until 26ka, succeeded by a period of consistently low marginality until 16ka. After 16ka, the climatic niche of modern humans became increasingly marginal relative to the average climatic conditions 46ka-11ka (Fig. 2B). The warmer and wetter conditions occupied by Palearctic modern humans at the end of the Pleistocene are far from the average conditions for the time extent of our study, which was colder and dryer during most of the time, reflected in our results by a final large increase in niche marginality.

Trends across all niche parameters roughly divide the 35,000 year temporal extent of our study into two main periods: a period of niche change associated with niche expansion from the beginning of the study, 46ka, until ~26ka and a period of larger niche stability associated with gradual niche contraction from ~26ka through the end of the Pleistocene, ~11ka. These periods partially coincide with two distinct paleoclimatic periods: Marine Isotope Stage 3 (MIS3; ~60ka-27ka), which was warmer than MIS2, and Marine Isotope Stage 2 (MIS2; ~27ka-11ka), which was characterized by cold and arid conditions throughout most of its extent, including the Last Glacial Maximum (~21ka: LGM) (31-32). The variances of niche overlap ($P=0.026$) and niche breadth ($P=0.041$) are significantly different between periods, and the variance of niche marginality is nearly so ($P=0.054$).

Cultural and climatic drivers of niche dynamics

Changes in climate, particularly precipitation, were found to be the strongest driver of niche parameters (see *Materials and Methods* and *SI Results*). Change in summer precipitation was the strongest predictor of niche overlap (deviance explained=57.8%; $P=0.001$). Niche breadth change was correlated with change in summer precipitation (deviance explained=57%; $P=0.001$). Niche marginality change was most strongly correlated with change in winter precipitation (deviance explained=52.8%; $P<0.001$) closely followed by change in spring precipitation (deviance explained=49%; $P<0.001$) (Table S3). Deviance explained increased from 5%-40% for all models when changes in cultural periods were added

119 to single-variable climatic models as a categorical factor with 5 levels (see *Materials and Methods* and Table S4). Changes
120 between cultural periods as a single predictor of the three niche parameters were not statistically significant (Table S5).

121 **Geographic overlap of climatically suitable areas**

122 Changes in geographic overlap of climatically suitable areas (hereafter geographic overlap) between consecutive time
123 intervals are consistent with the patterns for niche overlap, breadth and marginality in that the results roughly indicate two
124 periods (46ka-26ka and 26ka-11ka) differing both in direction and magnitude of change (Fig. 2A). There was a general
125 tendency for decreasing geographic overlap until 26ka, followed by a tendency for increase until 11ka. Apart from this general
126 trend, medium to large overlap within discrete intervals during these two periods suggest short intervals of relative stability in
127 the geographic distribution of suitable climatic conditions (Fig. 2A).

128 Comparing the geographic distribution of climatically suitable areas across all intervals (Fig. 3) indicates that a belt of
129 consistently suitable climatic conditions across South Siberia (5) may have allowed modern humans to disperse across western
130 Eurasia and Northeastern Eurasia/Beringia (33). We also found climatically stable areas that are isolated from other patches of
131 suitable conditions, which may have acted as potential climatic refugia (34), occurring in present day East China, Japan, Korea,
132 Kamchatka and submerged areas of Beringia (Fig. 3).

133 **Discussion**

134 We document the extent to which the expansion of modern humans across Eurasia occurred via colonization of novel
135 climatic conditions or by tracking specific climate conditions into new regions, along 35,000 years of climate change. We
136 found that modern humans followed both strategies: from 46ka to 26ka, changes in geographic distribution coincided with
137 expansion of the climatic niche, but after 26ka, they began to track a similar set of suitable conditions during the extreme cold
138 and arid conditions of the Last Glacial Maximum. The combined effect of both climate change and changes between cultural
139 periods are significant predictors of the shifts in the climatic niche of modern humans. In addition, we present evidence of a
140 potential dispersal route across South Siberia which retained suitable climatic conditions dating back to 32ka and persisting
141 until 18 ka, when this belt became more unsuitable, suggesting reduced potential for dispersal across a vast space of harsh
142 climate immediately following the onset of the LGM (5, 33, 35, 36). Our results are robust to the number of occurrences of
143 modern humans and to the temporal resolution of millennial versus bimillennial time periods (see *SI Results*).

During the niche-expansion phase, from 46ka to 26ka and roughly coinciding with MIS3, modern humans expanded their distribution across much of the Palearctic (6); evidence from archeology and human genetics show that modern humans had already reached parts of south-central Siberia -although there is also evidence for occupation of more northerly sites (2, 33, 37, 38). Our results reveal that from 46ka to 26ka, human expansion into new regions of Eurasia was accompanied by increased niche breadth and low niche overlap between intervals, reflecting the growing variety of climatic conditions that modern humans were able to inhabit and exploit.

In contrast, during MIS2, 26ka-11ka, the climatic niche of modern humans was more stable. During this period populations dispersed to and inhabited regions with similar climatic conditions; that is, modern humans entered a climate tracking phase, as suggested by higher niche overlap, lower niche expansion and higher geographical overlap between intervals (Fig. 2) than during the MIS3. These findings indicate that modern humans adjusted their geographical distribution to colonize suitable climatic conditions rather than expanding to fill new ones. There is evidence that human populations in high latitudes persisted in some pockets of suitable climate (33, 39) during MIS2. Coinciding with the LGM, modern humans experienced a decline in niche breadth. We presume that this decline reflects the decrease in the geographical availability of climatic conditions supporting the minimum levels of ecosystem productivity required to maintain viable populations of hunter-gatherers. Changes in seasonal water availability appear to be the key driver of change in climate niche parameters. In temperate and cold areas, the level of precipitation during the growing season could have played a critical role in plant productivity, driving the availability of a vital resource for hunter-gatherer populations, herbivores and food webs on which they may have depended (40-42). Climatic conditions have been used to predict Net Primary Productivity levels (43) for this period (36) with higher productivity associated with warmer and wetter conditions. During the cold and dry conditions of the LGM, the rate of gross terrestrial primary productivity was about $40 \pm 10 \text{ PgCyr}^{-1}$, half of the pre-industrial Holocene (44).

However, our results suggest also that transitions to more modern cultural periods also contributed, as a secondary factor, to the ability of modern humans to colonize new climatic conditions. Hunter-gatherers during the Late Pleistocene demonstrated a remarkable variety of cultural adaptations concurrent with a period of climatic and environmental changes, which may have played a key role in ensuring their survival and population growth. Cultural evolution is indeed suggested to have been affected by major episodes of unfavourable conditions (45, 46), population growth, intra and inter-population interactions (47) and subsistence practices (48). Upper Paleolithic hunting tools show a considerable variance and diversified rapidly both in time and space (48) exhibited by the similarities in 'cultural periods' between far apart Eurasian populations (49-51). This

171 diversification might stem from the different carrying capacities of ecosystems, variability of resources, seasonality and
172 demographic pressure. As a result, modern humans may have increased their dietary niche (52) and respectively the need for
173 more efficient resource uptake. The ‘cultural periods’ used in our study do not reflect specific technological changes per se, but
174 rather represent adaptations that would enable them to survive in a variety of climatic conditions, thus increasing their climatic
175 niche.

176 Existence of a persistent corridor with suitable climate across Southern Siberia suggests that modern human populations
177 inhabiting Europe and central-north Asia may have remained connected via dispersal along this route (Figs. 3 and S7). Our
178 results indicate that this corridor linking Europe and Asia emerged ~36ka, in agreement with recent findings based on ancient
179 DNA of European populations in the Middle Don River (53). Previous studies have also documented evidence of gene flow
180 between Europeans and central Asian populations (5, 53, 54). Despite the early emergence of this relatively continuous belt of
181 suitable climatic conditions surrounded by highly unsuitable areas, it is after 32ka that this route remains highly suitable until
182 18ka. Modern humans have been recorded in south Siberia as early as 45ka (55), and occupations have been detected at
183 relatively low frequencies in this region through 36ka, but with substantially increased frequency after 16ka (39). Recent
184 genomic sequencing of ancient DNA from two individuals from northeastern Siberia, dated 24ka and 17ka, suggests that this
185 region was occupied throughout the Last Glacial Maximum (54). Whether this indicates continuous occupation or the region
186 was mainly depopulated during the LGM is still debated (2, 37, 49). However, the climatic suitability along this corridor was
187 much lower than that of Western Europe, suggesting that the modern human populations within central Siberia may have
188 dispersed, occupied and survived in lower productivity ecosystems and lower population sizes than in Western Europe. All
189 regions above 61° latitude showed consistently low climatic suitability across the time extent of the study, although the models
190 include some archaeological localities from that latitudinal band. These localities have been reported in previous studies (33,
191 37, 49, 50), suggesting, in light of our results, that some pioneering human populations either survived in conditions of
192 extremely unsuitable climates and low productivity ecosystems, or in micro-refugia at a spatial scale that is poorly reflected by
193 the 2-degree spatial resolution of our paleoclimatic simulations. Surprisingly, the area south of this mid-latitude belt in Asia
194 (south of 48 ° latitude, apart from East China) also exhibited low climatic suitability for most time intervals of the analyses
195 (Fig. S7). This is due to a lack of well dated human occurrences for South Asia in our database. To more fully understand the
196 movement patterns of modern humans during the Late Pleistocene, key regions like the Arabian Peninsula, South Asia, East
197 Asia and Southeast Asia need to be more intensively surveyed and studied (56).

While all regions of our study area were affected to some extent by climate change, western and eastern Europe have numerous archaeological sites from the period (33, 57-59) and were probably continuously inhabited by modern humans (Fig. S7). Our results show that the climate of this region was consistently more suitable than that of eastern Eurasia, suggesting that the ecosystems had levels of primary productivity able to support dense modern human populations. Outside Western Europe and the Middle East, however, the presence of suitable climatic conditions in East China, Japan, Korea, Kamchatka and Beringia over lengthy periods suggests that these regions may have served as climatic refugia for modern humans. High latitude climatic refugia have been frequently reported for woolly rhinos, woolly mammoths, horses, reindeer, elk musk ox (16, 60-64) and small mammals (65-67). Furthermore, Beringia has been proposed as a potential refugium for modern humans and other animal and plant species based on records from sea-floor sediments (10, 40, 41) and from the presence of similar ecosystems in analogous latitudes (68-70).

Naturally, there are caveats associated with our analyses of the early biogeography of modern humans. First, our results are contingent on this particular data set, spatial resolution and temporal extent of the analyses. The addition of localities recording human presence across southern Asia, for example, would expand the area of suitable climatic conditions from the South Siberian belt towards these regions. In addition, the archaeological record may not accurately reflect the full geographic range of early humans within a particular time interval because of differences in sampling intensity among regions, taphonomic potential among sites, settlement size and potential for detection, or the existence of multiple dates among artifacts from a single site (71, 72). To reduce the impact of these biases on our results, niche parameter and climatic suitability estimates counted each occupied climatic grid cell only once per time interval regardless of the number of dated remains therein, avoiding the artificial weighting of a subset of climatic conditions toward that of a few well-sampled grid cells (Figs. S4-S6). The algorithm to estimate the climatic niche uses a kernel density function, reducing the impact of differences in sample size. Second, while high resolution climatic reconstructions vastly improve our ability to investigate the processes governing human range expansion, even 1,000 year intervals cannot capture the abrupt climatic events (i.e. Heinrich events and Dansgaard-Oeschger events; 73) which likely affected the distributions of humans and other species (59), and the spatial resolution of 2 degrees hampers the ability for detecting refugia of small extent. However, even if the temporal resolutions of these reconstructions were higher, the precision of ^{14}C dating does not permit more detailed interpretations. Our results may therefore underestimate the abrupt nature of climate niche dynamics. Third, while each techno-cultural transition was implemented in our analysis as a single event, transitions were in reality more gradual, reaching different parts of Eurasia at

different times, while multiple lithic industries existed simultaneously even within Europe (e.g. Solutrean in the Atlantic side and Lower Magdalenian in the rest of Europe). Nevertheless, we used a broad classification of cultural periods (see *Results*), as a proxy for cultural and technological advancement, and including more complex variables to estimate niche construction using technological developments may fine-tune our findings in the future.

Despite these caveats, our approach pulls together complementary sources of paleo records in a unique quantitative framework, and reveals significant changes to the climatic niche during modern human dispersal across Eurasia under severe climate change. These methods provide insights which are distinct from, but complementary to, other modes of inference such as archeology or population genetics: for example, the timing and magnitude of changes in the ecological niche and geographical range could be used to inform population genetic hypotheses. Linking our framework to genetic evidences will allow exploring the effects of climate change on genetic diversity, population size and genomic evolution in Palearctic modern humans as has been done for other megafauna species (17, 18). We can also identify potential climatic refugia that may be targeted for future field work exploration to find previously undiscovered settlement sites and human remains.

Additionally, this approach can provide further clues as to where early modern humans may have overlapped in geographic and/or environmental space with archaic populations, such as Neanderthals (55, 74, 75) or Denisovans, as evidenced by their contribution to our genetic heritage (76-78), and may shed new light on the mechanisms, such as competition for resources, underlying their gradual geographical replacement and extinction during our global expansion.

Materials and methods

Human occurrences

Localities of fossils and other archaeological remains span Eurasia, from western Europe to western Beringia (north of 31° and 38° N latitude for Europe and Asia, respectively), and include 3,993 radiocarbon dated finds from 46.5ka-10.5ka. The majority were associated with Upper Paleolithic archaeological sites in western and central Europe, Siberia, and China. We focused on this temporal extent because at ~50ka Anatomically Modern Humans in Eurasia were largely restricted to the Near East (but see 1, 9, 79), while by 11ka they had completely occupied Eurasia, and had replaced or absorbed all archaic populations, particularly the Neanderthals (5, 80).

Localities of archaeological remains were collated from Ugan & Byers (81) (all Eurasia), Hamilton & Buchanan (33) (Siberia and northern China) and the International Union for Quaternary Research (INQUA) Radiocarbon Paleolithic Database,

v.13, excluding any data from North America or associated with *Homo neanderthalensis* remains or tool traditions. The data were standardised by excluding all specimen localities i) not associated with lab codes, ii) without reported errors for ^{14}C determinations, iii) duplicate ^{14}C estimates or iv) with ^{14}C error >10% of the mean age.

Climatic Data

Paleoclimatic conditions were simulated under the HadCM3 (*Hadley Centre Coupled Model, version 3*) Atmospheric-Ocean coupled General Circulation Model (AOGCM). The model was driven by changes to incoming solar insolation due to variation in orbital configuration, atmospheric greenhouse gas changes derived from ice core records and ice-sheet and sea level changes. The simulations have a time step of 1,000 years between 22ka and 11ka and of 2,000 years before 22ka, resulting in 25 intervals between 11ka-46ka (see 73, 82 for further details of experimental details).

The climatic niche was characterised on the basis of mean temperature ($^{\circ}\text{C}$) and total precipitation (mm) during the spring, summer, fall, and winter seasons. Seasonal variables were used because they are more likely than annual means to capture the climatic boundaries of a species' niche. Previous studies have used seasonal variables that captured climatic variability, rather than annual means, to model the climatic niches of other megafauna species (16). Each climatic surface was cropped to the appropriate land surface area for that period based on estimated changes in sea level and land surface incorporated into the AOGCMs. Study area is shown in Fig. 3.

Climatic niche overlap, niche breadth and niche marginality

An adaptation of Schoener's D metric for niche overlap (83) from Broennimann and colleagues (22) was implemented to measure the similarity in climatic niche occupied by Palearctic populations of modern humans between consecutive time periods. This metric has been previously used to quantify niche overlap between sister species or between different (i.e., native and introduced) ranges of a single species. D ranges from 0 (no overlap; niches are completely different) to 1 (complete overlap; niches are identical). This framework compares species' niches directly in climatic, rather than geographic, space, and uses a kernel density function to determine the 'smoothed' density of occurrences in each cell in the niche space (22). Consequently, it reduces the effects of an uneven distribution of archaeological localities relative to the human climatic niche and the uneven availability of environmental conditions between the range between time periods (i.e., a particular set of conditions may occur over extensive geographic space during one period but occupy significantly less space during the subsequent period). Niche overlap (D) was calculated in R (R Development Core Team).

277 The Outlying Mean Index (OMI), an ordination technique, was used to assess changes in niche breadth and niche
278 marginality through time (23), utilising the package ade4 in R (R Development Core Team, CRAN) (84). OMI is used to
279 assess the niche separation and breadth of species assemblages or closely related species across the main environmental axes
280 (24) (see *SI Outlying Mean Index*) but it was applied here to the climatic niche of modern humans across time. OMI quantifies
281 the environmental conditions occupied by each species (niche breadth) and calculates the distance between the average
282 environmental conditions occupied by each species (i.e. niche centroid) and the average conditions of the study area, or niche
283 marginality. In this study we used a climatic niche space that encompasses all the available climatic conditions across the 25
284 climatic simulation intervals.

285 **Climate envelope models**

286 Climatic Envelope Models (CEMs) were used to reconstruct the distribution of climatic suitability for Paleolithic modern
287 humans in the Palearctic for each time interval, and to identify areas of relative climatic stability across the time extent of the
288 study. CEMs were constructed using Maximum Entropy (MaxEnt; 85, 86) in R (R Development Core Team, CRAN) and the
289 package Dismo (87). Modeling was only performed for time intervals with at least 2n localities (n= number of predictor
290 climatic variables). All other parameters for the models were implemented as the default settings. The human occurrence data
291 were randomly split to 70%-30%, using 70% of localities to calibrate the models, and 30% as a validation dataset to evaluate
292 the models' predictive accuracy (but see 88). This process was repeated 10 times for each time interval, and as the final
293 suitability map is the average of these 10 repetitions.

294 CEMs were validated using the Continuous Boyce Index ($B_{cont(W)}$), which does not require setting a threshold value for
295 environmental suitability, following the implementation by Hirzel and colleagues (89). After partitioning the climatic
296 suitability range into 10 evenly spaced bins, the *predicted frequencies* of archaeological localities for each climatic suitability
297 bin were correlated with the *expected frequency* based on the relative geographic area within each bin. The Spearman rank
298 correlation of *predicted/expected* frequencies against the mean climatic suitability of each bin provides the Boyce index. The
299 $B_{cont(W)}$ varies from -1 to 1, with higher values of suitability indicating good model fit (validation localities are predicted in
300 areas with high suitability values), values close to 0 indicating a model no better than random, and negative values indicating a
301 poor model (validation localities are predicted in areas with low suitability values). Model validation was performed for the
302 average model per time period.

303 Schoener's D metric as implemented by Warren et al (29) (i.e., in geographic, rather than climatic, space), was used to
304 quantify the degree of geographic overlap in climatically suitable areas between consecutive time intervals. To define regions
305 of long-term high climatic suitability, the cumulative climatic suitability per grid cell across all time intervals was divided by
306 the number of intervals each grid cell was above sea level and not covered by ice. The regions of consistently high climatic
307 suitability were measured as the 30% of grid cells with the highest mean climatic suitability, following an approach similar to
308 Graham et al (90).

309 **Cultural and climatic drivers of niche dynamics**

310 Generalized Additive Models (GAMs) were used to assess the effects of climate change and transitions between 'cultural
311 periods' as drivers of niche parameters and distribution changes in climatic suitability of Palearctic modern humans. Changes
312 in the seasonal medians across fossil localities between consecutive time intervals for temperature and total precipitation were
313 employed as single climatic predictors. Each GAM was run using a single climatic predictor because of the small number of
314 time periods relative to the number of climatic variables. This was performed both including and excluding changes between
315 'cultural periods', which were measured as a factor with 5 levels (1 is the earliest cultural period while 5 is the most recent).
316 Lastly, a series of GAMs was run using cultural changes as a single predictor.

317 **Cultural Periods**

318 We used a broad classification of 'cultural periods' identified by lithic industries: Initial Upper Paleolithic (UP) (45ka-
319 40ka); Aurignacian Europe/ Early UP Siberia, 40ka-32ka; Gravettian Europe /Middle UP Siberia, 32ka-24ka; Glacial
320 Maximum Europe/Middle UP Siberia, 24ka-17ka; Late Glacial Europe/Late UP Siberia, 17ka-11ka, partially modifying the
321 chronological periods defined by (35) for the European part. Even though Siberia has three main cultural periods within the
322 UP, they were divided into four based on the European archaeological record, where most of the fossil occurrences of our
323 database are found, permitting a common analysis of European and Asian data. Despite these differences between regions of
324 Eurasia, previous studies state that technological innovations spread quickly, suggesting population connections between
325 central-South Siberia and central-Eastern Europeans (49-51).

326

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499 Fig.1: Theoretical representation of niche parameters in climatic space. A) Change in the climatic niche (expansion
500 and contraction along climatic axes) and in the position of the niche centroid (black line) between two time periods.
501 The grey and brown ovals represent the climatic niche at two different time intervals, while the overlap between
502 intervals is shown in green. B) The brown circle represents the climatic niche in one time interval. Grey rectangles
503 represent climatic conditions of individual grid cells in paleoclimatic maps, while black dots and their relative sizes
504 indicate the presence and abundance of human occurrences within a particular set of climate conditions. G_{area} is the
505 center of gravity of the total climatic space from 46ka to 11ka of the study area, while G_{hum} is the average climatic
506 conditions of the climatic niche of modern humans, and the purple line represents the niche marginality, or the
507 distance between these two points. The blue dashed line represents the niche breath or the total climatic conditions
508 occupied by modern humans in a time period.
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Fig. 2: Temporal trends of climatic niche parameters for Anatomically Modern Humans between 46-11ka. The upper panel (2A) of the graph indicates overlap of the climatic niche (dashed brown line) and the geographic overlap of climatically suitable areas (light green line) between consecutive time intervals. The middle panel (2B) represents the changes in niche breadth (dashed light blue line) and niche marginality (purple line). The lower panel (2C) indicates the number of fossil occurrences used per time interval for all analyses. The climatic periods Marine Isotope Stage 3 and Marine Isotope Stage are indicated by a yellow bar, while different cultural periods are indicated by the grey bars behind the three panels (names are shown at the top).

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Fig. 3: Suitable climatic corridor for potential dispersal across Eurasia. The maximum extent of the study area, including all grid cells above sea level for at least one time interval, is outlined in black. The colored areas represent the 30 percent of the grid cells with the highest average suitability values across time bins between 46ka and 11ka. Grey areas inside the black outline represent the remaining 70 percent. Red pins represent locations of Late Pleistocene human findings; 1 - Yana RHS site, 2 - Mal'ta, 3 - Denisova cave, 4 - Ust'-Ishim, 5 - Kostyonki.