COMPARISON OF MANUAL AND AUTOMATED PRETREATMENT METHODS FOR AMS RADIOCARBON DATING OF PLANT FOSSILS

LEE-ANN BRADLEY

U. S. Geological Survey, Branch of Earthquake and Landslide Hazards, MS 966, Denver, Colorado 80225 USA

and

THOMAS W. STAFFORD, JR.

INSTAAR, The University of Colorado, Boulder, Colorado 80303 USA, and U.S. Geological Survey, Branch of Isotope Geology, MS 963, Denver, Colorado 80225 USA

ABSTRACT. A new automated pretreatment system for the preparation of materials submitted for accelerator mass spectrometry (AMS) analysis is less time-consuming and results in a higher sample yield. The new procedure was tested using two groups of plant fossils: one group was pretreated using the traditional method, and the second, using the automated pretreatment apparatus. We compared the time it took to complete the procedure and the amount of sample material remaining. The automated pretreatment apparatus proved to be more than three times faster and, in most cases, produced a higher yield. We also observed a darker discoloration of the KOH solutions, indicating that the automated system is more thorough in removing humates from the specimen compared to the manual method.

INTRODUCTION

Chemical pretreatment of routine samples for AMS ¹⁴C analysis is labor intensive, requiring as much as one-third of the time required for a complete sample analysis. Standard manual pretreatment methods (Gagnon and Jones 1993) are slow because a technician can process only 1 or 2 samples at a time. These methods also commonly result in a large percentage of sample loss. To speed pretreatment time and reduce sample loss, we designed and constructed an automated system controlled by a personal computer (Fig. 1). The system runs multiple samples simultaneously and yields a higher percentage of sample material. Here we compare pretreatment results of 11 samples using standard manual methods and our automated system. On average, the automated system reduced pretreatment time by 70% and sample loss by 4%.

METHODS

Eleven plant fossils of different ages were selected to demonstrate the effects of manual and automated pretreatment methods on the types of materials commonly submitted for AMS analysis (Table 1). All specimens were cleaned of foreign matter, and split into two samples weighing 0.01– 0.10 g. One sample was pretreated using the manual method (denoted by a suffix M on the sample number), and the other sample was pretreated using the automated system (suffix A). The samples were weighed before and after the pretreatment procedures to determine the amount of sample loss. Pretreatment times do not include drying time. All samples were processed through the same sequence of reagents (Table 2). Deionized water and distilled HCl were used for reagent mixing and to decrease contamination; all glassware was heated to 550°C for 1 h.

Automated Pretreatment Method

Each sample was placed in a vertical borosilicate glass column (10 mm ID \times 100 mm long) with a Teflon filter end-cap fixed to each end (Fig. 2). The filter end-caps were fitted with 25- μ polyethylene disposable frits that were replaced with each sample change. Teflon tubing leading from pressurized reagent containers were attached to the top filter end-cap to allow reagents to flow through

			Amt. C			Fraction		
		Pretreatment	dated	Sample		modern	FM	¹⁴ C Age
Lab no.	Sample no.	method	(mg)	type	Location	(FM)	SD	(yr BP)
	HB415-416A	Automated	:	Moss	Harberton, Argentina	-	ł	Not dated
ł	HB415-416M	Manual	ł	Moss	Harberton, Argentina	1	ł	Not dated
:	HB561-562A	Automated	1	Moss	Harberton, Argentina	ł	;	Not dated
ł	HB561-562M	Manual	ł	Moss	Harberton, Argentina	ł	ł	Not dated
CAMS-6343	HB310-311A	Automated	1.24	Moss	Harberton, Argentina	0.6372	0.004	3620 ± 60
CAMS-6342	HB310-311M	Manual	1.17	Moss	Harberton, Argentina	0.6422	0.005	3560 ± 60
CAMS-6340	CHEM-409A	Automated	1.28	Wood	Hiscock site, New York	0.3159	0.0024	9260 ± 70
CAMS-6341	CHEM-409M	Manual	0.74	Wood	Hiscock site, New York	0.32	0.003	9150 ± 80
CAMS-6345	NSRL-747A	Automated	1.23	Twig*	Rockfall Cave, New Mexico	0.716	0.009	2680 ± 100
CAMS-7556	NSRL-747A	Automated	1.20	Twig*	Rockfall Cave, New Mexico	0.7246	0.005	2590 ± 60
CAMS-7199	NSRL-747M	Manual	1.50	Twig*	Rockfall Cave, New Mexico	0.7279	0.0077	2550±90
CAMS-16812	NSRL-531A	Automated	1.22	Stems	Rampart Cave, Arizona	0.2576	0.0018	$10,900 \pm 60$
CAMS-16813	NSRL-531M	Manual	0.88	Stems	Rampart Cave, Arizona	0.2566	0.0016	$10,930 \pm 60$
CAMS-6369	NSRL-748A	Automated	1.11	Charcoal	Hermit's Cave, New Mexico	0.6631	0.007	3300 ± 90
CAMS-6383	NSRL-748A	Automated	1.11	Charcoal	Hermit's Cave, New Mexico	0.6572	0.005	3370 ± 70
CAMS-6365	NSRL-748M	Manual	1.41	Charcoal	Hermit's Cave, New Mexico	0.6636	0.007	3290 ± 90
CAMS-6364	NSRL-749A	Automated	1.66	Charcoal	Hermit's Cave, New Mexico	0.6454	0.005	3520±70
CAMS-6366	NSRL-749M	Manual	1.54	Charcoal	Hermit's Cave, New Mexico	0.6482	0.0052	3480 ± 70
CAMS-6368	NSRL-750A	Automated	0.98	Charcoal	Hermit's Cave, New Mexico	0.4988	0.004	5590 ± 70
CAMS-6381	NSRL-750M	Manual	1.23	Charcoal	Hermit's Cave, New Mexico	0.4972	0.004	5610 ± 70
CAMS-5897	NSRL-805A	Automated	1.30	Wood	Fairbanks Glacier, Alaska	Not reported	Not reported	5610 ± 140
CAMS-5898	NSRL-805A	Automated	1.30	Wood	Fairbanks Glacier, Alaska	Not reported	Not reported	5400 ± 130
CAMS-5899	NSRL-805A	Automated	1.72	Wood	Fairbanks Glacier, Alaska	Not reported	Not reported	5320 ± 130
CAMS-6344	NSRL-805M	Manual	1.06	Wood	Fairbanks Glacier, Alaska	0.5115	0.007	5380 ± 120
ł	NSRL-746A	Automated	ł	Wood	Rockfall Cave, New Mexico	ł	1	Not dated
CAMS-7555	NSRL-746M	Manual	1.2	Wood	Rockfall Cave, New Mexico	0.2131	0.0021	$12,420 \pm 80$
	•••							

*Archaeological sample; probably a yucca spine tied into a knot

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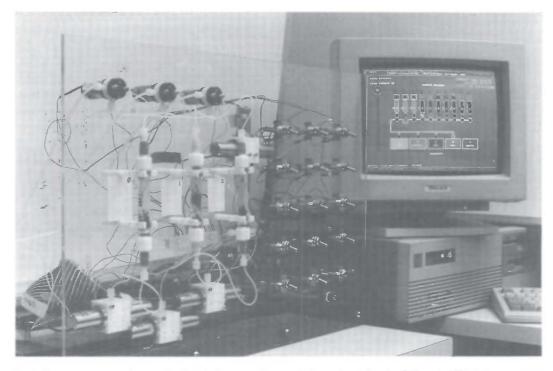


Fig. 1. Computer-automated system for chemically pretreating wood, charcoal and plant fossils for AMS 14 C dating. An IBM-compatible personal computer controls the movement of four solvents over samples contained in the glass columns.

the glass column. Each column drained through Teflon tubing attached to the bottom filter end-cap to a waste receptacle. Our system handles up to four different reagents (Table 2) and processes up to 3 sample columns simultaneously, but it could be modified to run as many as 5–10 samples. Each sample was weighed in the glass column including the filter end-caps and remained in its column throughout the pretreatment process.

We used an IBM-compatible personal computer to control the flow of reagents through the automated pretreatment apparatus. Specially designed computer software activates solenoid switches that control the flow of the reagents for time intervals specified by the program (Table 2). We used the same sequence for all samples except NSRL-750A (Table 1) because the discoloration of the KOH solution after its third treatment led us to infer that this sample needed additional pretreatment. Because discoloration of the KOH solution usually signifies contamination with humates, we ran sample NSRL-750A through the entire program sequence again (Table 2).

After drying, the final sample weight was calculated by total yield, and total recoverable yield. The total yield was calculated by subtracting the weight of the sample in the column from the initial weight of the column before pretreatment. The total recoverable yield was the weight of the dried sample after removal from the column. We noted only minor differences in these two values (see below and Fig. 3).

Manual Pretreatment Method

We pretreated the manual samples in the original sample vials, using the same sequence of reagents used in the automated pretreatment method (Table 2). Reagents were added and removed using

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TABLE 2. Program sequence for one complete sample run with the automated pretreatment system. Our solvent reagents are made from deionized H_2O and a stock solution of distilled, constant-boiling HCl

Sample			Time	
Step	Column		(sec)	
1	Fill	#1	H ₂ O pH6.5	50
2	Fill	#2	H ₂ O pH3	90
3	Purge			30
4	Fill	#3	0.5N HCl	60
5	Hold	#3	0.5N HCl	300
6	Purge			30
7	Fill	#1	$H_2O pH 6.5$	90
8	Purge			30
9	Fill	#4	0.5% KOH	60
10	Hold	#4	0.5% KOH	600
11	Purge			30
12	Fill	#1	H ₂ O pH 6.5	120
13	Purge			30
14	Fill	#2	H ₂ O pH 3	90
15	Purge			30
16	Fill	#1	H ₂ O pH 6.5	90
17	Purge			30
18	Fill	#4	0.5% KOH	60
19	Hold	#4	0.5% KOH	600
20	Purge			30
21	Fill	#1	H ₂ O pH 6.5	120
22	Purge			30
23	Fill	#2	H ₂ O pH 3	90
24	Purge			30
25	Fill	#1	H ₂ O pH 6.5	90
26	Purge			30
27	Fill	#4	0.5% KOH	60
28	Hold	#4	0.5% KOH	600
29	Purge			30
30	Fill	#1	$H_2O pH 6.5$	150
31	Purge			30
32	Fill	#2	H ₂ O pH 3	200
33	Purge			120
34	End			

pipettes with the tips modified to ca. 100μ in diameter to minimize the amount of sample loss. A single pipette was used for each reagent to reduce contamination of the reagents. The time for each step was kept as close as possible to the times of the automated sequence including the time needed to add and remove reagents. After drying, the total yield was calculated by subtracting the initial weight of the sample vial from the final weight.

RESULTS

Time

All 11 samples processed using the automated system were completed in a total of 4.7 h; processing with the manual method required 15.7 h, or 3.4 times longer than the automated method (Fig. 4). The main reasons for the time savings are: 1) multiple samples can be run simultaneously in the automated method; and 2) adding and draining the reagents takes just a few seconds with the automated system, whereas adding and removing reagents manually using pipettes is very timeconsuming.

Sample Yield

In 9 of 11 specimens treated, the percent of total yield was higher for the automated system than the manual method. However, after the samples were removed from the glass columns, the total recoverable yield using the automated system was greater only in 6 of 11 specimens (Fig. 3). Apparently, fibrous or minute amounts of sample material were trapped in the frits at the end of the columns and could not be removed. The differences in yield in the automated system ranged from a

minimum of 0.5% for HB561-562A to a maximum of 7% for NSRL-531A (Fig. 3). For one sample (NSRL-746), the wood completely dissolved during automated pretreatment, whereas the yield during manual pretreatment was 45.54%. The reason for higher manual yield was that the solvents were added very slowly to lessen physical and chemical deterioration of the very friable wood. Although slow, gentle pretreatment yielded wood for dating, it is very possible that contaminants (*e.g.*, humates) could remain. Because no test exists presently to determine when contamination removal is complete and sample dissolution begins, we favor pretreating all samples to completion. If the material disintegrates totally, the sample is labeled "not datable due to dissolution during pretreatment".

Quality of Pretreatment

In the first steps of the manual pretreatment process, some samples (especially mosses) float, thus reducing surface contact with the reagents. With the manual method, these buoyant samples were submerged only after special attention from the technician. The automated samples do not suffer

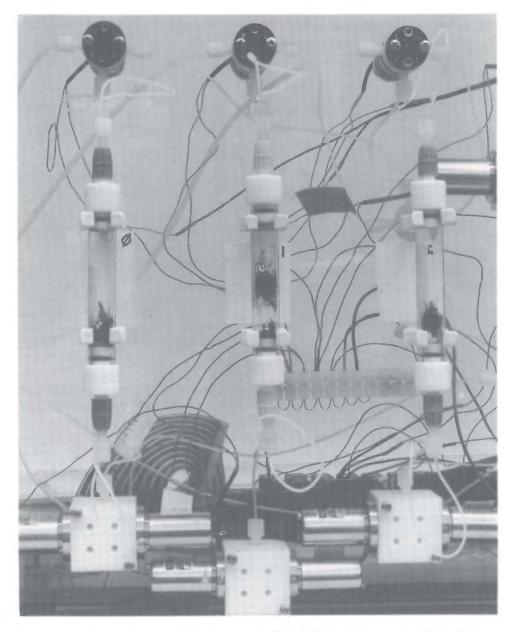


Fig. 2. Close-up of glass columns containing plant macrofossils ready for chemical pretreatment. Solenoid valves control the flow of the reagents upwards into the columns during pretreatment, and purge downward by using nitrogen gas pressure.

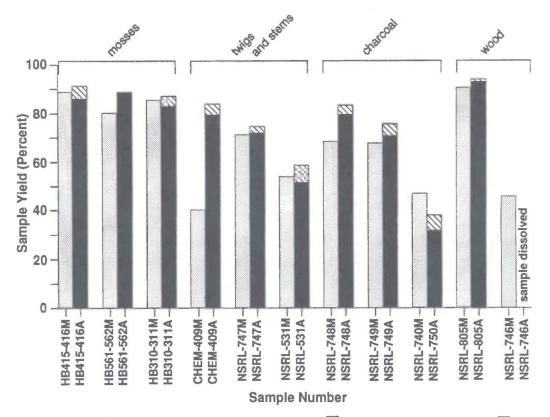


Fig. 3. Sample yields for manual and automated pretreatment methods. \square = total yield from manual method; \square = total recoverable yield from automated method; \square = percent of sample not lost during pretreatment but not recoverable from the glass column; \square + \blacksquare = total yield from automated method. Note higher percentage of total yield in automated samples. There was no significant difference between total yield and recoverable yield for most materials.

from this problem because they are sealed in glass columns and thus are completely submerged during the entire pretreatment process. The reagents used with the automated system showed more discoloration than the manually treated samples; in particular, the degree of brown discoloration from humates leached from the sample during the KOH steps is an indication of the degree of surface contact between the sample and the reagents. Thus, the automated process appears to be more effective in removing humic acid contaminants from the sample. There was also a much greater chance for human error with the manual method. A distracted technician can easily lose track of the sequence and add the wrong reagent. Contamination of the reagents is more likely because the containers are unsealed and pipettes can be inadvertently switched between reagents. With the automated system, the sample is only minimally disturbed because the sample is in contact only with the reagents flowing in and draining from the column.

CONCLUSIONS AND RECOMMENDATIONS

Because labor costs are a large part of the cost of pretreatment, time savings reduce the cost of pretreatment. The automated method was more than three times faster than the manual method, making the automated method more cost-effective.

													THORE-740M
	I LIDA1E.	-416M	HB561-4	562M HB31	0-311M C	HEM-409M	NSRL-747M	NSRL-531M	NSRL-748M	NSRL-749M	NSRL-750M	NSRL-805M	NODI -74614
1	r										Manua		nent Method me = 15.7 h
	NSRL-74	46A NS	RL-750A	NSRL-805A	HB310-31	A							
	NSRL-74	48A	NSRL	-750A	HB561-56		ated Pretrea ime = 4.7 h		hod				
	NSRL-53	31A CH	EM409A	NSRL-747A	HB415-41								
					Batch 4								

Fig. 4. Cumulative time required to complete pretreatments on 11 specimens using the automated system and the manual method. Using the automated system, three samples were processed simultaneously, permitting all samples to be run in 4 batches. NSRL-750A was run twice because discolored KOH reagents indicated incomplete pretreatment. The total time to process all 11 samples using the automated system was 4.7 h, compared to 15.7 h using the manual method.

The total recoverable yield is dependent on the method used and the type of sample material. In most cases, the automated method produced a higher yield, probably because less material is lost draining the reagents through filters than by siphoning off reagents manually. Extremely fragile materials that require special care, such as very soft wood or fine-grained charcoal, showed a higher yield by manual pretreatment. For most materials (plants, mosses, twigs and stems, most charcoal and fibrous wood) the yield is higher using the automated method. A necessary modification would be to use finer porosity frits in the filter end-caps. Smaller pore sizes would be smoother and occlude less of the fibrous sample. The second modification would be to decrease the gas pressure that moves the solutions in and out of the system. By decreasing the flow rate, the automated system would more closely mimic the manual pretreatment of delicate samples.

The darker discoloration of the KOH solutions we observed during the automated pretreatments indicates that this method is more thorough in removing humates from the specimens. Although there was no statistically significant difference in the paired ¹⁴C ages, 6 out of 7 pairs of samples gave older ages for the automated samples, thus supporting our observation that the automated pre-treatment process was slightly better (Table 1).

The next generation instrument is under development and will provide more control over pretreatment conditions. The improvements will include: 1) ability to vary linearly the KOH concentrations from 0.01% to 0.5%; 2) change flow rates to accommodate friable samples, 3) provisions for treating samples from room temperature to 80° C; 4) improvement of software to enable rerunning of specific, severely contaminated samples; and 5) rewriting the software for Macintosh systems.

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