



ELPAT Program Report Background and Current Status

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Paul C. Schlecht and Jensen H. Groff, Column Editors

Introduction

The Environmental Lead Proficiency Analytical Testing (ELPAT) Program is administered by the American Industrial Hygiene Association (AIHA), in cooperation with researchers at the Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH), and the U.S. Environmental Protection Agency (EPA), Office of Pollution Prevention and Toxics, to evaluate and improve the performance of laboratories conducting analyses associated with lead abatement.^(1,2) Proficiency test samples are prepared by an AIHA contractor, Research Triangle Institute (RTI), using real-world paint chips, dusts, and soils. Quarterly samples are sent to participating laboratories by RTI and the performance of the laboratories is evaluated at NIOSH, with sufficient time for laboratories to obtain repeat samples and to correct analytical problems before the next round of samples is sent.

The ELPAT Program is open to all interested laboratories, including laboratories outside the United States, laboratories seeking accreditation by various private or state laboratory accreditation systems, and laboratories that do not intend to seek laboratory accreditation. The ELPAT Program is part of an EPA Program, the National Lead Laboratory Accreditation Program (NLLAP), to recognize private and state laboratory accreditation systems.⁽³⁾ NLLAP requirements include successful participation in the ELPAT Program for EPA recognition of accreditation. Two organizations, the American Association for Laboratory Accreditation (A2LA)⁽⁴⁾ and AIHA,⁽⁵⁾ are recognized as accrediting organizations under NLLAP and have environmental lead laboratory accreditation systems in place. Each of these accreditation systems requires participation in ELPAT for environmental lead analysis of paint chips, dusts, and soils. Information on specific A2LA or AIHA laboratory accreditation requirements can be obtained from

A2LA and AIHA at the addresses listed at the end of this column.

ELPAT Performance Evaluation

The evaluation of the individual laboratories in the ELPAT Program is based upon consensus values from reference laboratories and is modeled after the evaluation procedures currently used in an industrial hygiene proficiency testing program, the Proficiency Analytical Testing (PAT) Program. Reference laboratories are preselected to provide the performance limits for each sample. These laboratories must meet the following criteria: the laboratory was proficient in the previous PAT round for a wide variety of industrial hygiene laboratory air analyses, including airborne lead, and industrial hygiene laboratory operations must be accredited by AIHA.⁽⁶⁾ Once a significant history of ELPAT performance is available, reference laboratories will include all participating laboratories in the ELPAT Program that have previously demonstrated proficient performance in analyzing all matrices of the ELPAT Program. Eventually, a requirement will be added that ELPAT reference laboratories must be accredited by an EPA NLLAP-recognized accrediting organization.

After data from reference laboratories are collected and extreme reference laboratory data have been statistically treated, the mean ± 3 standard deviations of the treated reference laboratory data become the acceptable performance range. Laboratory results are acceptable if they fall within the performance limits. Results falling outside the performance limits are designated as outliers. This is the same criterion used by NIOSH to establish acceptable and outlier performance of industrial hygiene laboratories in the PAT Program.⁽⁶⁾

Laboratories are rated based upon performance in the ELPAT Program over the last year (i.e., four rounds) for each lead matrix—paint chips, soil, and dust wipes. The laboratory is proficient for the lead matrix if the following occurs:

1. all four results have been reported and all are designated as acceptable for the last two consecutive rounds; or
2. three-fourths or more of the results reported in the last four consecutive rounds are designated as acceptable.

However, if a laboratory does not report values for the lead matrix on the round being evaluated, the laboratory is not rated.

Initial criteria for proficient performance are similar to the procedure used in the PAT Program.⁽⁶⁾ However, the ELPAT statistical protocol and related computer programs have been designed to permit future change to harmonize these proficiency test requirements with internationally harmonized proficiency test protocols. An international protocol for consensus values from reference laboratories using z-scores is being developed by the International Organization for Standardization, the Association of Official Analytical Chemists International, and the International Union of Pure and Applied Chemists.⁽⁷⁾

ELPAT Round 8, August 1994

Paint samples for round 8 were prepared from paint chips collected from a variety of sites, including commercial lead abatement, building renovation, and demolition sites. The chips were ground to a maximum particle size of 120 μm .

Soil samples came from driplines around older houses in North Carolina and from industrial sites in Colorado and Louisiana. Soil samples were dried and then sterilized by heating the soil to 325°F for a minimum of 2 hours, and finally sieved to a maximum particle size of 150 μm .

Round 8 dust wipes were prepared by gravimetrically loading Whatman 40 filter paper with sterilized (gamma-irradiated) household and postabatement dust, sieved to a maximum particle size of 150 μm . The loaded filters were moistened with 0.5 ml of 3 percent hydrogen peroxide solution. The blank wipe was prepared from a Whatman filter moistened with the same hydrogen peroxide solution. Whatman filters are easier to digest

TABLE 1. ELPAT Program Summary Statistics of Reference Laboratories for Round 008

Sample Type	Sample	N	Mean	Minimum	Maximum	STD	RSD (%)	Acceptable Range
Paint chips (%)	1	31	0.322	0.294	0.3444	0.018	5.5	0.2685–0.3756
	2	31	3.5563	3.2137	3.823	0.186	5.2	0.9992–4.1133
	3	31	0.0784	0.0698	0.0888	0.006	8.2	0.0591–0.0977
	4	31	0.7435	0.67	0.875	0.058	7.8	0.5692–0.9177
Soil (mg/kg)	1	31	1620.1	1480	1729.4	78.6	4.9	1384.1–1856.1
	2	31	44.2	31.2	61.3	8.58	19.4	18.4–70
	3	31	251.1	211	286.2	25.5	10.1	174.6–327.6
	4	31	791.9	724.7	847.6	43.4	5.5	661.6–922.1
Dust wipes (μg)	1	31	469.1	409	521.2	35.8	7.6	361.8–576.4
	2	31	104.6	86.2	119.8	10.2	9.7	74–135.2
	3	31	1302.1	1112.7	1454.5	107	8.2	980–1624.1
	4	31	241.9	213	278	20.5	8.5	180.3–303.5

than other wipe media (e.g., baby wipes, hand wipes) used by many laboratories. In the future, the wipe medium may be changed from the Whatman filter to a commercially available wipe that more closely represents field sample media, if a single sample medium is recommended by various lead methods.

A total of 324 laboratories were enrolled for round 8 of the ELPAT Program, with 308 (95%) laboratories submitting results either by paper or by the ELPAT automated data entry system. Table 1 lists summary statistics of reference laboratories for each matrix and sample number. Agreement among reference laboratories is demonstrated by relative standard deviations ranging from 5.2 to 8.2 percent for paint chips, 4.9 to 19.4 percent for soils, and 7.6 to 9.7 percent for dust wipes. This is similar to the agreement among reference laboratories on previous ELPAT rounds for each matrix.

Table 2 shows the number of all participating laboratory analyses that were identified as outliers. The percentage of outliers for all analyses was under 15.4 percent (7.7 to 15.4% for paint chips, 5.8 to 9.7% for soils, and 6.1 to 8.2% for dust wipes). This is also similar to the frequency of outliers reported on the earlier rounds of ELPAT for each matrix.

Sample digestion techniques are grouped into the most common: hotplate, microwave, and all other techniques reported by participants. Hotplate digestion categories are: NIOSH 7082/7105 (a nitric acid/hydrogen peroxide digestion method modified from NIOSH Method 7082⁽⁸⁾), EPA SW846–3050A⁽⁹⁾

(an EPA nitric acid/hydrogen peroxide method), and other hotplate techniques.

Microwave digestion categories are: EPA SW846–3051⁽¹⁰⁾ (a nitric acid digestion method), EPA AREAL⁽¹¹⁾ [a nitric/hydrochloric acid digestion method from AREAL (RTP-MRDD-037) standard operating procedure], and other microwave techniques. The “all other” category includes nonmicrowave and non-hotplate techniques such as X-ray fluorescence sample preparation, leaching techniques, muffle furnace, and Parr bomb.

Instrumental methods are categorized into flame atomic absorption (FAA), graphite furnace atomic absorption (GFAA), inductively coupled plasma—atomic emission spectroscopy (ICP-

AES), laboratory X-ray fluorescence (lab XRF), and “others,” which includes ICP-mass spectroscopy (ICP-MS).

Table 3 shows a summary of failures (outliers) for the three lead matrices by digestion technique and analytical method used by participating laboratories. A series of Fischer’s exact tests (non-parametric tests) were used to compare the various combinations of digestion techniques (hotplate and microwave) and analytical methods (FAA, GFAA, ICP-AES) for statistically significant differences in the ability of the digestion techniques/analytical method combinations to meet ELPAT performance limits.⁽¹²⁾ To detect differences in performance, a criterion was then used where participat-

TABLE 2. ELPAT Round Program Summary of Performance—All Laboratories Participated for Round 008

Sample Type	Sample No.	No. of Labs Rated	Acceptable Labs	Low Outlier	High Outlier
Paint chips (%)	1	299	273	17	9
	2	299	253	29	17
	3	299	273	11	15
	4	299	276	11	12
Soil (mg/kg)	1	259	234	18	7
	2	259	236	6	17
	3	259	244	5	10
	4	259	237	16	6
Dust wipes (μg)	1	279	256	14	9
	2	279	260	9	10
	3	279	262	10	7
	4	279	262	14	3

TABLE 3. ELPAT Program Labs Performance Summary for Round 008

Method	Sample	Preparation	Paint Chips (%)				Soil (mg/kg)				Dust Wipes (µg)			
			Acceptable		Failures		Acceptable		Failures		Acceptable		Failures	
			N	%	N	%	N	%	N	%	N	%	N	%
FAA	Hotplate	NIOSH-7082/7105	147	90	17	10	76	90	8	10	231	96	9	4
		EPA-SW846-3050A	319	91	33	9	373	93	27	7	254	92	22	8
		Other-hotplate	91	91	9	9	30	94	2	6	38	95	2	5
	Microwave	EPA AREAL	3	75	1	25	7	88	1	13	3	75	1	25
		EPA-SW846-3051	44	100	0	0	43	98	1	2	31	97	1	3
		Other-microwave	20	100	0	0	0	0	0	0	16	100	0	0
	Other	All others	22	92	2	8	11	92	1	8	28	100	0	0
GFAA	Hotplate	NIOSH-7082/7105	5	63	3	38	4	100	0	0	9	75	3	25
		EPA-SW846-3050A	14	58	10	42	22	79	6	21	30	83	6	17
ICP-AES	Hotplate	NIOSH-7082/7105	50	96	2	4	33	83	7	18	77	96	3	4
		EPA-SW846-3050A	235	90	25	10	249	93	19	7	206	92	18	8
		Other-hotplate	31	97	1	3	18	90	2	10	49	94	3	6
	Microwave	EPA AREAL	7	88	1	13	8	100	0	0	8	100	0	0
		EPA-SW846-3051	46	96	2	4	43	90	5	10	42	95	2	5
		Other-microwave	18	90	2	10	12	100	0	0	13	81	3	19
	Other	All others	8	100	0	0	4	100	0	0	0	0	0	0
LAB-XRF	Other	All others	2	50	2	50	2	50	2	50	0	0	0	0
	—	—	0	0	4	100	2	50	2	50	0	0	0	0
Others	Hotplate	EPA-SW846-3050A	4	100	0	0	10	83	2	17	3	75	1	25
	Microwave	EPA-SW846-3051	1	25	3	75	4	100	0	0	2	50	2	50
		Other-microwave	8	100	0	0	0	0	0	0	0	0	0	0
Total			1075	90	121	10	951	92	85	8	1040	93	76	7

— = Not reported.

ing laboratories were classified into two groups: those that had no outliers on the four ELPAT samples of the matrix and those that had one or more outliers. Fischer's exact test was then repeated for each ELPAT matrix. No statistically significant differences were detected for soils ($p = 0.16$) or dust wipes ($p = 0.76$). However, statistically significant differences ($p = 0.012$) were detected in the paint chip digestion techniques/analytical method combinations. Of these combinations, the EPA SW846-3051 digestion technique with the FAA analytical method was performed by 11 laboratories with no outliers. The combination of EPA SW846-3050A hotplate with GFAA was performed by six laboratories with five outliers. Thirteen laboratories used the combination of the NIOSH 7082/7105 hotplate digestion technique with ICP with only one outlier. Twelve laboratories used the combination of

EPA SW846-3051 with ICP with only two outliers.

Inspection of Table 3 shows that the predominant analytical methods were FAA and ICP-AES. A total of 8 laboratories analyzed paint chips and soils and 12 laboratories analyzed dust wipes with GFAA. One laboratory used ICP-MS, one used DC plasma, and one used dithizone spectrophotometry. The laboratory that used ICP-MS had three of four samples as outliers for paint and two of four dust wipe samples as outliers. The laboratory that used DC plasma had four of four paint samples acceptable and the laboratory that used dithizone spectrophotometry analyzed for paint chips, soil, and dust wipes and had only one outlier on the dust wipes. Two laboratories used lab XRF, and had only limited success meeting the performance limits for paint chips and soils.

A more complete comparison of biases

and interlaboratory precision differences among digestion techniques and instrumental methods is being undertaken at NIOSH.

ELPAT Round 7 Bias Analysis

Statistical significance tests are performed for investigating differences in bias among the principal sample preparation and instrumental methods and among the combinations of these two factors. The tests are performed for each matrix (paint chips, soils, and dust wipes) and ELPAT sample (sample numbers 1, 2, 3, and 4) whenever at least three laboratories use the sample preparation and instrumental method.

Analysis of variance (ANOVA) is used if the data meet the general assumptions of the ANOVA procedure, homogeneity of variances, and normality. Bartlett's test is used for testing homogeneity of variances and the Shapiro-Wilk test is used



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for testing normality.^(13,14) If the ANOVA assumptions are violated, the Box-Cox transformation procedure is used to examine the data for possible transformations to correct the problem.⁽¹³⁾ If the transformed data meet the ANOVA assumptions, then the ANOVA tests are performed on the transformed data. If homogeneity of variance and normality are not achieved by transformation of the data, then a non-parametric approach is used.

In instances where variances are homogeneous and data are normally distributed (either before or after transformation), a one-way ANOVA followed by the Scheffe's multiple comparison test procedure is performed to test for differences in bias among the combinations of the principal sample preparation techniques and instrumental methods.⁽¹⁵⁾ A two-way ANOVA followed by Scheffe's multiple comparison test procedure to test for any difference among principal sample preparation techniques and prin-

cipal instrumental methods is also performed. Two-way ANOVAs separate bias that may be the result of sample preparation, instrumental method, or an interaction of these two factors.

In instances where ANOVA cannot be performed on either the original data or transformed data, one of two nonparametric tests is performed. If transformed data meet the homogeneity of variances but not the normality assumptions, then the Kruskal-Wallis rank sums test followed by the Mann-Whitney-Wilcoxon test with a Bonferroni adjustment is used.⁽¹⁶⁾ If no transformation can equalize the variances, then the median scores test followed by the sign test with Bonferroni adjustment is used.⁽¹⁷⁾

Further analyses of ELPAT round 7 were performed to determine if there were any statistically significant bias differences among sample preparation techniques and instrumental methods. The analyses compared the following principal sample preparation techniques:

NIOSH 7082/7105, EPA SW846-3050A, EPA SW846-3051, and EPA AREAL, and the following principal instrumental methods: FAA, GFAA, and ICP-AES.

One-way ANOVA procedures found bias among sample preparation/instrumental method combinations for paint chip samples 2 and 3 and dust wipe sample 2. Subsequent Scheffe's multiple comparison tests for differences in mean lead levels among any of the combinations found the following. For paint chip sample 2, EPA SW846-3051 and EPA SW846-3050A with ICP analysis gave lower results than EPA SW846-3050A, NIOSH 7082, and EPA SW846-3050 with FAA analysis. For paint chip sample 3; SW846-3051 with ICP analysis gave lower results than any other combination tested. For dust wipe sample 2; SW846-3050A with ICP analysis gave lower results than any other combination tested.

Two-way ANOVA procedures separate bias that may be the result of sample

preparation, instrumental method, or the interaction of these two factors. Two-way ANOVAs found three instances where bias was identified on ELPAT round 7: paint chip samples 1, 2, and 3. Subsequent Scheffe's multiple comparison test either could not identify the reason for the difference (paint chip sample 1) or attributed the difference to ICP, giving lower results than FAA.

ELPAT round 7 is consistent with findings on previous rounds that ICP sometimes gives lower results than FAA. For paint chip sample 2, the maximum difference in means is about 0.02 percent lead, or about 0.5 percent of the corresponding ELPAT reference laboratory mean. For paint chip sample 3, the maximum difference is about 0.013 percent lead, or about 16 percent of the corresponding reference laboratory mean.

The finding that ICP can sometimes give lower results than FAA is consistent with ANOVA findings on previous ELPAT rounds, and is consistent with the findings of an EPA-sponsored collab-

orative test of microwave and hotplate digestion techniques conducted by RTI. The RTI study found that ICP-AES consistently gave 3 to 18 percent lower results than FAA. The RTI study showed that failure to use background correction causes a positive bias in results for laboratories using FAA. The RTI study also found that suppression of the ICP signal for lead believed to be associated with matrix effects could be minimized by either: (1) diluting final solutions to a lead concentration of 10 μg of lead/ml; (2) using standard additions; or (3) using internal standards. Laboratories that do not do FAA background corrections or take ICP signal suppression into account may find it difficult to consistently meet ELPAT proficiency requirements.

While the magnitude of mean ICP-AES and FAA bias appears to be small, the bias for individual laboratories that do not perform FAA background correction or ICP-AES matrix effect minimization may be much larger. This is because FAA and ICP means used from ELPAT data

are based upon a composite performance of laboratories. FAA means may include both laboratories that do FAA background correction and those that do not. Similarly, ICP-AES laboratories may include laboratories that take ICP signal suppression into account and those that do not. It is therefore reasonable to assume that the actual bias, when analytical methods are not followed, may be much larger than the bias found in this ELPAT study.

Lead Reference Materials

The ELPAT program is designed to supplement, but not replace, a laboratory's internal quality control program. Use of materials of known lead content in suitable matrices is important in obtaining accurate and reliable lead results. Such materials should be used to validate methods when sample preparation techniques or instrumental methods are adopted or modified. In addition, the materials should be used for daily quality control charting of laboratory/analyst

TABLE 4. Certified Reference Materials

NIST Standard Reference Materials (SRMs)	Lead
SRM 1579a Powdered paint	$11.995 \pm 0.031\%$
SRM 2580 Powdered paint (to be released in 1995)	Nominal value 4%
SRM 2581 Powdered paint (to be released in 1995)	Nominal value 0.5%
SRM 2582 Powdered paint (total lead by weight)	$208.8 \pm 4.9 \text{ ppm}$
SRM 2709 Lead in soil	$18.9 \pm 0.5 \text{ ppm}$
SRM 2710 Lead in soil	$5532 \pm 80 \text{ ppm}$
SRM 2711 Lead in soil	$1162 \pm 31 \text{ ppm}$
SRM 2583 Lead in household dust (to be released in 1995) nominal value 100–200 ppm	
SRM 2579 Lead paint film on Mylar (set of 5)	$3.53 \pm 0.24 \text{ mg/cm}^2$ $1.63 \pm 0.08 \text{ mg/cm}^2$ $1.02 \pm 0.04 \text{ mg/cm}^2$ $0.29 \pm 0.01 \text{ mg/cm}^2$ Less than 0.001 mg/cm^2
(Intended for checking the calibration of portable X-ray fluorescence analyzers when testing for lead in paint coatings on interior and exterior building surfaces in the field)	
SRM 1648 Urban particulate matter	$0.655 \pm 0.008\%$
SRM 2704 Buffalo River sediment (total lead by weight)	$161 \pm 17 \text{ ppm}$
EPA/A2LA Certified Reference Materials	
Commercial supplier: RT Corporation through Fisher Scientific	
SRS014-50 Bag house dust	$1914 \pm 180 \text{ ppm}^*$
SRS013-50 Paint blasting waste	$643 \pm 56 \text{ ppm}^*$
SRS006-50 Paint sludge	$753 \pm 51 \text{ ppm}^*$

*The concentrations of lead determined in a sample following digestion by EPA Method 3010, 3020, and 3050. All concentrations expressed on dry weight basis. The 50-g samples should be mixed well before removing subsamples.

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performance. ELPAT paint chip, soil, and dust wipe samples from completed ELPAT rounds are available from AIHA at the address listed at the end of this column. ELPAT materials differ from the certified reference materials listed below. Either ELPAT materials are destroyed in one analysis (dust wipes), or the amount of material in bottles is limited to reduce the number of times that analyses can be repeated by laboratories reporting in the proficiency test round. National Institute of Standards and Technology (NIST) standard reference material (SRM) values report lead as total lead, whereas ELPAT- and EPA-certified reference materials report extractable lead.

Certified reference materials are commercially available from NIST and from commercial reference material suppliers participating in the EPA/A2LA environmental reference material certification program.⁽¹⁸⁾ The materials listed in Table 4 are useful for daily quality control of analyses and initial evaluation of methods associated with residential or steel structure lead abatement. Since work continues on developing additional reference materials, this list of certified reference materials is subject to significant change.

Updated lists of available certified reference materials are available from NIST, EPA-EMSL Cincinnati, and A2LA at the addresses listed at the end of this column.

EPA NLLAP

Under Title X of the Housing and Community Development Act of 1992, EPA, in consultation with the Department of Health and Human Services, has the responsibility to review and determine if effective voluntary laboratory accreditation systems are in place. If EPA determines that effective voluntary laboratory accreditation systems are not in place, EPA is responsible to establish a federal laboratory certification system.⁽¹⁹⁾

The EPA has established an NLLAP to recognize laboratories performing analysis associated with lead abatement. NLLAP recognition of laboratories analyzing lead in paint chips, soils, and dusts has two requirements: (1) successful participation in proficiency testing using real-world matrices; and (2) laboratory accreditation including on-site assessment of laboratory operations. NLLAP requirements are based upon the recommendations of a Federal Interagency Taskforce on Lead Based Paint, a group

of 17 federal agencies involved with lead issues, that recognition should be based upon both proficiency testing and laboratory accreditation.⁽²⁰⁾ Similarly, proficiency testing and laboratory accreditation requirements were also part of the recommendations for environmental laboratories of a 1991 National Conference on Laboratory Issues in Childhood Lead Poisoning Prevention sponsored by the Association of State and Territorial Public Health Laboratory Directors, the CDC, and EPA. NLLAP requirements for laboratories are based upon Guide 25-1990, "General Requirements for the Competence of Calibration and Testing Laboratories,"⁽²¹⁾ a guide already in use by many national laboratory accreditation systems worldwide.

The ELPAT Program began providing paint chip, soil, and dust audit samples to evaluate laboratory performance in the fall of 1992 and has grown to over 300 participating laboratories. In December 1993, the first two laboratory accreditation organizations, A2LA and AIHA, were recognized by NLLAP. Laboratories interested in obtaining accreditation information such as the program requirements, time needed to complete the

process, and cost should contact the recognized laboratory accreditation organizations. If other laboratory accreditation organizations are recognized, this information will be included in subsequent ELPAT columns.

Laboratory accreditation takes some time to achieve. Laboratory accreditation involves submittal of a description of a laboratory's quality system and manual to the accrediting organization and the on-site evaluation by NLLAP-qualified assessors of laboratory operations, including equipment, facilities, analytical methods, staff, and internal quality control.

Lists of laboratories that have performed successfully (rated proficient) in the ELPAT Program are prepared at NIOSH and are provided upon request to the public via a toll-free number by the Lead Information Clearinghouse (1-800-424-LEAD). The accreditation status of laboratories is included on the list of NLLAP-recognized laboratories provided to the public.

Once a sufficient number of laboratories (several hundred) geographically dispersed across the United States have received accreditation, only accredited laboratories which are ELPAT proficient will be NLLAP recognized and included on the list of laboratories provided to the public by the Lead Information Clearinghouse. Given the capacity of cooperating laboratory accreditation organizations to perform on-site assessments and initial laboratory demand for accreditation, it is projected that this will occur in 1995.

Participation in the ELPAT proficiency testing program would continue to be open to all interested laboratories. That means laboratories outside the United States and laboratories that do not wish to be accredited can continue to participate in ELPAT. However, starting in 1995, laboratories that are not accredited by an NLLAP-recognized laboratory accrediting organization will not be included on lists of NLLAP-recognized laboratories provided to the public by the federal government.

Upcoming ELPAT Round Information

Round 9 ELPAT samples were sent to participants on November 1, 1994. The reporting date of the laboratories was December 8, 1994. The dust wipes were preserved with 0.5 ml of 3 percent hydrogen peroxide solution. This is to re-

tard the formation of any fungal growth in the samples, and should not have any effect on the digestion and analysis of them. The ELPAT automated data entry system is available to laboratories that want to submit their laboratory data over a modem for faster and more reliable means of transmittal.

Information

A2LA Laboratory Accreditation, A2LA Certified Reference Materials, and A2LA/AIHA seminars on Environmental Lead Laboratory Accreditation:

American Association for Laboratory Accreditation (A2LA)
656 Quince Orchard Road
Gaithersburg, MD 20878
Phone: (301) 670-1377
FAX: (301) 869-1495

AIHA Laboratory Accreditation, ELPAT Program information, ELPAT sample orders, and A2LA/AIHA seminars on Environmental Lead Laboratory Accreditation:

ELPAT Coordinator
American Industrial Hygiene Association (AIHA)
2700 Prosperity Avenue, Suite #250
Fairfax, VA 22031
Phone: (703) 849-8888
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Orders for NIST SRMs:
National Institute of Standards and Technology
Standards Reference Materials Program
Room 204, Building 202
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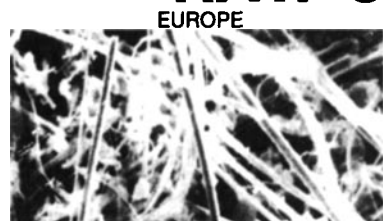
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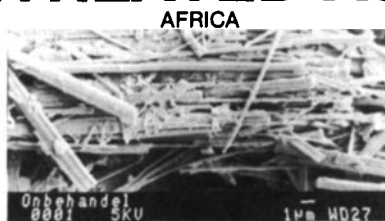
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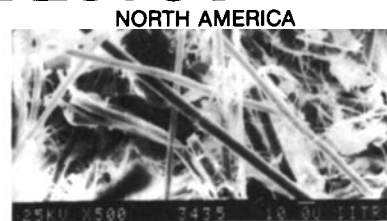
A Scanning Electron Microscope looks at **RAW UNTREATED ASBESTOS**



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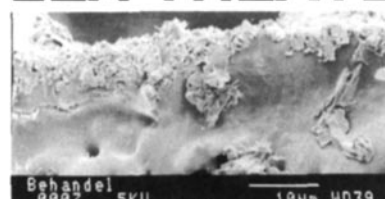


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