

DISENTANGLING GEOMAGNETIC AND PRECIPITATION SIGNALS IN AN 80-KYR CHINESE LOESS RECORD OF ^{10}Be

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ABSTRACT. The cosmogenic radionuclide ^{10}Be is produced by cosmic-ray spallation in Earth's atmosphere. Its production rate is regulated by the geomagnetic field intensity, so that its accumulation rate in aeolian sediments can, in principle, be used to derive high-resolution records of geomagnetic field changes. However, ^{10}Be atmospheric fallout rate also varies locally depending on rainfall rate. The accumulation rate of ^{10}Be in sediments is further complicated by overprinting of the geomagnetic and precipitation signals by ^{10}Be attached to remobilized dust, which fell from the atmosphere at some time in the past. Here, we demonstrate that these signals can be deconvoluted to derive both geomagnetic field intensity and paleoprecipitation records of Asian Monsoon intensity in an 80,000-yr-long ^{10}Be record from Chinese loess. The strong similarity between our derived paleomagnetic intensity record and the SINT 200 (Guyodo and Valet 1996) and NAPIS 75 (Laj et al. 2002) stacked-marine records suggests that this method might be used to produce multimillion-yr-long records of paleomagnetic intensity from loess. This technique also reveals a new method for extracting quantitative paleoprecipitation records from continental interior regions. Our derived precipitation record is broadly similar to the speleothem $\delta^{18}\text{O}$ -based records of paleo-Asian Monsoon intensity from Dongge (Yuan et al. 2004) and Hulu (Wang et al. 2001) caves, and suggests that the paleo-Asian Monsoon intensity may be responding to a combination of both Northern and Southern Hemisphere insolation forcing.

INTRODUCTION

Wind-blown loess sediments have been accumulating in central China nearly continuously for the last 2.6 million yr (Liu and Ding 1998). These sediments are a storehouse of very long and potentially high-resolution records of both variations in the geomagnetic field, and of variability in the East Asian Monsoon. While there have been numerous previous efforts to extract climate or magnetic intensity records from Chinese loess (An et al. 2000; Beer et al. 1993, 2002; Evans and Heller 2001; Gu et al. 1996; Kukla et al. 1990; Maher and Thompson 1995; Pan et al. 2001; Porter et al. 2001; Shen et al. 1992; Zheng et al. 1995), the exercise has proved challenging because the climate and geomagnetic signals contained in these sediments are tightly intertwined. This is true for both magnetic susceptibility as well as for records of cosmogenic isotopes found in loess, such as ^{10}Be , both of which have been used in attempts to reconstruct climate and geomagnetic field variations. For paleomagnetic reconstructions based on ^{10}Be , some ^{10}Be comes from remobilized dust that fell to Earth at some time in the past, whereas a second component derives from atmospheric fallout of new ^{10}Be recently generated by cosmic-ray interactions in the atmosphere. It is the flux of this fallout component that is regulated by the geomagnetic field intensity. Unfortunately, interpretation of fallout ^{10}Be is further complicated by the fact that some of it makes its way to the ground via adsorption onto dry particulates, but the rest arrives by wet precipitation (Wallbrink and Murray 1994). While the dry fallout fraction is usually less than 10% (Wallbrink and Murray 1994), the local ^{10}Be fallout flux to the ground depends strongly on the wet precipitation amount. To further complicate matters, solar magnetic field variations can also influence atmospheric ^{10}Be production via fluctuations in solar wind rigidity (Masarik and Beer 1999), though solar forcing mainly acts on shorter periods than are being considered here.

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In this paper, we argue that the amount of magnetic susceptibility and ^{10}Be associated with recycled dust can be calculated and eliminated by using magnetic coercivity and magnetic susceptibility measurements together. Once the recycled dust effects are removed, we will show that the residual susceptibility and ^{10}Be signals are controlled by a combination of both climate and geomagnetic field intensity. Because both signals are overprinted by climate in a similar manner, their cross-correlation offers a simple way to remove the climate contamination from the ^{10}Be record, yielding a signal dependent only on geomagnetic field intensity. Once the geomagnetic field signal is extracted, we are then able to recover a record of paleorainfall intensity by the separation of variables.

Other authors have previously noted that magnetic susceptibility (intensity of magnetization/unit mass in response to a defined applied field) and ^{10}Be concentration in Chinese loess are correlated with rainfall (Beer et al. 1993, 2002; Heller et al. 1993; Maher and Thompson 1995; Porter et al. 2001; Shen et al. 1992), and have used this fact in various approaches to deduce records of paleoprecipitation or geomagnetic variability. In one instance (Maher and Thompson 1995), the correlation between modern rainfall and total magnetic susceptibility in surficial loess/paleosols was used to derive a paleoprecipitation record. The chief difficulty with this method is in first constraining the effects of recycled dust on magnetic susceptibility. In a second approach (Porter et al. 2001), an attempt to remove the dust component of magnetic susceptibility is first made using a grain size model, then the residual susceptibility is correlated to modern precipitation patterns. In a third method, Beer et al. (1993) and Heller et al. (1993) used loess ^{10}Be concentrations to first constrain and remove the fractions of magnetic susceptibility associated with recycled dust in surface loess/paleosols, then they correlate the residual signal with modern precipitation for various regions of the loess plateau to establish a modern calibration, which they then use to reconstruct paleoprecipitation. While this last technique does remove the component of susceptibility associated with inherited dust, it does not account for variations in the ^{10}Be signal associated with geomagnetic modulation of ^{10}Be , which accounts for nearly half of the total ^{10}Be signal.

In our method, we modify these approaches by first removing the inherited dust signal from both magnetic susceptibility and ^{10}Be concentration records, then we extract the rainfall effect using the correlation between the inherited-dust-free ^{10}Be and magnetic susceptibility fractions. The residual ^{10}Be signal is then converted to a flux using the sediment accumulation rate, divided by the range in flux, and then normalized to the modern production flux. This signal is then used to provide a record of geomagnetic fluctuations using the production rate dependence on magnetic field intensity (Masarik and Beer 1999). Once the time function describing the variations in ^{10}Be flux due to geomagnetic-field fluctuations is established, we solve for the variations in ^{10}Be flux due to changes in precipitation by dividing the dust-free ^{10}Be fluxes by the geomagnetic-field production-rate modulation function to obtain a record of variations in ^{10}Be flux due to changes in wet precipitation. Finally, we use the correlation between ^7Be in modern precipitation (Caillet et al. 2001; Ishikawa et al. 1995; Wallbrink and Murray 1994) and tropospheric $^{10}\text{Be}/^7\text{Be}$ ratio to derive quantitative estimates of paleoprecipitation.

METHODS

Sampling and Measurements

Analyses were performed on a 970-cm-long core taken from the Luochuan loess section in central China ($35^{\circ}45'\text{N}$, $109^{\circ}25'\text{E}$) (Figure 1). This core was sampled at approximately 100-yr intervals (1 cm) for grain size analysis, dry bulk density, and magnetic susceptibility measurements at the Xi'an Laboratory of Loess and Quaternary Geology (Figure 2 and Appendix: Table 2). The core was sampled at 4-cm intervals for ^{10}Be measurements, except during the interval representing the last

glacial maximum (LGM), where it was sampled at 1-cm intervals. Additional details on sampling methodology are given in the Appendix. The ^{247}BeO samples were chemically prepared at the Xi'an Laboratory of Loess and Quaternary Geology. The majority of these samples were measured for ^{10}Be on the 3MV accelerator mass spectrometer of VERA (Vienna Environmental Research Accelerator) at the University of Vienna (Priller et al. 2000). Approximately 20% of the ^{10}Be analyses were made on the 3MV accelerator mass spectrometer at the NSF-Arizona AMS Facility at the University of Arizona. All ^{10}Be concentrations have been corrected for radioactive decay (Appendix: Table 2). ^{10}Be AMS measurements at both VERA and Arizona were made at 3MV accelerating voltage using 1–3 μA $^9\text{Be}^{16}\text{O}^-$ target injection currents. VERA used gas stripping in the terminal to 3+ for high-energy analysis, whereas Arizona used gas stripping to 2+ in the terminal and then foil stripping of the ions to 3+ after high-energy magnetic and electrostatic separation at 2+, but prior to analysis through a second 1.2-m-radius electrostatic analyzer (ESA) and switching magnet. Both labs used a 2-stage dE-dx coincidence gas analyzer in which the interfering isobar ^{10}B is completely stopped in the first stage. These analysis techniques are described in greater detail in Priller et al. (2004) (VERA) and McHargue et al. (2000) (Arizona).

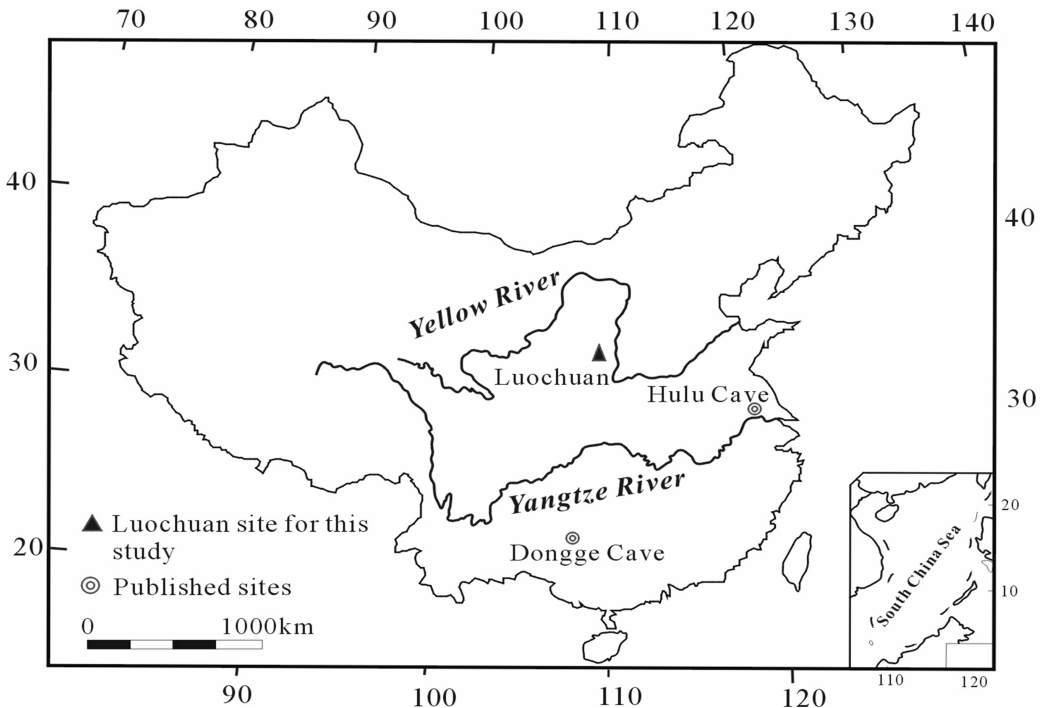


Figure 1 Map showing the location of Luochuan where the sample core was taken, in the context of the regional setting of the Chinese Loess Plateau. Also shown are the locations of Hulu Cave and Dongge Cave, from which $\delta^{18}\text{O}$ records representing the East Asian Monsoon intensity were used for comparison with the paleoprecipitation reconstruction from Luochuan.

Time Scale

A calendar age model for this loess sequence was generated using a grain size vs. accumulation rate model (Porter and An 1995; Wu 2004). This calendar age scale was then compared to a combination of optical luminescence (OSL) and ^{14}C ages for the younger portion of this time scale (Figure 2 and

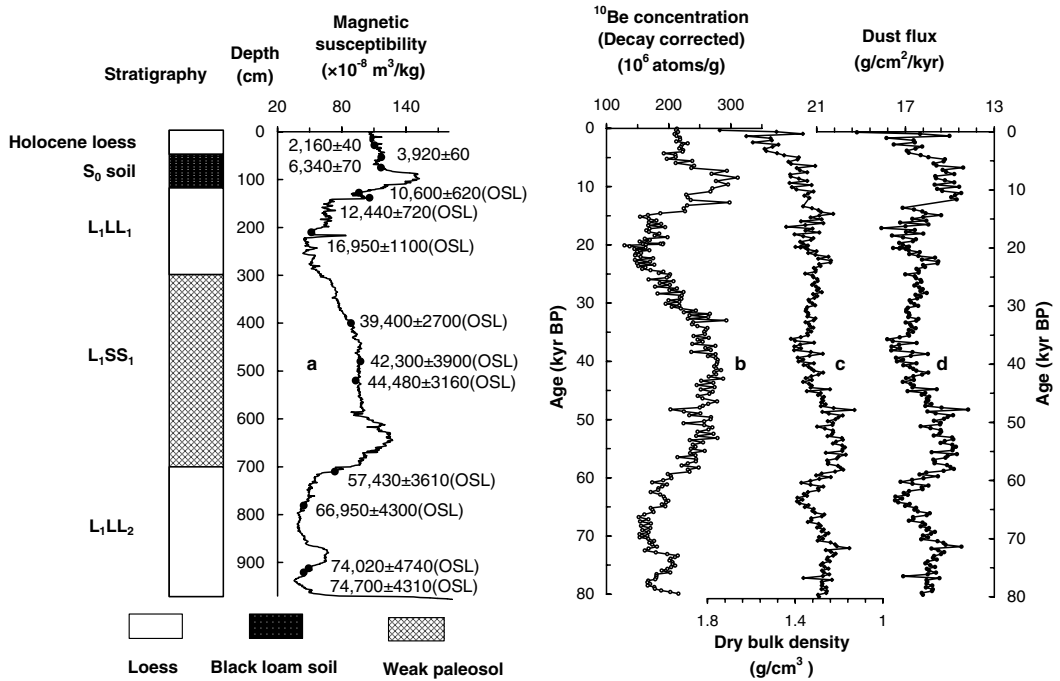


Figure 2 Loess magnetic and ^{10}Be stratigraphy at Luochuan, China ($35^{\circ}45'\text{N}$, $109^{\circ}25'\text{E}$). Shown from left to right: loess/paleosol stratigraphy; sample depths; magnetic susceptibility (magnetic dipole moment/unit volume in SI units) with locations of OSL and ^{14}C dates superimposed (black dots); uncalibrated ages (kyr BP) (calibrated ages listed in Appendix: Table 3); ^{10}Be concentration (10^6 atoms/g); dry bulk density; and dust flux ($\text{g/cm}^2/\text{kyr BP}$). Dust flux is calculated from the age model and dry bulk density data (see Appendix).

Appendix: Tables 3, 4). For the older portion of the record, we cross-checked the age model using a correlation (Porter 2001) between the magnetic susceptibility (SUS) curve from the loess sequence with the SPECMAP marine oxygen isotope (MIS) curve, and using inferred L1/S0 (Gu et al. 1996) (12.3 kyr BP) and S1/L1 (GRIP ^{18}O age $79 \pm 1 \text{ kyr BP}$ [Johnsen et al. 2001]) stratigraphic boundaries (Liu and Ding 1998), which correlate with MIS 2/1 and 5/4 transitions (Porter and An 1995). The calendar ages provided by this grain size model were found to be in good agreement with these external cross-checks.

RESULTS AND DISCUSSION

Geomagnetic Field Intensity Deconvolution

To isolate the ^{10}Be component associated with reworked dust, we first define the measured magnetic susceptibility (SUS(M)) as the sum of 2 components, SUS(D) and SUS(P), of which the former is the inherited dust-borne fraction of SUS and the latter is pedogenic fraction of SUS, which is acquired through secondary mineral reactions after deposition of the loess, respectively (see Table 1 for a summary of term definitions).

To solve for SUS(D), we note that a comparison of loess magnetic susceptibility versus coercivity (Evans and Heller 2001) from a wide range of locations on the Loess Plateau for the last 135 kyr reveals a pattern suggestive of 2-component mixing between a low-SUS/high-coercivity inherited-dust component, with admixtures of high-SUS/low-coercivity magnetic domains associated with

Table 1 Definition of terms used in this paper.

Terms	Explanation
Be(M)	The measured adsorbed ^{10}Be concentration in loess corrected for radioactive decay.
Be(D)	The concentration of recycled ^{10}Be adsorbed on remobilized dust.
Be(P,GM)	Equal to Be(M) – Be(D). Equivalent to the ^{10}Be concentration in loess coming from atmospheric fallout recently produced in the atmosphere by cosmic rays.
SUS(M)	The measured magnetic susceptibility.
SUS(D)	The fraction of SUS(M) inherited from recycled dust.
SUS(P)	Equal to SUS(M) – SUS(D). Equivalent to the pedogenic fraction of SUS(M), which is acquired through secondary mineral reactions after deposition of the loess.
Be(GM)	That fraction of the Be(P,GM) concentration signal that is due to fluctuations in the geomagnetic field, as opposed to variations in wet precipitation.

pedogenesis (Figure 3). These data asymptotically approach a well-defined susceptibility value of $(25 \pm 3.5) \times 10^{-8}$ (1σ) m^3/kg for the high-coercivity endmember (Evans and Heller 2001). In accord with the proposal made by these authors that this endmember represents a dry dust component of the loess susceptibility, samples at this end of the mixing curve corresponds to those from the driest periods, typical of the last glacial maximum (LGM) and marine isotope stage 4 (MIS4) when rainfall was thought to be very low in this region and pedogenesis was very slow. The fact that samples close to this endmember come from a wide geographic distribution from both the western and central Loess Plateau suggests that this dust endmember is spatially homogeneous with regards to susceptibility (Evans and Heller 2001). Recalling that susceptibility does not depend on dust flux rate but rather only on the concentration of magnetic domains and their size distribution, Evans and Heller (2001) found that for the most weakly weathered loess (which they identify as being from the driest periods), the mass fraction of hematite, magnetite, and maghemite were on average 0.9%, 0.02%, and 0.02%, respectively, and that these concentrations are quite uniform over the geographic extent of the Chinese Loess Plateau. Furthermore, long records from the Loess Plateau show that loess magnetic susceptibility has been similar for each dry climate phase during the last 1 Myr (Evans and Heller 2001). We use these arguments collectively to contend that there is a dry dust endmember component of SUS in the loess that is spatially and temporally homogeneous with a well-defined susceptibility value on a plot of coercivity versus susceptibility. We define a value for this SUS(D) component as the high-coercivity asymptotic endmember on this plot, yielding a value of $\text{SUS(D)} = (25 \pm 3.5) \times 10^{-8}$ (1σ) m^3/kg . We then solve for SUS(P) for all times from the difference between SUS(D) and measured SUS(M) (Figure 4a).

Next, to solve for Be(D) (the adsorbed ^{10}Be concentration associated with remobilized dust), we define the total measured ^{10}Be concentration in the loess (Be(M)) as the sum of 2 sources: Be(D), which is ^{10}Be from atmospheric fallout produced at some time in the past, which was then adsorbed on dust, buried, and later remobilized; and Be(P,GM), which is ^{10}Be fallout recently produced in the atmosphere by cosmic rays (modulated by the geomagnetic field intensity) and brought to the ground primarily by wet precipitation. We argue that Be(D) (i.e. the adsorbed ^{10}Be concentration in dust, *not* the ^{10}Be dust flux) should be highly uniform, because the dust falling on the Loess Plateau has been derived from a very large area (e.g. the Gobi and Taklimakin deserts and Loess Plateau) representing landscapes of many ages, and transported by a variety of aeolian processes over large distances during which substantial mixing occurs.

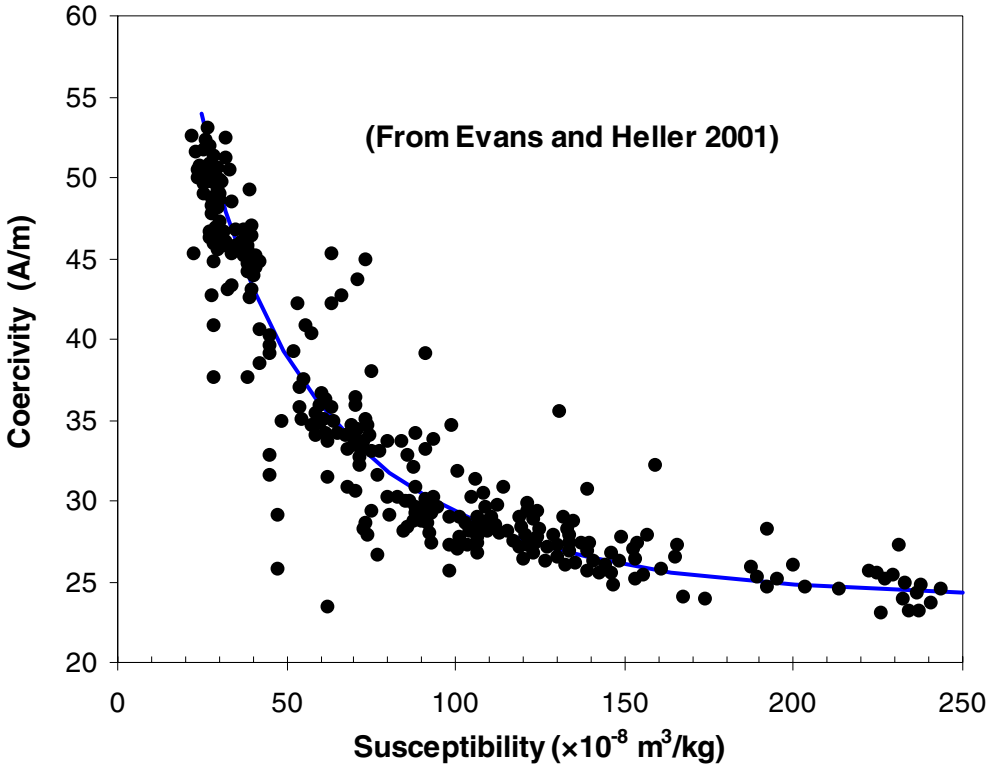


Figure 3 Loess magnetic susceptibility versus coercivity for samples from the western and central Chinese Loess Plateau. Note that the data array is suggestive of 2-component mixing between a low-susceptibility/high-coercivity recycled dust component, with admixtures of high-susceptibility/low-coercivity mineral grains added by pedogenesis of the loess during wetter epochs. We define a susceptibility value for the (dry) dust endmember component (SUS(D)) as the high-coercivity asymptotic endmember on this plot, yielding a value of $SUS(D) = 25 \pm 3.5$ (1σ). Reprinted from Evans and Heller (2001), with additional thanks to Elsevier for copyright permission.

As in the case for susceptibility, we expect that during extremely dry periods, loess ^{10}Be concentration is dominated by the remobilized dust fraction $\text{Be}(D)$, because nearly all (>90%) atmospheric fallout of new ^{10}Be recently generated by cosmic-ray interactions results via wet precipitation (Wallbrink and Murray 1994). We show that $SUS(M)$ and $\text{Be}(M)$ are highly linearly correlated (Figure 4b). We determine the dry dust endmember $\text{Be}(D)$ by using this correlation and the previous observation that $SUS(D) = 25 \times 10^{-8} \text{ m}^3/\text{kg}$. This yields a value of $\text{Be}(D) = (136.4 \pm 6.25) \times 10^6$ (1σ) atoms/g. As we already showed for $SUS(P)$, we solve for $\text{Be}(P,GM)$ by calculating the difference between $\text{Be}(M)$ and $\text{Be}(D)$ for the entire record (Figure 4c).

After subtracting the effects of reworked dust, we observe that both $SUS(P)$ and $\text{Be}(P,GM)$ form a linear array (Figure 4d). We assert that this linear correlation is due to the effect of variations in wet precipitation rate on the fallout flux of ^{10}Be on the one hand, and on the growth rate of pedogenic magnetic domains in the loess on the other hand. This linear relationship has a correlation coefficient of 0.81, which indicates there is some other source of variance. We believe that much of this dispersion is due to the fact that $\text{Be}(P,GM)$ is also a function of geomagnetic field intensity in addition to the wet precipitation rate. Since it is likely that the variance associated with geomagnetic field variations is independent of climate, we may then remove the precipitation effect from

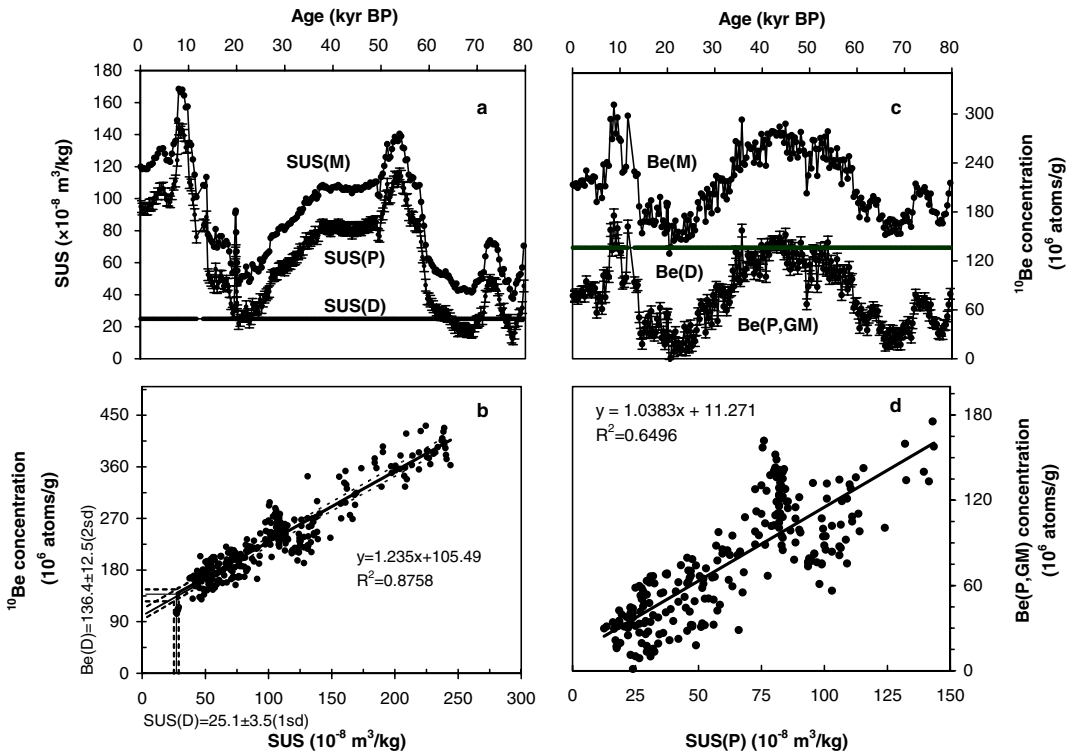


Figure 4 Deconvolution of loess magnetic susceptibility and ^{10}Be records. a) Measured magnetic susceptibility (SUS(M)) vs. time, decomposed into SUS(D) (dust) and SUS(P) (pedogenic) fractions. b) Measured magnetic susceptibility (SUS(M)) vs. measured ^{10}Be concentration (Be(M)). Be(D) is derived from this plot, from the coincidence of SUS(D) with the model linear regression through the data. The uncertainty in Be(D) is derived graphically from the intersection of the (2σ) uncertainty envelope around the regression with the (1σ) errors on SUS(D) determined from regression analysis of Figure 3. c) Be(M) vs. time decomposed into Be(D) and Be(P,GM) (associated with atmospheric fallout and modulated by rainfall and geomagnetic field strength). d) Linear correlation between SUS(P) and Be(P,GM). This correlation defines the fraction of variance associated with the wet precipitation amount effect. Deviation from this line for each datum defines the residual variance assumed to be associated with geomagnetic modulation.

Be(P,GM) using the correlation between SUS(P) and Be(P,GM). After removal of the precipitation effect by subtraction of this linear correlation, the residual variations in Be(P,GM) (i.e. Be(GM)) due to geomagnetic modulation are revealed (not shown). Finally, we convert these residual Be(GM) concentration variances to flux using sediment accumulation rates, normalize them by dividing by the range, then scale them to the modern production rate (Masarik and Beer 1999), yielding a curve of ^{10}Be production rate relative to today (Figure 5a).

Reconstructed Magnetic Field Record

Figure 5a shows the record of relative variations in ^{10}Be production rate for the last 80 kyr BP, which we derived above. In Figure 5b, we convert this ^{10}Be production rate record to paleomagnetic field intensity (Masarik and Beer 1999), with the tacit assumption that solar modulation of ^{10}Be production rate is small relative to geomagnetic modulation at least for long period fluctuations. This plot shows that the low-frequency features of our paleomagnetic intensity record compare favorably with the SINT200 composite reconstruction (Guyodo and Valet 1996) (Figure 5b). These same broad features are also observed in the NAPIS 75 record (Laj et al. 2002), though its features are offset because

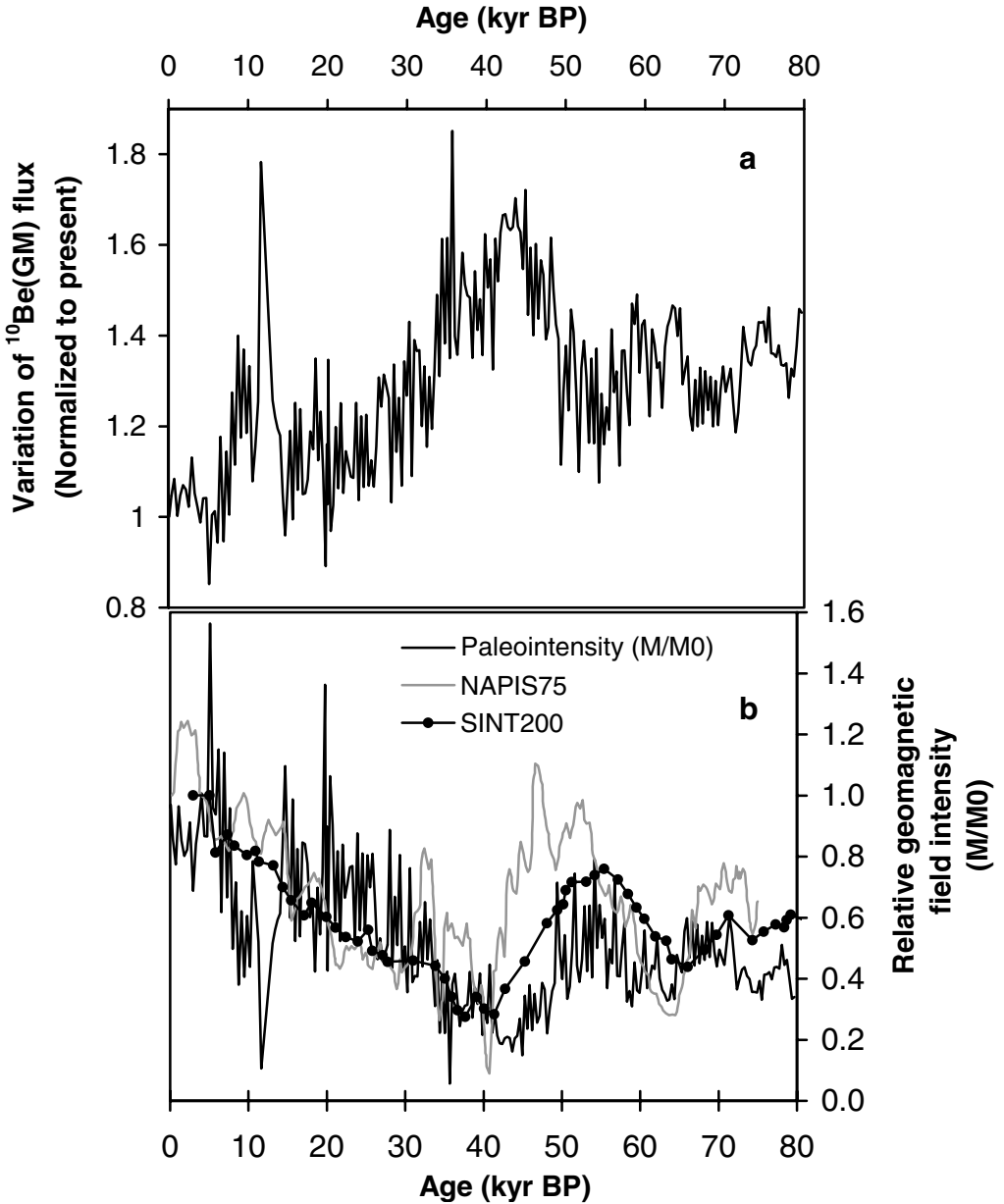


Figure 5 Deconvoluted ^{10}Be production rate variations and magnetic field variations. a) Loess-based ^{10}Be production rates normalized to the modern rate, after removal of the reworked dust signal and variations due to changes in wet precipitation. The data are presented as ^{10}Be flux, using sediment accumulation rate to convert from concentration. b) Inferred variations in geomagnetic field (black) normalized to the modern field, compared to the composite records NAPIS75 (Laj et al. 2002) (gray) and SINT200 (Guyodo and Valet 1996) (black with dots). Our magnetic field record was derived from inversion of the ^{10}Be production rate record shown in (a), using the model provided by Masarik and Beer (1999), and the assumption that all fluctuations in production rate are due to changes in geomagnetic field. As discussed in the Appendix in "Propagation of Errors," mean uncertainty in calculated M/Mo is 21%.

it utilizes a different calendar age model. Our record clearly shows the Laschamp geomagnetic minima events at about 42 kyr BP, as well as a smaller local minima at about 60–65 kyr BP, and geomagnetic maxima at about 55 and 70 kyr BP observed in these 2 composite records. The long rise in geomagnetic intensity common to most records observed between about 40 and 5 kyr BP is also observed in our record, as is an abrupt drop in intensity (reflected by an increased ^{10}Be flux in Figure 1a) at about 33 kyr BP, which we tentatively assign to the Mono Lake event (Wagner et al. 2000). There is also a prominent late Holocene paleomagnetic maximum similar to that seen in archaeological records (Yang et al. 2000), though in our record this maxima occurs $\sim 1\text{--}2$ kyr earlier.

One feature found in our record not expressed in either the SINT200 or NAPIS75 records is a period of apparently enhanced ^{10}Be concentration between about 15.5 and 8 kyr BP. If this feature were due to increased ^{10}Be production rate, we might expect a similar anomaly in the record of atmospheric ^{14}C (Reimer et al. 2005). However, while atmospheric ^{14}C concentrations were indeed significantly increased during the Younger Dryas–Bølling–Allerød part of this time frame (13.8 and 11.3 kyr; Hughen et al. 2004), this period of elevated ^{14}C is much shorter in duration than is seen in our loess record. This suggests that there may still be climatic influences in this record that we have failed to isolate. Nevertheless, the strong similarity among SINT 200, NAPIS 75, and the long wavelength features of our paleomagnetic intensity record suggests that this method of generating paleomagnetic intensity records is generally robust. If so, this technique could be used to generate an extremely long-term high-resolution record of paleomagnetic field intensity since the loess record extends back about 2.6 Myr BP, and this is less than just 2 half-lives of ^{10}Be .

Paleoprecipitation Reconstruction

We also can generate a paleoprecipitation record by beginning with Be(P,GM) , i.e. the inherited-dust-free concentration of ^{10}Be signal in loess, as derived above. We recall that Be(P,GM) varies in response to the product of the wet precipitation amount and the ^{10}Be production rate (modulated by geomagnetic field). We first convert Be(P,GM) to a ^{10}Be flux by multiplying Be(P,GM) by the loess accumulation rate (derived from the age model), then we remove the variations in $\text{Be(P,GM)}_{\text{flux}}$ due to geomagnetic modulation by dividing $\text{Be(P,GM)}_{\text{flux}}$ by the geomagnetic modulation function derived above from subtraction of the cross-correlation between SUS(P) and Be(P,GM) . This removes flux variations that are due to magnetic field fluctuations. The residual variations in ^{10}Be flux (i.e. $\text{Be(P)}_{\text{flux}}$) are then considered to be due to fluctuations in wet precipitation.

We derive quantitative estimates of paleoprecipitation from this residual signal using ^7Be as an analogue for ^{10}Be , and using the relationship between ^7Be concentration and precipitation amount in modern rainfall. While ^7Be has a similar atmospheric production pathway to ^{10}Be , its very short half-life means that there is no recycled dust component to confuse its relationship with rainfall amount. In 3 different ^7Be vs. precipitation amount studies on 3 continents (Caillet et al. 2001; Ishikawa et al. 1995; Wallbrink and Murray 1994), a similar linear relationship is found. Using the average slope in these studies, we then convert from ^7Be to ^{10}Be using the known $^{10}\text{Be}/^7\text{Be}$ flux ratio in modern tropospheric precipitation (Priller et al. 2004). We use the slope of this resulting ^{10}Be vs. precipitation line to calculate precipitation amounts, but we first adjust the intercept (corresponding to the local dry fallout fraction) to match the ^{10}Be flux vs. precipitation observed at Luochuan today. In this way, a precipitation record can be extracted from $\text{Be(P)}_{\text{flux}}$, which accounts for both wet and dry fallout fractions.

Figure 6 shows the resulting paleoprecipitation record that we have derived from the loess ^{10}Be record after removal of the dust and geomagnetic modulation influences. The general shape of this curve appears well correlated with the speleothem $\delta^{18}\text{O}$ records from Dongge (Yuan et al. 2004) and

Hulu (Wang et al. 2001) caves in SE China, which are widely regarded as a robust record of Asian Monsoon intensity (Figure 6a). Our record indicates extremely low rainfall for the LGM and marine isotope stage 4 (MIS4), with precipitation rising gradually out of MIS4 to near-modern levels during MIS3. After about 35 kyr BP, precipitation began a long decline culminating in a 5-kyr-long minima during the LGM ending at about 20 kyr BP. Precipitation then increased stepwise in dramatic fashion at ~20 kyr, and again at 14.3 kyr. Precipitation was constant or decreased slightly between 13 and 11 kyr, followed by another abrupt precipitation increase culminating in a Holocene maximum at 9 kyr BP. Precipitation then dropped rapidly to a Holocene minimum at 5.8 kyr BP, followed by a modest rise to 4 kyr, then a gentle decay and another small rise to the present during the late Holocene.

While our precipitation record is broadly similar in shape to the Dongge and Hulu $\delta^{18}\text{O}$ records, there are some surprising differences. Both records show large millennial-scale fluctuations during the deglacial and Holocene; however, during MIS3 these are significantly damped in our record relative to that seen in the Dongge/Hulu $\delta^{18}\text{O}$ records. This is surprising considering that today, Luochuan is located near the northern boundary of SE Asian Monsoon influence, in a region of high rainfall gradient. Consequently, we had expected to see even higher sensitivity to variations in summer monsoon intensity at Luochuan than at Hulu or Dongge, which lie ~1000 km to the east and south, respectively. One could explain this observation if our record were smoothed at millennial scales by some geologic process such as post-depositional wind mobilization of fallout ^{10}Be .

Some large millennial-scale fluctuations are observed in our record, such as that seen at about 48 kyr BP—potentially linked with Heinrich event H5. Several other abrupt changes are seen during the deglacial, and abrupt drops in precipitation are also seen at about 28 and 35 kyr BP, which could be correlative with H3 or H4. Of particular interest are the deglacial and Holocene changes, which express many similarities with the Dongge/Hulu records. Two large precipitation increases are observed that are roughly coeval with meltwater pulses 1 and 2, suggesting a coupling between high and low latitude climate dynamics during the deglacial (Zhou et al. 2001). It is interesting to note, however, that at Luochuan the ~9 kyr BP Holocene precipitation maximum peaks at the same time as the Dongge record, but falls off much more rapidly, reaching a minimum at about 5.8 kyr BP, whereas Dongge continues to drop until ~2 kyr BP. This asynchronous behavior may be understood in terms of the penetration depth of summer monsoon moisture into the continental interior being linked to insolation-driven land-sea temperature differential (An et al. 2000).

Like the Dongge and Hulu records, the low-frequency component of our precipitation record resembles the summer (JJA) solar insolation curve for 30°N, except during MIS3, when it more strongly resembles the insolation differential between 30°N and 30°S (Figure 6b). This can be understood in terms of two of the chief elements of monsoon forcing: 1) land/sea temperature differential and 2) the cross-hemispheric atmospheric pressure gradient exhibited over the tropical ocean (Clemens and Prell 2003). Land-sea differential is largely governed by sensible heating over the Asian continent during boreal summer, which is strongly influenced by snow cover over the Tibetan Plateau centered on ~30°N. Similarly, according to some estimates, 80–90% of the moisture flux in the Asian Monsoon originates from the southern subtropical Indian Ocean (Clemens et al. 1996). This flux is driven by the interhemispheric pressure gradient, which is coupled to the tropical interhemispheric insolation gradient. It is perhaps not surprising then that the paleo-Asian Monsoon intensity appears to be responding to some combination of both Northern and Southern Hemisphere insolation forcing.

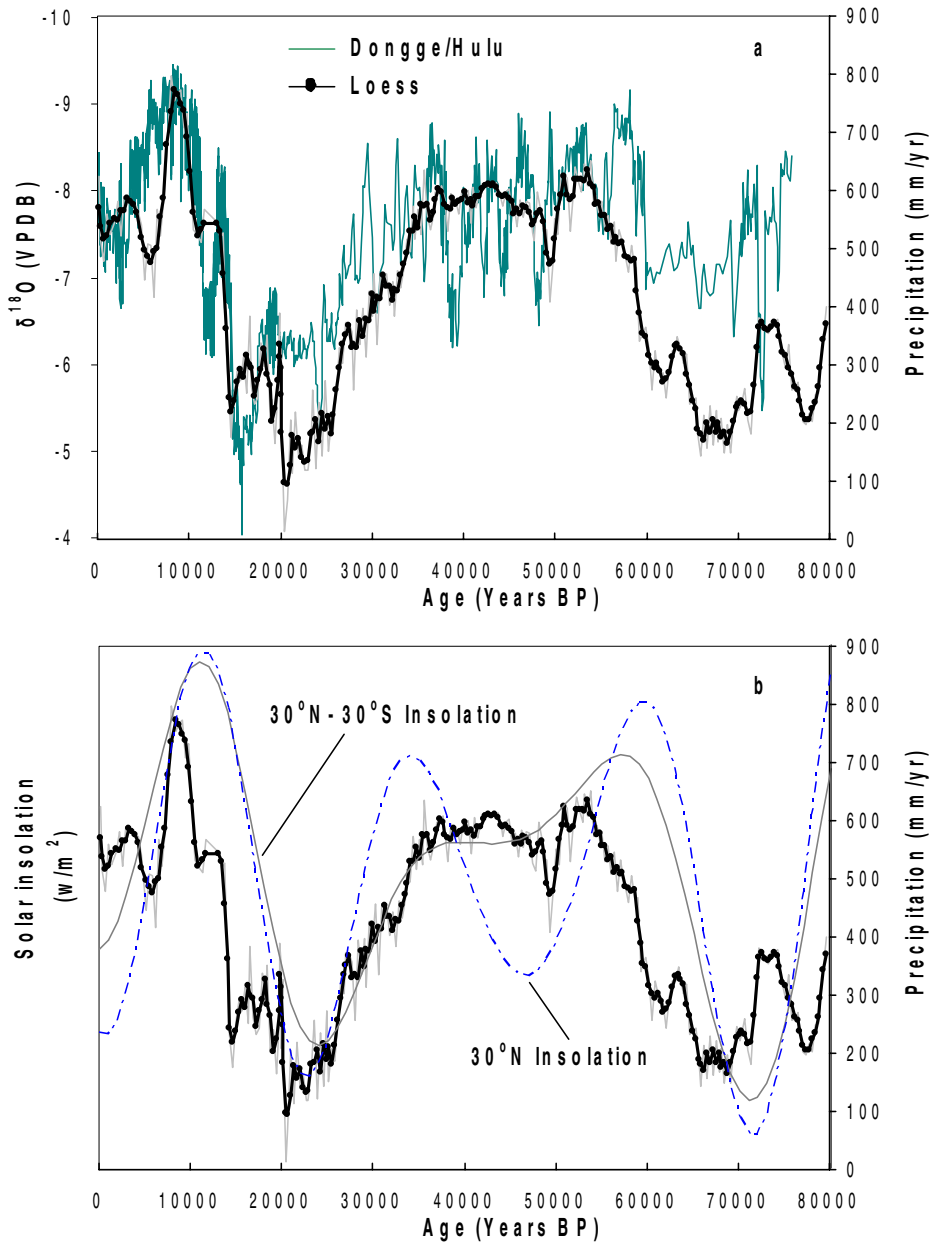


Figure 6 Paleoprecipitation reconstruction (mm/yr) derived from $\text{Be}(\text{P})$ after removal of the dust and geomagnetic modulation components of the ^{10}Be flux signal. a) Loess precipitation record is highly correlated with speleothem $\delta^{18}\text{O}$ records from Dongge and Hulu caves (Wang et al. 2001; Yuan et al. 2004), which are thought to be representative of variations in the Asian Monsoon intensity. As discussed in the Appendix in "Propagation of Errors," the mean uncertainty in calculated annual precipitation is 19.5%. b) Loess precipitation record compared to records of solar insolation (Berger and Loutre 1991). Like the Dongge and Hulu records, the low-frequency component of our precipitation record resembles the summer (JJA) solar insolation curve for 30°N , except during MIS3, when it more strongly resembles the insolation differential between 30°N and 30°S , suggesting that interhemispheric insolation gradient is also important in forcing the Asian Monsoon.

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APPENDIX

Loess Coring Procedures

The cores collected at Luochuan comprised a series of overlapping short cores from 2 parallel transects. Each core had a rectangular profile of about $15 \times 15 \text{ cm}^2$ and a length of about 40 cm. They were collected so as to overlap by about 5 cm at each end. During subsampling for individual measurements, the overlapping 5 cm at each core section end was discarded.

Magnetic Susceptibility (SUS) Measurement

SUS is a measure of induced magnetization of objects in an artificial weak magnetic field. The principal carrier of the magnetization in the loess and paleosol is fine-grained magnetite and maghemite.

The main carrier of the SUS signal is magnetic grains formed as an inorganic or biogenetic product of in situ pedogenesis (Maher and Taylor 1988; Maher and Thompson 1991; Zhou et al. 1990). SUS values directly reflect the intensity of pedogenesis and indirectly the amount of precipitation and intensity of the paleomonsoon (An et al. 1991). The SUS of the samples was measured in the laboratory using a Bartington MS2 susceptibility meter.

¹⁰Be Sample Preparation

¹⁰Be was extracted from loess samples using the following procedure (Wu 2004):

1. After drying in oven at 70 °C, approximately 1 g of sample was leached in 6N HCl along with 3 mL (30%) H₂O₂ and 0.5 mg ⁹Be carrier for 24 hr in an ultrasonic bath.
2. Centrifuge sample solution at 4000 rpm for 10 min.
3. Dry the supernatant at 80 °C in a Teflon[®] beaker, then dissolve salts with 1N 2 mL HCl, then centrifuge.
4. Add above supernatant to pretreated cation resin (Dowex 50WX–X8) column (plastic Bio-Rad column). The Be is eluted with 1N HCl. Discard the first 30 mL, then collect the next 80 mL.
5. Adjust pH of this solution to 8–9, then 14, then 8–9, in turn, with (25%) NH₄OH, (16%) NaOH, and (25%) NH₄OH, respectively. The Be should become a whitish Be(OH)₂ gel at this pH.
6. Centrifuge, discarding supernate.
7. Heat the Be(OH)₂ in a platinum crucible to about 850–900 °C in an oven for 2 hr. This oxidizes the Be(OH)₂ to BeO.
8. Mix the BeO powder with 200-mesh Cu powder (Cu:BeO = 3:1) and press into a Cu cathode for AMS analysis.

Depth, age, magnetic susceptibility, dry bulk density (DBD), and ¹⁰Be concentration data are tabulated in Table 2.

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ¹⁰Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ¹⁰Be concentrations to calculate ¹⁰Be fluxes.

Depth (cm)	Age (kyr BP)	¹⁰ Be concentration (Be(M)) (10 ⁶ atoms/g)	Magnetic susceptibility (SUS(M)) (10 ⁻⁸ m ³ /kg)	Dry bulk density (g/cm ³)
1	0.09	213	120	1.74
4	0.36	212	118.5	1.49
8	0.73	215	119	1.37
12	1.09	210	118.5	1.62
16	1.45	213	118.5	1.51
20	1.82	218	121	1.51
24	2.18	219	121	1.59
28	2.54	212	122	1.48
32	2.91	231	125.5	1.5
36	3.27	221	125.5	1.54
40	3.64	220	128	1.53
44	4	217	131	1.48
48	4.36	224	131	1.45
52	4.73	222	131.5	1.38
56	5.09	192	128	1.39
60	5.45	211	124	1.43
64	5.82	212	124	1.42
68	6.18	197	123	1.31

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ^{10}Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ^{10}Be concentrations to calculate ^{10}Be fluxes. (*Continued*)

Depth (cm)	Age (kyr BP)	^{10}Be concentration (Be(M)) (10^6 atoms/g)	Magnetic susceptibility (SUS(M)) (10^{-8} m ³ /kg)	Dry bulk density (g/cm ³)
72	6.54	238	128	1.39
76	6.91	211	134	1.38
80	7.27	241	136	1.35
84	7.63	237	149	1.42
88	8	294	168.5	1.42
92	8.36	269	166.5	1.38
96	8.73	311	168	1.35
100	9.09	276	164.5	1.43
104	9.45	296	157	1.32
108	9.82	270	157.5	1.42
112	10.18	267	135	1.39
116	10.54	228	133.5	1.32
120	10.91	231	124.5	1.34
124	11.27	235	115	1.36
128	11.63	298	101	1.34
132	13.07	230	108.5	1.36
136	13.39	224	108	1.32
140	13.71	226	113.5	1.29
144	14.03	187	77.5	1.28
148	14.35	167	76	1.23
152	14.67	154	74	1.29
156	14.99	168	72.5	1.33
160	15.3	180	70	1.28
164	15.62	168	81	1.37
168	15.94	188	69.5	1.27
172	16.26	167	72.5	1.35
176	16.58	194	76.5	1.44
180	16.9	169	76.5	1.32
184	17.22	164	70.5	1.35
188	17.53	168	72	1.29
192	17.85	185	73.5	1.4
196	18.17	178	71.5	1.36
200	18.49	198	67	1.32
204	18.81	162	59.5	1.33
208	19.13	169	53.5	1.38
212	19.45	154	54.5	1.36
216	19.77	165	91	1.34
217	19.84	191	92.5	1.35
218	19.92	187	79	1.37
219	20	159	67	1.4
220	20.08	190	60	1.38
221	20.16	164	55	1.35
224	20.4	129	47	1.34
228	20.72	137	49	1.3
232	21.04	163	51.5	1.34
236	21.36	145	51.5	1.31
240	21.68	169	51.5	1.25

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ^{10}Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ^{10}Be concentrations to calculate ^{10}Be fluxes. (*Continued*)

Depth (cm)	Age (kyr BP)	^{10}Be concentration (Be(M)) (10^6 atoms/g)	Magnetic susceptibility (SUS(M)) (10^{-8} m ³ /kg)	Dry bulk density (g/cm ³)
244	22	149	57.5	1.3
248	22.31	155	52.5	1.24
252	22.63	147	48	1.24
256	22.95	146	50	1.29
260	23.27	155	59	1.28
264	23.59	175	57	1.31
268	23.91	146	56	1.31
272	24.23	170	56	1.31
276	24.54	149	54.5	1.35
280	24.86	177	62	1.32
284	25.18	151	57	1.32
288	25.5	163	61	1.3
292	25.82	158	63.5	1.3
296	26.14	171	65	1.31
300	26.46	192	66	1.32
304	26.78	184	67	1.3
308	27.09	203	75.5	1.29
312	27.41	201	77	1.3
316	27.73	197	77	1.28
320	28.05	168	77.5	1.32
324	28.37	208	78	1.3
328	28.69	186	81	1.31
332	29.01	202	81.5	1.34
336	29.32	178	82.5	1.33
340	29.64	214	82.5	1.32
344	29.96	202	80.5	1.31
348	30.28	224	82	1.35
352	30.6	182	83.5	1.35
356	30.92	221	84.5	1.35
360	31.24	218	84.5	1.34
364	31.55	221	86.5	1.34
368	31.87	200	87	1.34
372	32.19	217	87.5	1.31
376	32.51	195	88.5	1.32
380	32.83	217	90.5	1.35
384	33.15	205	93	1.34
388	33.47	220	91.5	1.33
392	33.79	244	93.5	1.32
396	34.1	225	97.5	1.34
400	34.42	267	100	1.33
404	34.74	234	97.5	1.35
408	35.06	264	97.5	1.34
412	35.38	231	99	1.37
416	35.7	293	100.5	1.42
420	36.02	240	101.5	1.4
424	36.33	238	104	1.31
428	36.65	246	100	1.38

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ^{10}Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ^{10}Be concentrations to calculate ^{10}Be fluxes. (*Continued*)

Depth (cm)	Age (kyr BP)	^{10}Be concentration (Be(M)) (10^6 atoms/g)	Magnetic susceptibility (SUS(M)) (10^{-8} m ³ /kg)	Dry bulk density (g/cm ³)
432	36.97	263	102	1.4
436	37.29	260	107	1.37
440	37.61	257	108	1.4
444	37.93	259	108.5	1.33
448	38.25	240	107	1.27
452	38.57	260	107.5	1.34
456	38.88	242	106.5	1.39
460	39.20	250	106	1.36
464	39.52	234	105	1.38
468	39.84	271	107	1.32
472	40.16	255	107	1.36
476	40.48	263	107.5	1.34
480	40.80	232	107	1.33
484	41.11	271	108.5	1.32
488	41.43	259	107.5	1.27
492	41.75	271	106.5	1.29
496	42.07	274	105.5	1.33
500	42.39	273	105.5	1.35
504	42.71	272	107	1.32
508	43.03	268	105.5	1.36
512	43.34	271	106.5	1.34
516	43.66	279	106	1.32
520	43.98	268	103.5	1.33
524	44.30	271	105	1.24
528	44.62	259	107	1.35
532	44.94	282	105.5	1.28
536	45.26	248	107	1.30
540	45.58	267	106	1.27
544	45.89	240	105.5	1.28
548	46.21	268	107	1.27
552	46.53	247	107	1.28
556	46.85	265	108.5	1.26
560	47.17	260	108	1.28
564	47.49	242	107.5	1.23
568	47.81	249	110	1.13
572	48.12	272	109	1.28
576	48.44	258	109.5	1.25
580	48.76	250	110.5	1.18
584	49.08	245	110.5	1.20
588	49.40	198	102.5	1.22
592	49.72	220	101.5	1.25
596	50.04	245	113.5	1.23
600	50.36	228	115.5	1.23
604	50.67	262	120.5	1.30
608	50.99	262	126	1.26
612	51.31	251	134	1.23
616	51.63	219	125	1.23

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ^{10}Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ^{10}Be concentrations to calculate ^{10}Be fluxes. (*Continued*)

Depth (cm)	Age (kyr BP)	^{10}Be concentration (Be(M)) (10^6 atoms/g)	Magnetic susceptibility (SUS(M)) (10^{-8} m ³ /kg)	Dry bulk density (g/cm ³)
620	51.95	251	126	1.23
624	52.27	264	131	1.25
628	52.59	258	135.5	1.19
632	52.90	241	138.5	1.18
636	53.22	266	137.5	1.21
640	53.54	239	136	1.23
644	53.86	272	140.5	1.19
648	54.18	228	139	1.17
652	54.50	249	132	1.18
656	54.82	233	131.5	1.19
660	55.13	237	124	1.26
664	55.45	230	124.5	1.17
668	55.77	252	115	1.20
672	56.09	229	111.5	1.21
676	56.41	237	113.5	1.25
680	56.73	209	113	1.25
684	57.05	238	108.5	1.25
688	57.37	237	107.5	1.21
692	57.68	225	108	1.19
696	58.00	215	108.5	1.18
700	58.32	242	98	1.22
704	58.64	225	88.5	1.28
708	58.96	227	83	1.30
712	59.28	193	71	1.24
716	59.60	198	62.5	1.29
720	59.91	199	62.5	1.33
724	60.23	186	61	1.37
728	60.55	169	59	1.33
732	60.87	192	57.5	1.27
736	61.19	184	54.5	1.30
740	61.51	179	55.5	1.29
744	61.83	178	53.5	1.34
748	62.14	166	54	1.36
752	62.46	181	52.5	1.35
756	62.78	187	53	1.39
760	63.10	189	52.5	1.37
764	63.42	194	54.5	1.39
768	63.74	190	51.5	1.38
772	64.06	184	52	1.34
776	64.38	189	50	1.36
780	64.69	166	48	1.32
784	65.01	169	46.5	1.32
788	65.33	170	44.5	1.31
792	65.65	156	47	1.27
796	65.97	147	42.5	1.28
800	66.29	162	43	1.33
804	66.61	149	43	1.33
808	66.92	166	43.5	1.34

Table 2 Measured quantities used for deducing geomagnetic and precipitation signals. Magnetic susceptibility, ^{10}Be concentration measurements, and time scale derivation are discussed in the Methods and Appendix sections. The age model and dry bulk density calculations were used to determine mass accumulation rate, which was then combined with ^{10}Be concentrations to calculate ^{10}Be fluxes. (*Continued*)

Depth (cm)	Age (kyr BP)	^{10}Be concentration (Be(M)) (10^6 atoms/g)	Magnetic susceptibility (SUS(M)) (10^{-8} m ³ /kg)	Dry bulk density (g/cm ³)
812	67.24	151	44	1.29
816	67.56	165	43.5	1.28
820	67.88	153	42	1.30
824	68.20	163	42.5	1.27
828	68.52	148	42.5	1.28
832	68.84	160	41.5	1.25
836	69.15	149	42.5	1.26
840	69.47	162	45.5	1.29
844	69.79	171	48	1.27
848	70.11	164	48	1.30
852	70.43	168	49	1.24
856	70.75	176	53	1.22
860	71.07	162	50	1.21
864	71.39	157	54	1.15
868	71.70	174	64	1.26
872	72.02	191	70.5	1.24
876	72.34	208	72.5	1.22
880	72.66	203	74	1.23
884	72.98	196	72	1.25
888	73.30	195	71.5	1.26
892	73.62	199	72	1.24
896	73.93	200	71	1.27
900	74.25	203	68	1.28
904	74.57	197	62	1.27
908	74.89	191	55.5	1.25
912	75.21	182	53	1.28
916	75.53	196	56	1.27
920	75.85	175	48	1.24
924	76.17	174	48	1.27
928	76.48	174	49	1.36
932	76.80	172	43.5	1.23
936	77.12	161	38.5	1.28
940	77.44	160	37.5	1.28
944	77.76	164	41	1.28
948	78.08	160	47	1.26
952	78.40	171	49	1.28
956	78.71	171	52	1.26
960	79.03	181	53	1.26
964	79.35	195	57	1.30
968	79.67	208	70.5	1.29

Age Calibration Using Radiocarbon Dates

^{14}C dating of the samples used for establishing the time scale in the section <12 kyr cal BP was performed at the NSF-Arizona AMS Facility (Table 3). Sample preparation took place at the Institute of Earth Environment in Xi'an. The ^{14}C ages were calibrated using OxCal v 3.10 (Bronk Ramsey 2001) and the IntCal04 calibration curve (Reimer et al. 2004).

Table 3 ^{14}C ages used for calibrating the younger part of the investigated loess sequence (depth <130 cm).

Material	Lab code	Depth (cm)	$\delta^{13}\text{C}$ (‰)	^{14}C age $\pm 2\sigma$ (yr BP)	Calibrated range (yr BP)	Mean calibrated (yr BP)
XLLQ1056	AA-44668	28	-25	2160 \pm 40	2040–2320	2180 \pm 140
XLLQ1087	AA-44696	53	-25	3920 \pm 65	4150–4530	4340 \pm 190
XLLQ1986B	AA-44695	75	-25	6340 \pm 70	7150–7430	7290 \pm 140

Age Calibration Using OSL Dates

Luminescence measurements (Table 4) of all quartz aliquots were undertaken using an automated Daybreak 2200 TL/OSL reader with a $^{90}\text{Sr}/^{90}\text{Y}$ beta source for irradiation. Optical stimulation with blue LEDs (470 ± 5 nm) was performed at 125°C , and OSL emission was detected through two 3-mm U-340 filters. For all samples, neutron activation analysis (NAA) was used to measure the uranium and thorium concentrations, and K content was determined by flame spectrum analysis. Dose rate was calculated according to (Aitken 1998).

Table 4 OSL ages of loess samples used for calibrating the older portion of the investigated loess sequence (deeper than 130 cm).

Lab code	Depth (cm)	U (ppm)	Th (ppm)	K (%)	Water content (%)	Dose (Gy/kyr)	Equi. dose (Gy)	Age (kyr BP)
IEE630	128	2.83 \pm 0.14	12.45 \pm 0.27	1.87	15 \pm 3	3.72 \pm 0.2	39.42 \pm 0.84	10.6 \pm 0.62
IEE631	138	2.73 \pm 0.15	11.87 \pm 0.27	1.71	15 \pm 3	3.5 \pm 0.19	43.55 \pm 0.98	12.44 \pm 0.74
IEE170	210	2.57 \pm 0.11	12.21 \pm 0.13	1.81	15 \pm 5	3.58 \pm 0.19	60.7 \pm 2.2	16.95 \pm 1.1
IEE395	400	2.51 \pm 0.13	12.04 \pm 0.26	1.61	15 \pm 3	3.34 \pm 0.19	131.6 \pm 5.4	39.4 \pm 2.7
IEE171	480	2.58 \pm 0.1	11.71 \pm 0.14	1.95	15 \pm 5	4 \pm 0.37	173.3 \pm 8.2	42.3 \pm 3.9
IEE396	520	2.15 \pm 0.11	11.14 \pm 0.25	1.69	15 \pm 5	3.21 \pm 0.17	142.6 \pm 6.5	44.48 \pm 3.16
IEE172	710	2.49 \pm 0.11	11.65 \pm 0.14	1.83	22 \pm 3	3.16 \pm 0.18	181.2 \pm 5.3	57.43 \pm 3.61
IEE397	780	2.46 \pm 0.11	11.33 \pm 0.25	1.53	15 \pm 5	3.15 \pm 0.18	210.8 \pm 6.4	66.95 \pm 4.3
IEE398	910	2.71 \pm 0.12	11.91 \pm 0.26	1.39	22 \pm 3	2.88 \pm 0.17	212.8 \pm 4.8	74.02 \pm 4.74
IEE173	920	2.75 \pm 0.1	12.48 \pm 0.14	1.86	25 \pm 3	3.1 \pm 0.15	231.7 \pm 7.8	74.7 \pm 4.31

The fine-grained (4–11 μm) quartz grains were extracted from the loess samples for luminescence dating. Equivalent dose values were determined by sensitivity-corrected multiple aliquot regenerative-dose protocol, which enabled us to recover equivalent dose values with high accuracy and precision in Chinese loess (Lu et al. 2007). Preheating conditions are 260°C (10 seconds) and 220°C (10 seconds) for natural/regeneration OSL (L_i) and test dose OSL (T_i) responses, respectively. The OSL signal from the first 5 seconds and subtracted that from the last 5 seconds of the decay curve has been applied to construct dose regenerative growth curve of the corrected OSL intensity (L_i/T_i) and to determine equivalent dose value.

Propagation of Errors

We report measurement uncertainties on ^{10}Be concentrations of 3%, and magnetic susceptibility uncertainties of ± 1 ($\times 10^{-8} \text{m}^3/\text{kg}$). Propagation of uncertainties in calculated results were made using standard statistical methods (Bevington and Robinson 1969; Jeter 2003). An estimate and uncertainty for SUS(D) of 25.1 ± 3.5 ($\times 10^{-8} \text{m}^3/\text{kg}$, 1σ) was obtained from linear least-square hyperbolic mixing model through the coercivity vs. susceptibility data shown in Figure 3. These uncertainties were graphically propagated through the linear least-squares model (2σ) uncertainty envelope surrounding the plot of Be(M) vs. SUS(M) (Figure 4b), to yield a value and uncertainty for the ^{10}Be dust

endmember Be(D) ($136.4 \pm 6.25 [\times 10^6 \text{ atoms/g}, 1 \sigma]$). As by convention, for the regression calculations of SUS(M) vs. Be(M), all analytical errors were assigned to the dependent-variable (Be(M)) assuming the errors were not covariant. Uncertainties in SUS(D) and Be(D) were quadratically combined with measurement errors on SUS(M) and Be(M) to determine uncertainties for SUS(P) and Be(P,GM). Average propagated (1σ) uncertainties on SUS(P) are 3.64 (SI Units) and 7.1% for Be(P,GM) values. A linear model dependence between SUS(P) and Be(P,GM) is assumed (Figure 4d). This linear model is subtracted from Be(P,GM) to yield values for Be(GM) and Be(P). We assert the linear trend in this plot is due to precipitation amount, but that much of the observed scatter is not noise but rather is due to the second independent variable, i.e. geomagnetic field variations. As it is not possible to independently determine how much scatter is due to noise vs. field variations, the errors in the model results Be(GM) and Be(P) are assumed to be equivalent to those determined for Be(P,GM). Be(GM) concentrations were converted to fluxes using the linear age model-based accumulation rates (see Figure 7), with errors on $\text{Be(GM)}_{\text{flux}}$ expanded to account for standard error in the slope of the age vs. depth model relationship (2.3%). Normalization of $\text{Be(GM)}_{\text{flux}}$ to modern ^{10}Be production rate is assumed to not increase fractional uncertainty on normalized $\text{Be(GM)}_{\text{flux}}$. Errors on normalized $\text{Be(GM)}_{\text{flux}}$ were propagated through the global mean ^{10}Be power law production rate equation of Masarik and Beer (1999). Uncertainties of 7% (1σ) in ^{10}Be production rates (Masarik and Beer 1999) were used in combination with calculated model errors on normalized $\text{Be(GM)}_{\text{flux}}$. Median uncertainties on calculated M/Mo (Figure 5b) is 14% with a mean uncertainty of 21%. The large difference between mean and median uncertainties is due to a relatively small number (~ 15 of 116) of model M/Mo values from the LGM and MIS4 with very large uncertainties (between 30–100%, 1σ). The reason for the large uncertainties in these few samples is due to low precipitation during the LGM and MIS4, resulting in low Be(P,GM) fluxes, which after subtraction of Be(P) resulted in even lower calculated Be(GM) fluxes with proportionally high uncertainties.

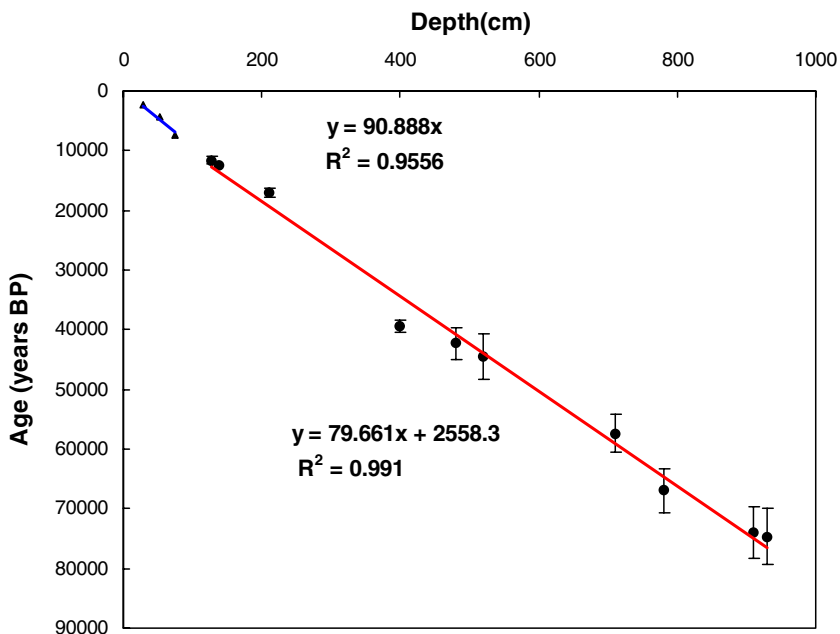


Figure 7 Plot of all data used for age calibration of the loess sequence. ^{14}C and OSL data are shown together with their corresponding regression lines (<10 and >10 kyr BP, respectively). The ^{14}C ages were used for calibrating the younger part (depth <130 cm). The OSL ages were used for calibrating the older part (depth >130 cm).

Uncertainties for paleorainfall amount (Figure 6) used calculated uncertainties for $\text{Be(P)}_{\text{flux}}$ (average 7.03%, 1 σ), which were calculated using the same method as for $\text{Be(GM)}_{\text{flux}}$ uncertainties. The uncertainty in tropospheric $^{10}\text{Be}/^7\text{Be}$ is not well known. As such, we arbitrarily assigned a 5% (1 σ) uncertainty to this ratio. These uncertainties were combined with uncertainties in slope and intercept of the plot of ^7Be vs. annual rainfall for the 3 modern sites evaluated to yield average uncertainties for model paleoprecipitation estimates of 19.5% (1 σ).