

SMALL SAMPLE DATING IN CHINA

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ABSTRACT. The Xi'an Laboratory of Loess and Quaternary Geology has developed a small sample ¹⁴C dating facility consisting of a Wallac 1220 Quantulus™ liquid scintillation spectrometer, and a miniature benzene synthesis line based on the synthesis procedures used at the Australian National University (ANU). This line can produce *ca.* 0.3-ml benzene samples, which are then measured for ¹⁴C activity using 0.3-ml Teflon vials developed by Wallac Oy. The counting performance of the Quantulus™ spectrometer using 0.3-ml vials has been evaluated, and a potential age limit of *ca.* 45,000 BP has been obtained for samples containing up to 250 mg carbon. This dating facility fills the gap between large sample (2.4–6 g carbon) and microsample (<1 mg carbon) handling to form a ¹⁴C dating method sequence.

INTRODUCTION

The establishment of a relatively accurate ¹⁴C chronology for many of the well-defined loess-paleosol sequences in northern China is an important component of any investigation of global climate change over the past 30,000 yr. The carbon content of these sediments ranges between 1 and 2%. Intensive agriculture has introduced a relatively high concentration of young water-soluble organic compounds, which are then adsorbed onto the clay component of the paleosols. Thus, it is advisable to subject the organic component of the paleosols to extensive pretreatment and chemical fractionation to obtain reliable ¹⁴C ages (Head, Zhou and Zhou 1989; Zhou, Zhou and Head 1990; Zhou *et al.* 1992). The final components for dating are usually too small for conventional liquid scintillation counting (LSC). Because of the scarcity of ¹⁴C accelerator mass spectrometry (AMS) facilities in China, and the expense of obtaining AMS dates from overseas, the concept of upgrading an existing radiometric ¹⁴C dating facility to handle carbon sample sizes between 25 and 250 mg is very attractive.

Polach *et al.* (1988) have documented LSC performance on very small samples (25–250 mg carbon) for a variety of vial types ranging from 0.3 ml to 3.0 ml using carefully diluted aliquots of a 200% modern standard. They found that, for samples containing <250 mg carbon, minivials produced more precise results than larger vials. For 0.3-ml Teflon vials developed by Wallac Oy (Kaiholä, Kojola and Heinonen 1992), a figure of merit (FM) of 11.9 was obtained by the ANU Radiocarbon Dating Research Unit, producing a theoretical dating limit of 43,230 BP for a 3000-min counting period (Polach *et al.* 1988). Further studies by Hogg (1992) have shown that the accuracy of results obtained using this type of vial falls within acceptable limits for samples containing 100–250 mg carbon. This method thus enables us to determine the ¹⁴C activity of samples containing 250 mg carbon with acceptable precision, even though the counting is slow.

METHODS

We decided that using a full-scale benzene synthesis apparatus for samples of <250 mg carbon decreased efficiency of preparation and yield of benzene. Thus, we designed a miniature system based on the procedures developed at the ANU Radiocarbon Dating Research Unit (Gupta and Polach,

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1985). The miniature benzene synthesis apparatus also enabled us to keep possible memory effects to a minimum.

Because of the difficulty in transferring the sample benzene from the collection trap to a storage vial without incurring significant losses, actual yield measured was as low as 75%, which is representative of a total yield of *ca.* 95%. Sample benzene was added to the 0.3-ml Teflon vials together with butyl PBD scintillant at a relative concentration of 15 g liter⁻¹ (0.0045 g). It was found that the 0.3-ml vial as supplied (Kaihola, Kojola and Heinonen 1992) tended to lose up to 1 mg benzene by evaporation over a period of 7–8 days. The use of 2 Tuffbond silicone/Teflon seals instead of 1 caused benzene loss to be as low as 0.3 mg per week. The vials were positioned for maximum quantum yield (Kaihola, Kojola and Heinonen 1992) by lining up a mark on each vial with a mark on the Quantulus sample lift mechanism. The value of the modern standard obtained as a mean of the two vials is 2.9853 cpm, and background is 0.07 cpm, giving an FM of 11.28. With this value, the theoretical age limit for the vials is 43,270 BP, using the equations listed by Gupta and Polach (1985), and by Kaihola, Kojola and Heinonen (1992). These values were obtained with a counting efficiency of 88%. This theoretical age limit compares favorably with that obtained by Polach *et al.* (1988).

CROSS-CHECK SAMPLES

To test the accuracy of the technique, a series of previously dated samples was collected from the ANU Laboratory and the Xi'an Laboratory archives. Table 1 shows the results. All samples were measured for 3000 min, and are expressed as conventional ages (Stuiver and Polach 1977). To test whether the small sample ages are significantly different from the original ages obtained for the samples, the T statistic for one degree of freedom was calculated (Gupta and Polach 1985). Because T has a chi-squared distribution, we can then use the relevant tables for one degree of freedom at the 95% confidence level. T needs to be greater than 3.841 for the results to be significantly different.

TABLE 1. Comparison Between Small Sample Results and Dates Obtained Previously for the Same Sample

Lab no.	Age 1 (BP) (0.3 ml)	Lab no.	Age 2 (BP)	T value*
XLLQ-58	39,650 ± 2210	XLLQ-58	30,930 ± 320	53.9
XLLQ-106	14,110 ± 530	ANU-6393	14,000 ± 1700	.004
XLLQ-336	5,330 ± 180	XLLQ-336	5100 ± 130	1.073
XLLQ-9	2,410 ± 130	XLLQ-9	2300 ± 50	.624
XLLQ-120	2,340 ± 130	XLLQ-120	2180 ± 90	1.024
XLLQ-14	2,470 ± 130	XLLQ-14	2390 ± 80	.275
XLLQ-13	2,460 ± 130	XLLQ-13	2620 ± 85	1.061
XLLQ-483	29,660 ± 680	ANU-5827	29,220 ± 640	.222
XLLQ-484	18,390 ± 450	ANU-4655	18,720 ± 220	.178
XLLQ-485	10,560 ± 160	ANU-6613	10,800 ± 220	1.528
XLLQ-486	6,255 ± 150	ANU-7099	6150 ± 80	.381
XLLQ-487	4,450 ± 140	ANU-7852	4360 ± 80	.312
XLLQ-489	3,550 ± 140	ANU-5085	3130 ± 80	6.785
XLLQ-490	1,650 ± 130	ANU-6088	1550 ± 70	.459
XLLQ-491	220 ± 140	ANU-5425	330 ± 80	.465

*T statistic (Gupta and Polach 1985)

The small sample age for XLLQ-58 is significantly older than the previous result obtained, and the only explanation for this is the fact that further pretreatment was carried out before the sample was

rerun (this was a sample of wood holocellulose). The small sample age would be considered to be the most reliable. The only other pair of results showing a significant difference is XLLQ-489 and ANU 5085. We have no explanation for the perceived difference in age other than the fact that this was a shell sample and there may be a small amount of heterogeneity in the sample. There may also have been exchange between shell carbonate and atmospheric CO₂ since the original shell material was dated *ca.* 5 yr earlier. For the remaining pairs, the T statistic is much less than the chi-squared value, indicating that the results are extremely close.

CONCLUSION

This exercise has indicated that the Xi'an Laboratory small sample facility for ¹⁴C dating using liquid scintillation spectrometry can produce ¹⁴C ages with precision comparable to other counting techniques.

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