Adirondack
Environmental Services Inc.

Experience is the solution

PCB sampling, laboratory analysis, report and drawings: SUNY New Paltz

POLYCHLORINATED BIPHENYL (PCB) SAMPLING

September 1997

Prepared for:

SUNY New Paltz
75 South Manheim Blvd., Suite 9
New Paltz, New York 12561

Prepared by:

Adirondack Environmental Services, Inc.
260 Broadway, Suite 201
Saratoga Springs, NY 12866

Adirondack Project No. 970815EA

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EXECUTIVE SUMMARY

On August 18 and 19, 1997, Adirondack Environmental Services, Inc. (Adirondack) performed Polychlorinated Biphenyl (PCB) area air and wipe sampling in the Residence Halls located on the State University of New York (SUNY) New Paltz Campus. All monitoring and analysis activities were performed in accordance with the SUNY New Paltz Polychlorinated Biphenyl Air and Wipe Sampling Strategy, dated August 1997. The purpose of the sampling was to ensure that the four residence halls located on the campus were free of PCB contamination within a statistical confidence limit of 95%.

A total of 29 wipe samples and three field blanks and 30 air samples and three blanks were collected in randomly selected rooms in the four residence halls. Both the wipe sample locations and the area air sample locations were selected with a random number table. Samples were collected, labeled and placed on ice and delivered to the laboratory. Wipe samples were analyzed by the EPA 8080 method and the air samples were analyzed by the New York State Department of Health (NYDOH) method DOH 311-1.

The analysis of the 32 wipe samples collected indicated all results to be less than the detection limit of the analysis <0.1 ug/100cm^2 (<1ug per 900 cm^2). The analysis of the 33 air samples collected indicated results to be less than the detectable limits of the analysis (<0.2 ug/m^3) for all of the samples except one. Field sample identification number A24 (laboratory identification number 970820EA12) indicated a concentration of 0. 3 ug/m^3 for Aroclor 1242 and 0.2 ug/m^3 for Aroclor 1254. This sample was obtained outside Gage Hall.

The results of the wipe and air sampling indicate that all sample results were below Environmental Protection Agency, Occupational Safety and Health Administration and National Institute for Occupational Safety and Health published exposure limits for PCBs.

1.0 *INTRODUCTION*

On August 18 and 19, 1997, Mr. Paul Watson, CIH, CET and Mr. John Frisone, Industrial Hygiene Technician of Adirondack Environmental Services, Inc. (Adirondack) performed Polychlorinated Biphenyl (PCB) area air and wipe sampling in the four residence halls located on the State University of New York (SUNY), New Paltz Campus. The sampling team was escorted by Mr. Peter Betley of the SUNY New Paltz, Health and Safety Department. All monitoring and analysis activities were performed in accordance with the SUNY New Paltz Polychlorinated Biphenyl Air and Wipe Sampling Strategy, dated August 1997. A copy of the sampling strategy is located in Appendix A. The purpose of the sampling was to ensure that the four residence halls were free of PCB contamination that may have resulted from a release that occurred on December 29, 1991. The PCB material may have entered the residence halls by way of open windows, doors, and the ventilation systems. These residence halls were subsequently cleaned after the release and this sampling was conducted to ensure that the residence halls did not contain any residual material at a statistical confidence limit of 95%.

The sampling strategy and methodology used to complete the sampling is described in Section 2. The sample results and observations are presented in Section 3 and discussed in Section 4. The SUNY New Paltz Air and Wipe Sampling Strategy is included in Appendix A and the exact wipe and air sample locations are identified in Appendix B. The remaining Appendices present the laboratory results and field sampling data sheets.

2.0 SAMPLING STRATEGY AND METHODOLOGY

2.1 SAMPLING STRATEGY

The purpose of the exposure assessment was to obtain statistically representative wipe and air samples of the various residence hall rooms. A sampling strategy was designed to meet this objective. The sampling strategy developed for the facility is presented in Appendix A. This strategy served as a guide for the quantitative monitoring performed in this assessment.

2.2 SAMPLING AND ANALYTICAL METHODOLOGY

2.2.1 *Wipe Sampling*

Wipe samples were obtained utilizing 900 cm^2, disposable cardboard templates. Sample technicians utilized latex gloves and laboratory prepared swabs. Wipe sampling was performed by wiping a laboratory prepared filter paper over a 900 square centimeter area. The filter paper was prepared with 10 milliliters (ml) of reagent grade hexane. The pretreated wipes were replaced in the glass vials for storage and transport to the laboratory. All of the samples collected were assigned a sequential sample identifier and sample locations were plotted on site plans supplied by SUNY New Paltz. Wipe sample locations were selected with a random number table. Floor or desk top locations were selected in a similar fashion. All floor samples were obtained in the center of the residence hall rooms, except where otherwise noted on the field sample data sheets. All desk top samples were obtained in the area defined as the student's work area (that area directly in front of where a student may sit at the desk). All wipe samples were documented on a sampling record. This record included the location (identified by name and at least two measurements with a tape measure), date, room number, building and samplers name. All samples were assigned a unique sequential sample number. All samples were labeled, placed on ice and driven to the laboratory for analysis. Samples were analyzed by the EPA 8080 method for the following Aroclors; 1016, 1221, 1232, 1242, 1248, 1254 and 1260.

Identification of any PCB concentrations above 500 micrograms was to be confirmed by gas chromatography/mass spectrometry (GC/MS), however no peaks were identified above this level. The selection of this method was consistent with the methods previously utilized at SUNY New Paltz.

2.2.2 *Air Sampling*

Thirty -three air samples were collected using Mine Safety Appliances (MSA) Flow-lite air sampling pumps. All air sampling pumps were calibrated before and after sampling with a BIOS Dry Cal primary standard. All air samples were obtained in the center of the residence hall rooms with the doors and windows closed. All air samples were documented on an air sampling record. This record included the exact run time, pre and post calibration information, sample location and sampler's name. All samples were assigned a unique sequential identification number. Three (3) field blanks were submitted as per the sampling strategy. Blanks were not identified to the analytical laboratory and were assigned a sequential sample number.

Samples were analyzed by the DOH 311-1 method Analysis targeted the following Aroclors; 1016, 1221, 1232, 1242, 1248, 1254, and 1260. This method was selected to be consistent with previous air monitoring performed at SUNY New Paltz.

All air and wipe samples were submitted under chain of custody documentation to Adirondack's laboratory in Albany, New York for analysis. Wipe and air samples were stored in separate ice chests for delivery to the laboratory. Adirondack is accredited by the American Industrial Hygiene Association (AIHA), accreditation number 490.

3.0 *RESULTS AND OBSERVATIONS*

3.1 *RESULTS*

3.1.1 *PCB Wipe Sampling*

The analysis of the 32 wipe samples collected indicated all results to be less than the detection limit of the analysis, <0.1 ug/100cm² (<1ug per 900 cm²). These results can be compared to the Environmental Protection Agency's clean up criteria for a high contact residential/commercial areas of 10 ug/100 cm². Wipe sample locations are presented in Appendix B, Table 1.

3.1.2 *PCB Air Sampling*

The analysis of the 33 air samples collected indicated results to be less than the detectable limits of the analysis (<0.2 ug/m³) for all of the samples except one. Field sample identification number A24 (laboratory identification number 970820EA12) indicated a concentration of 0.3 ug/m³ for Aroclor 1242 and 0.2 ug/m³ for Aroclor 1254. This sample was obtained outside Gage Hall. These results can be compared to the Occupational Safety and Health Administration's (OSHA) Permissible Exposure Limit (PEL) of 500 ug/m³ for Aroclor 1254 and 1000 ug/m³ for Aroclor 1242. These are the only two Aroclors that OSHA has established PELs. The National Institute for Occupational Safety and Health (NIOSH) Recommended Exposure Limit (REL) for all PCBs is 1 ug/m³. Air sample locations are listed in Appendix B, Table 2.

3.2 *OBSERVATIONS*

Wipe sampling was conducted in unoccupied residence hall rooms. Where possible, SUNY New Paltz identification tags were utilized to identify desk tops that were sampled. However the identification tags were not always readily visible on the furniture and therefore were not documented.

All air monitoring was conducted with the doors and windows closed. All rooms

were unoccupied except for the following; Bliss Hall rooms 114, 223; Gage Hall 203, 325; Scudder Hall B3; and Capen Hall room 121. Students were not present in the rooms during the air monitoring periods.

Sample number A31 (room 214, Bliss Hall) was voided due to an air sampling pump failure. A31 was replaced with sample identifier A31A and was resampled.

4.0 *DISCUSSION AND CONCLUSION*

Based upon the findings, all sample results were less than the detection limits of the analysis except for one air sample, field number A24. This sample was obtained outside Gage Hall. This sample indicated two Aroclors (1242 and 1254) at the detection limit of the analysis.

It may be likely that the two Aroclors were components in the transformer oil (this oil was identified as Aroclor 1260) from the release incident. Commercial PCBs were generally mixtures of many different chlorinated biphenyls that were manufactured to meet specific requirements. These mixtures varied from batch to batch. Another possible explanation of the results is that the source of the material was from an off site release. PCBs can be present as solid or liquid aerosols and as vapors and depending on the type of PCB they can remain airborne for up to ten days. While dispensed in the air PCBs can travel fairly long distances.

The New York State Department of Health is currently conducting quarterly air monitoring on campus. These results should be compared to previously obtained results to determine if these Aroclors have been identified and are, as suspected, transient in nature.

Adirondack recommends that SUNY New Paltz maintain this and all previous sampling results for future reference.

APPENDIX A
SUNY NEW PALTZ AIR AND WIPE SAMPLING STRATEGY

Adirondack Environmental Services Inc

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POLYCHLORINATED BIPHENYL AIR AND WIPE SAMPLING STRATEGY

August 1997

Prepared for:

State University of New York, New Paltz Campus
Mr. Peter Betley
75 South Manheim Blvd., Suite 9
New Paltz, New York 12561

Prepared by:

Adirondack Environmental Services, Inc.
260 Broadway, Suite 201
Saratoga Springs, NY 12866

Adirondack Project No. 970805EB

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APPENDICES

1.0 INTRODUCTION

The following sampling plan has been developed to ensure that the four residence halls located on the State University of New York (SUNY) New Paltz campus are free of potential Polychlorinated Biphenyl (PCB) contamination. Area air monitoring and wipe sampling on horizontal surfaces have been prescribed. The number of air and wipe samples have been selected to provide a statistical confidence limit of 95%. Sampling methodology, selection of sample locations, and analytical methods are described in the text below.

2.0 *SAMPLING STRATEGY AND METHODOLOGY*

The purpose of the air and wipe sampling plan is to devise a statistically significant approach to the evaluation of approximately 422 residence hall rooms for PCB contamination.

In order to develop this sampling plan a few assumptions have been made. The first is that the four residence halls (approximately 422 rooms total, a room listing appears in Appendix A) have the same risk of potential exposure and are therefore defined as one homogeneous risk group. This assumption was based upon the information provided from SUNY New Paltz that the PCB release that occurred on December 29, 1991, may have emitted airborne PCBs. The PCB material may have entered the residence halls by way of open windows, doors, and the ventilation systems.

The second assumption is that PCB contamination is most likely present in the form of dust contamination and that horizontal surfaces (floors and desk tops) would be the most likely locations to detect any PCB material.

The final assumption that has been made is that the largest population that spends the greatest amount of time in the residence halls is the students. The majority of the students time spent would most likely be in the residence hall rooms, therefore sampling will concentrate on these areas and will exclude hallways, kitchen and common areas. Student contact with any potential PCB material would most likely be on the working area of the desk (approximately center) and the open areas of the floor (approximately center).

In determining that the four residence halls are free of PCB material, it will be necessary to randomly select rooms to be sampled. The sample group must be of sufficient size to ensure that the random sample will have a high probability to contain at least one sample containing PCB contamination if one exists. The number of samples (represented by n) to be collected from the homogenous risk group is dependent on the number of areas in the risk group (represented by N),

and the statistical confidence required (in this instance 95% confidence to collect at least one sample in the highest 10% of potential contamination).

Sample size n can be determined from a complex formula or by using the Table 1, Calculation of Sample Size for a Maximum Risk Subgroup from a Homogenous Exposure Risk Group. Prior to using the table, the following variables must be determined:

N = Size of the population with potential contamination (422 rooms in this case).
n = Size of the sample subgroup to be selected from the homogenous risk group.
α = Allowed probability of missing all of the areas in the top contamination group (α=0.05 for the 95% confidence level).
τ = Proportion of the group included as the high exposures-desired high in the exposure subgroup percentage (t=0.1 for the top 10% of all exposures).

From Table 1, the number of air and wipe samples for a population of 422 rooms, with a 95% confidence and a top 10% desired high exposure subgroup yields 29 samples each.

Table 1 - Calculation of Sample Size for a Maximum Risk Subgroup From a Homogeneous Exposure Risk Group

Size of partial Sample for Top 10% (τ=0.1) and Confidence 0.95 (α=0.05) (use n-N if N<11)

Size of Group, N	Number of Required Samples, n
12	11
13-14	12
15-16	13
17-18	14
19-21	15
22-24	16
25-27	17
28-31	18
32-35	19
36-41	20
42-50	21
∞	29

2.1 *SAMPLING STRATEGY*

2.1.1 *Wipe Samples*

Wipe sample locations (29) have been selected by assigning sequential numbers to each residence hall room starting with the number one (approximately 422 rooms, a room listing appears in Appendix A). Identified wipe sample locations appear in Appendix D. Locations must be indicated on drawings supplied by SUNY New Paltz. A random number table generator was utilized to select twenty-nine wipe sample locations. Once the rooms have been identified, the random number generator was utilized to determine whether a floor or desk top surface will be wiped. This was accomplished by assigning a unique number to the desk top location and one to the floor location. The random number table was then used to select the sample location. The sample location (the floor or desk top) must be documented on the SUNY New Paltz supplied site plan.

Samples will be obtained from the approximate center of the desk top or floor surface.

2.1.2 *Air Samples*

Area air sample locations were determined by assigning sequential number to each residence hall room starting with the number one (approximately 422 rooms, a room listing appears in Appendix A). Sample locations and sample numbers must be identified on drawings supplied by SUNY New Paltz. A random number table was utilized to select twenty-nine sample locations for the area air samples. Samples will be obtained at approximately four to five feet above the floor in order to represent a person's breathing zone. Samples will be taken in the center of the room with the doors and windows closed. One ambient outside air sample will also be obtained for comparison purposes.

2.2 *SAMPLING AND ANALYTICAL METHODOLOGY*

2.2.1 *Wipe Samples*

Wipe sampling will be performed by wiping a laboratory prepared four inch square gauze or glass wool wipe over a 900 square centimeter area. The filter paper will be prepared with 10 milliliters of reagent grade hexane. The pretreated wipes will be placed in a glass vial for storage and use. A pre cut 900 square centimeter reusable stainless steel templates will be utilized. This template will be supplied by SUNY New Paltz. The template will be decontaminated in-between each sample by wiping the entire surface with hexane and disposable towels. Disposable cardboard templates of the same size may also be used. These templates must be discarded after each sample. The sample technician will wear disposable gloves during the collection and handling of each sample to prevent potential sample cross contamination and skin exposure. All wipe samples will be documented on a sampling record. This record shall include at a minimum the location (to be identified by name and at least two measurements with a tape measure), date, room number, building and samplers name. All samples will be assigned a unique sequential sample number. Ten percent field blanks (3) will be submitted at the time of analysis. Blanks must not be identified to the analytical laboratory and must be assigned a sequential sample number. Blanks will be unopened laboratory prepared filters. Wipe samples will be obtained on horizontal surfaces only (floors and desk tops).

The analytical method will be EPA Method 8080 utilizing gas chromatography with an electron capture detector (GC/ECD). Identified PCB concentrations above 500 micrograms will be confirmed by gas chromatography/mass spectrometry (GC/MS). The following Aroclors must be targeted and quantified; 1016, 1221, 1232, 1242, 1248, 1254, and 1260. A copy of the method is included in Appendix B. The selection of this method is consistent with the methods previously utilized at SUNY New Paltz.

2.2.2 *Air Samples*

The New York State Department of Health Method 311-1, Polychlorinated

Biphenyls in Ambient Air, will be utilized as the air sampling and analytical method. Samples will be obtained with an air sampling pump, flexible tubing and appropriate media as outlined in the method. Sample time will be from four to eight hours at one liter per minute. A copy of the method is included in Appendix C. All samples will be pre and post calibrated with a National Institute of Standards and Technology (NIST) traceable primary standard such as a Gillian Gilabrator, Bios DRY CAL or equivalent. All air samples will be documented on an air sampling record. This record shall include at a minimum the run time, pre and post calibration information, sample location and samplers name. All samples will be assigned a unique sequential sample number. Ten percent field blanks (3) will be submitted at the time of analysis. Blanks must not be identified to the analytical laboratory and must be assigned a sequential sample number. Analysis must identify and quantify the following Aroclors; 1016, 1221, 1232, 1242, 1248, 1254, and 1260. This method was selected to be consistent with previous air monitoring performed at SUNY New Paltz.

2.3 SAMPLING HANDLING

2.3.1 Wipe samples

Immediately after wipe sampling, return the sample to the glass vial. Label the vial with an unique sample number and secure the sample for transport. Samples will be shipped to the laboratory in an ice chest in order to maintain the samples at 4^0 Celsius.

All samples must be accompanied to the laboratory with a properly completed chain of custody record.

All wipe samples must be shipped to the laboratory in a separate container from any air samples.

2.3.2 Air Samples

The air sampling train will be disassembled after completion of sampling. The

florisil tube will be capped immediately and labeled with a unique sample number and will be placed in a clean zip lock bag. The bag will be labeled with the unique sample number and secured for transport to the laboratory.

All samples must be accompanied to the laboratory with a properly completed chain of custody record. Samples will be shipped to the laboratory in an ice chest in order to maintain the samples at 4^0 Celsius

All air samples must be shipped to the laboratory in a separate container from any wipe samples.

2.4 PRESENTATION OF RESULTS

All supporting sampling documentation including chain of custody records, air sampling data forms, wipe sampling forms, and laboratory analytical results will be presented in a written report. The format will be determined by SUNY New Paltz. Wipe sample results will be compared to the Environmental Protection Agency's clean up criteria for a high contact residential/commercial area of 10 micrograms/100 square centimeters. Air sample results will be compared to the outside sample, sample blanks, and to National Institute for Occupational Safety and Health Administration airborne limits.

3.0 *REFERENCES*

Environmental Protection Agency, 40 CFR Part 761, Polychlorinated Biphenyls Spill Cleanup Policy: Final Rule

Environmental Protection Agency, *Field Manual For Grid Sampling of PCB Spill Sites to Verify Cleanup*, EPA Document Number 560/5-86-017

Environmental Protection Agency, *Verification of PCB Spill Cleanup By Sampling and Analysis*, EPA Document Number 560/5-86-026

National Institute for Occupational Safety and Health, *Occupational Sampling Strategy Manual*, DHEW Document Number 77-173

SCUDDER HALL, BUILDING #20

Basement
B1
B2
B3
B4
B5
B6
B7
B8
B9
B10
B11
B12
B13
B14
B15
B16
B17 Overflow

First Floor
103
105
107
108
109
Director's Apartment
113
115
116
117
119
121
123
101
102
104
106
110
111
Guest Room
112
114
118
120
122
124

Second Floor
203
205
207
208
209
211
213
215
216
217
219
221
223
224
225
227
229
201
202
204
206
210
212
214
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220
222
226
228
230
231

Third Floor
302
304
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308
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311
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315
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317
319
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324
325
327
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301
303
305
307
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312
314
318
320
322
326
328
330
331

Capen Hall, Building #9

Basement
B4
B5
B6
B7
B8
B9
B10
B11
B12
B13
B14
B15

First Floor
103
105
107
108
RD's Apartment
113
115
116
117
119
121
123
101HC
102HC
104HC
110
118HC
120HC
122HC
124HC

Second Floor
203
205
207
208
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211
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223
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227
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202
204
206
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222
226
228
230
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Third Floor
302
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311
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315
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317
319
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323
324
325
327
329
301
303
305
307
310
312
314
318
320
322
326
328
330
331

GAGE HALL, BUILDING #21

Basement East
B1
B2
B3
B4
B5
B6
B7
B8
B9
B10

Basement West
B11
B12
B13
B14
B15
B16
B17
B18
B19
B20
B21

First Floor East
101
103
105
107
108
108 Overflow
109
110
111
112
113
114
100 Overflow
102
104
106
RD's Apartment

First Floor West
115
117
119
121
127
129
131
133 Overflow
116
118
120
122
123
124
124 Overflow
125
126
128
130
132

Second Floor East
202
204
206
208
209
210
211
212
213
215
217
219
223
224
201
203
205
207
214
216

218
220
221
222

Second Floor West
227
229
231
233
235
237
243
245
247
249
225
226
228
230
232
234
236
238
239
240
241
242
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246
248

Third Floor East
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304
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308
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317
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321
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Third Floor West

327
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333
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337
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345
347
349
325
326
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348

BLISS HALL, BUILDING #22

BASEMENT

B00 Overflow
B1 Overflow
B2 Overflow
B3
B4
B5
B6
B7
B8
B9
B10
B11
B13
B14
B15
B16
B17 Overflow

FIRST FLOOR

101
102
104
106
110
111
Guest Room
112 Overflow
112
114
118
120
122
124
103
105
107
108
109
Director's Apartment
113
115
116
117
119
121
123

Second Floor

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Third Floor

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METHOD 8080A

ORGANOCHLORINE PESTICIDES AND POLYCHLORINATED BIPHENYLS BY GAS CHROMATOGRAPHY

1.0 SCOPE AND APPLICATION

1.1 Method 8080 is used to determine the concentration of various organochlorine pesticides and polychlorinated biphenyls (PCBs). The following compounds can be determined by this method:

Compound Name	CAS No.[a]
Aldrin	309-00-2
α-BHC	319-84-6
β-BHC	319-85-7
δ-BHC	319-86-8
γ-BHC (Lindane)	58-89-9
Chlordane (technical)	12789-03-6
4,4'-DDD	72-54-8
4,4'-DDE	72-55-9
4,4'-DDT	50-29-3
Dieldrin	60-57-1
Endosulfan I	959-98-8
Endosulfan II	33212-65-9
Endosulfan sulfate	1031-07-8
Endrin	72-20-8
Endrin aldehyde	7421-93-4
Heptachlor	76-44-8
Heptachlor epoxide	1024-57-3
4,4'-Methoxychlor	72-43-5
Toxaphene	8001-35-2
Aroclor-1016	12674-11-2
Aroclor-1221	1104-28-2
Aroclor-1232	11141-16-5
Aroclor-1242	53469-21-9
Aroclor-1248	12672-29-6
Aroclor-1254	11097-69-1
Aroclor-1260	11096-82-5

a Chemical Abstract Services Registry Number.

1.2 Table 1 lists the method detection limit for each compound in organic-free reagent water. Table 2 lists the estimated quantitation limit (EQL) for other matrices.

2.0 SUMMARY OF METHOD

2.1 Method 8080 provides gas chromatographic conditions for the detection of ppb concentrations of certain organochlorine pesticides and PCBs. Prior to the use of this method, appropriate sample extraction techniques must be used. Both neat and diluted organic liquids (Method 3580, Waste Dilution) may be analyzed by direct injection. A 2 to 5 µL sample is injected into a gas chromatograph (GC) using the solvent flush technique, and compounds in the GC effluent are detected by an electron capture detector (ECD) or an electrolytic conductivity detector (HECD).

2.2 The sensitivity of Method 8080 usually depends on the concentration of interferences rather than on instrumental limitations. If interferences prevent detection of the analytes, Method 8080 may also be performed on samples that have undergone cleanup. Method 3620, Florisil Column Cleanup, by itself or followed by Method 3660, Sulfur Cleanup, may be used to eliminate interferences in the analysis.

3.0 INTERFERENCES

3.1 Refer to Methods 3500, 3600, and 8000.

3.2 Interferences by phthalate esters can pose a major problem in pesticide determinations when using the electron capture detector. These compounds generally appear in the chromatogram as large late-eluting peaks, especially in the 15% and 50% fractions from the Florisil cleanup. Common flexible plastics contain varying amounts of phthalates. These phthalates are easily extracted or leached from such materials during laboratory operations. Cross contamination of clean glassware routinely occurs when plastics are handled during extraction steps, especially when solvent-wetted surfaces are handled. Interferences from phthalates can best be minimized by avoiding contact with any plastic materials. Exhaustive cleanup of reagents and glassware may be required to eliminate background phthalate contamination. The contamination from phthalate esters can be completely eliminated with a microcoulometric or electrolytic conductivity detector.

4.0 APPARATUS AND MATERIALS

4.1 Gas chromatograph

4.1.1 Gas Chromatograph: Analytical system complete with gas chromatograph suitable for on-column injections and all required accessories, including detectors, column supplies, recorder, gases, and syringes. A data system for measuring peak heights and/or peak areas is recommended.

4.1.2 Columns

4.1.2.1 Column 1: Supelcoport (100/120 mesh) coated with 1.5% SP-2250/1.95% SP-2401 packed in a 1.8 m x 4 mm ID glass column or equivalent.

4.1.2.2 Column 2: Supelcoport (100/120 mesh) coated with 3% OV-1 in a 1.8 m x 4 mm ID glass column or equivalent.

4.1.3 Detectors: Electron capture (ECD) or electrolytic conductivity detector (HECD).

4.2 Kuderna-Danish (K-D) apparatus:

4.2.1 Concentrator tube: 10 mL, graduated (Kontes K-570050-1025 or equivalent). A ground-glass stopper is used to prevent evaporation of extracts.

4.2.2 Evaporation flask: 500 mL (Kontes K-570001-500 or equivalent). Attach to concentrator tube with springs, clamps, or equivalent.

4.2.3 Snyder column: Three ball macro (Kontes K-503000-0121 or equivalent).

4.2.4 Snyder column: Two ball micro (Kontes K-569001-0219 or equivalent).

4.2.5 Springs - 1/2 inch (Kontes K-662750 or equivalent).

4.3 Boiling chips: Solvent extracted, approximately 10/40 mesh (silicon carbide or equivalent).

4.4 Water bath: Heated, with concentric ring cover, capable of temperature control (±5°C). The bath should be used in a hood.

4.5 Volumetric flasks, Class A: sizes as appropriate with ground-glass stoppers.

4.6 Microsyringe: 10 µL.

4.7 Syringe: 5 mL.

4.8 Vials: Glass, 2, 10, and 20 mL capacity with Teflon-lined screw caps or crimp tops.

4.9 Balances: Analytical, 0.0001 g and Top loading, 0.01 g.

5.0 REAGENTS

5.1 Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available. Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

5.2 Organic-free reagent water - All references to water in this method refer to organic-free reagent water, as defined in Chapter One.

5.3 Solvents

5.3.1 Hexane, C_6H_{14} - Pesticide quality or equivalent.

5.3.2 Acetone, CH_3COCH_3 - Pesticide quality or equivalent.

5.3.3 Toluene, $C_6H_5CH_3$ - Pesticide quality or equivalent.

5.3.4 Isooctane, $(CH_3)_3CCH_2CH(CH_3)_2$ - Pesticide quality or equivalent.

5.4 Stock standard solutions:

5.4.1 Prepare stock standard solutions at a concentration of 1000 mg/L by dissolving 0.0100 g of assayed reference material in isooctane and diluting to volume in a 10 mL volumetric flask. A small volume of toluene may be necessary to put some pesticides in solution. Larger volumes can be used at the convenience of the analyst. When compound purity is assayed to be 96% or greater, the weight can be used without correction to calculate the concentration of the stock standard. Commercially prepared stock standards can be used at any concentration if they are certified by the manufacturer or by an independent source.

5.4.2 Transfer the stock standard solutions into vials with Teflon-lined screw caps or crimp tops. Store at 4°C and protect from light. Stock standards should be checked frequently for signs of degradation or evaporation, especially just prior to preparing calibration standards from them.

5.4.3 Stock standard solutions must be replaced after one year, or sooner if comparison with check standards indicates a problem.

5.5 Calibration standards: Calibration standards at a minimum of five concentrations for each parameter of interest are prepared through dilution of the stock standards with isooctane. One of the concentrations should be at a concentration near, but above, the method detection limit. The remaining concentrations should correspond to the expected range of concentrations found in real samples or should define the working range of the GC. Calibration solutions must be replaced after six months, or sooner, if comparison with check standards indicates a problem.

5.6 Internal standards (if internal standard calibration is used): To use this approach, the analyst must select one or more internal standards that are similar in analytical behavior to the compounds of interest. The analyst must further demonstrate that the measurement of the internal standard is not affected by method or matrix interferences. Because of these limitations, no internal standard can be suggested that is applicable to all samples.

5.6.1 Prepare calibration standards at a minimum of five concentrations for each analyte of interest as described in Sec. 5.5.

5.6.2 To each calibration standard, add a known constant amount of one or more internal standards, and dilute to volume with isooctane.

5.6.3 Analyze each calibration standard according to Sec. 7.0.

5.7 Surrogate standards: The analyst should monitor the performance of the extraction, cleanup (when used), and analytical system and the effectiveness of the method in dealing with each sample matrix by spiking each sample, standard, and organic-free reagent water blank with pesticide surrogates. Because GC/ECD data are much more subject to interference than GC/MS, a secondary surrogate is to be used when sample interference is apparent. Two surrogate standards (tetrachloro-m-xylene (TCMX) and decachlorobiphenyl) are added to each sample; however, only one need be calculated for recovery. Proceed with corrective action when both surrogates are out of limits for a sample (Sec. 8.3). Method 3500 indicates the proper procedure for preparing these surrogates.

6.0 SAMPLE COLLECTION, PRESERVATION, AND HANDLING

6.1 See the introductory material to this chapter, Organic Analytes, Sec. 4.1. Extracts must be stored under refrigeration and analyzed within 40 days of extraction.

7.0 PROCEDURE

7.1 Extraction:

7.1.1 Refer to Chapter Two for guidance on choosing the appropriate extraction procedure. In general, water samples are extracted at a neutral, or as is, pH with methylene chloride, using either Method 3510 or 3520. Solid samples are extracted using Method 3540, 3541, or 3550.

7.1.2 Prior to gas chromatographic analysis, the extraction solvent must be exchanged to hexane. The exchange is performed during the K-D procedures listed in all of the extraction methods. The exchange is performed as follows.

7.1.2.1 Following K-D of the methylene chloride extract to 1 mL using the macro-Snyder column, allow the apparatus to cool and drain for at least 10 min.

7.1.2.2 Increase the temperature of the hot water bath to about 90°C. Momentarily remove the Snyder column, add 50 mL of hexane, a new boiling chip, and reattach the macro-Snyder column. Concentrate the extract using 1 mL of hexane to prewet the Snyder column. Place the K-D apparatus on the water bath so that the concentrator tube is partially immersed in the hot water. Adjust the vertical position of the apparatus and the water temperature, as required, to complete concentration in 5-10 min. At the proper rate of distillation the balls of the column will actively chatter, but the chambers will not flood. When the apparent volume of liquid

reaches 1 mL, remove the K-D apparatus and allow it to drain and cool for at least 10 min.

7.1.2.3 Remove the Snyder column and rinse the flask and its lower joint into the concentrator tube with 1-2 mL of hexane. A 5 mL syringe is recommended for this operation. Adjust the extract volume to 10.0 mL. Stopper the concentrator tube and store refrigerated at 4°C, if further processing will not be performed immediately. If the extract will be stored longer than two days, it should be transferred to a vial with a Teflon-lined screw cap or crimp top. Proceed with gas chromatographic analysis if further cleanup is not required.

7.2 Gas chromatography conditions (Recommended):

7.2.1 Column 1:

Carrier gas (5% methane/95% argon) flow rate:	60 mL/min
Column temperature:	200°C isothermal

When analyzing for the low molecular weight PCBs (PCB 1221-PCB 1248), it is advisable to set the oven temperature to 160°C.

7.2.2 Column 2:

Carrier gas (5% methane/95% argon) flow rate:	60 mL/min
Column temperature:	200°C isothermal

When analyzing for the low molecular weight PCBs (PCB 1221-PCB 1248), it is advisable to set the oven temperature to 140°C.

7.2.3 When analyzing for most or all of the analytes in this method, adjust the oven temperature and column gas flow to provide sufficient resolution for accurate quantitation of the analytes. This will normally result in a retention time of 10 to 12 minutes for 4,4'-DDT, depending on the packed column used.

7.3 Calibration: Refer to Method 8000 for proper calibration techniques. Use Table 1 and especially Table 2 for guidance on selecting the lowest point on the calibration curve.

7.3.1 The procedure for internal or external calibration may be used. Refer to Method 8000 for a description of each of these procedures.

7.3.2 Because of the low concentration of pesticide standards injected on a GC/ECD, column adsorption may be a problem when the GC has not been used for a day. Therefore, the GC column should be primed or deactivated by injecting a PCB or pesticide standard mixture approximately 20 times more concentrated than the mid-concentration standard. Inject this prior to beginning initial or daily calibration.

7.4 Gas chromatographic analysis:

7.4.1 Refer to Method 8000. If the internal standard calibration technique is used, add 10 µL of internal standard to the sample prior to injection.

7.4.2 Method 8000 provides instructions on the analysis sequence, appropriate dilutions, establishing daily retention time windows, and identification criteria. Include a mid-concentration standard after each group of 10 samples in the analysis sequence.

NOTE: A 72 hour sequence is not required with this method.

7.4.3 Examples of GC/ECD chromatograms for various pesticides and PCBs are shown in Figures 1 through 5.

7.4.4 Prime the column as per Sec. 7.3.2.

7.4.5 DDT and endrin are easily degraded in the injection port if the injection port or front of the column is dirty. This is the result of buildup of high boiling residue from sample injection. Check for degradation problems by injecting a mid-concentration standard containing only 4,4'-DDT and endrin. Look for the degradation products of 4,4'-DDT (4,4'-DDE and 4,4'-DDD) and endrin (endrin ketone and endrin aldehyde). If degradation of either DDT or endrin exceeds 20%, take corrective action before proceeding with calibration, by following the GC system maintenance outlined in of Method 8000. Calculate percent breakdown as follows:

$$\%\ \text{breakdown for 4,4'-DDT} = \frac{\text{Total DDT degradation peak area (DDE + DDD)}}{\text{Total DDT peak area (DDT + DDE + DDD)}} \times 100$$

$$\%\ \text{breakdown for Endrin} = \frac{\text{Total endrin degradation peak area (endrin aldehyde + endrin ketone)}}{\text{Total endrin peak area (endrin + endrin aldehyde + endrin ketone)}} \times 100$$

7.4.6 Record the sample volume injected and the resulting peak sizes (in area units or peak heights).

7.4.7 Using either the internal or external calibration procedure (Method 8000), determine the identity and quantity of each component peak in the sample chromatogram which corresponds to the compounds used for calibration purposes.

7.4.8 If peak detection and identification are prevented due to interferences, the hexane extract may need to undergo cleanup using Method 3620. The resultant extract(s) may be analyzed by GC directly or may undergo further cleanup to remove sulfur using Method 3660.

7.5 Cleanup:

7.5.1 Proceed with Method 3620, followed by, if necessary, Method 3660, using the 10 mL hexane extracts obtained from Sec. 7.1.2.3.

7.5.2 Following cleanup, the extracts should be analyzed by GC, as described in the previous sections and in Method 8000.

7.5.3 If only PCBs are to be measured in a sample, the sulfuric acid/permanganate cleanup (Method 3665), followed by Silica Cleanup (Method 3630) or Florisil Cleanup (Method 3620), is recommended.

7.6 Calculations (excerpted from U.S. FDA, PAM):

7.6.1 Calculation of Certain Residues: Residues which are mixtures of two or more components present problems in measurement. When they are found together, e.g., toxaphene and DDT, the problem of quantitation becomes even more difficult. In the following sections suggestions are offered for handling toxaphene, chlordane, PCB, DDT, and BHC. A 10% DC-200 stationary phase column was used to obtain the chromatograms in Figures 6-9.

7.6.2 Toxaphene: Quantitative calculation of toxaphene or Strobane is difficult, but reasonable accuracy can be obtained. To calculate toxaphene on GC/ECD: (a) adjust sample size so that toxaphene major peaks are 10-30% full-scale deflection (FSD); (b) inject a toxaphene standard that is estimated to be within $\pm$10 ng of the sample; (c) construct the baseline of standard toxaphene between its extremities; and (d) construct the baseline under the sample, using the distances of the peak troughs to baseline on the standard as a guide (Figures 7, 8, and 9). This procedure is made difficult by the fact that the relative heights and widths of the peaks in the sample will probably not be identical to the standard. A toxaphene standard that has been passed through a Florisil column will show a shorter retention time for peak X and an enlargement of peak Y.

7.6.3 Toxaphene and DDT: If DDT is present, it will superimpose itself on toxaphene peak V. To determine the approximate baseline of the DDT, draw a line connecting the trough of peaks U and V with the trough of peaks W and X and construct another line parallel to this line which will just cut the top of peak W (Figure 61). This procedure was tested with ratios of standard toxaphene-DDT mixtures from 1:10 to 2:1 and the results of added and calculated DDT and toxaphene by the "parallel lines" method of baseline construction were within 10% of the actual values in all cases.

7.6.3.1 A series of toxaphene residues have been calculated using total peak area for comparison to the standard and also using area of the last four peaks only in both sample and standard. The agreement between the results obtained by the two methods justifies the use of the latter method for calculating toxaphene in a sample where the early eluting portion of the toxaphene chromatogram is interfered with by other substances.

7.6.3.2 The baseline for methoxychlor superimposed on toxaphene (Figure 8b) was constructed by overlaying the samples on a toxaphene standard of approximately the same concentration (Figure 8a) and viewing the charts against a lighted background.

7.6.4 Chlordane is a technical mixture of at least 11 major components and 30 or more minor ones. Gas chromatography-mass spectrometry and nuclear magnetic resonance analytical techniques have been applied to the elucidation of the chemical structures of the many chlordane constituents. Figure 9a is a chromatogram of standard chlordane. Peaks E and F are responses to trans- and cis-chlordane, respectively. These are the two major components of technical chlordane, but the exact percentage of each in the technical material is not completely defined and is not consistent from batch to batch. Other labelled peaks in Figure 9a are thought to represent: A, monochlorinated adduct of pentachlorocyclopentadiene with cyclopentadiene; B, coelution of heptachlor and α-chlordene; C, coelution of β-chlordene and γ-chlordene; D, a chlordane analog; G, coelution of cis-nonachlor and "Compound K," a chlordane isomer. The right "shoulder" of peak F is caused by trans-nonachlor.

7.6.4.1 The GC pattern of a chlordane residue may differ considerably from that of the technical standard. Depending on the sample substrate and its history, residues of chlordane can consist of almost any combination of constituents from the technical chlordane, plant and/or animal metabolites, and products of degradation caused by exposure to environmental factors such as water and sunlight. Only limited information is available on which residue GC patterns are likely to occur in which samples types, and even this information may not be applicable to a situation where the route of exposure is unusual. For example, fish exposed to a recent spill of technical chlordane will contain a residue drastically different from a fish whose chlordane residue was accumulated by ingestion of smaller fish or of vegetation, which in turn had accumulated residues because chlordane was in the water from agricultural runoff.

7.6.4.2 Because of this inability to predict a chlordane residue GC pattern, it is not possible to prescribe a single method for the quantitation of chlordane residues. The analyst must judge whether or not the residue's GC pattern is sufficiently similar to that of a technical chlordane reference material to use the latter as a reference standard for quantitation.

7.6.4.3 When the chlordane residue does not resemble technical chlordane, but instead consists primarily of individual, identifiable peaks, quantitate each peak separately against the appropriate reference materials and report the individual residues. (Reference materials are available for at least 11 chlordane constituents, metabolites or degradation products which may occur in the residue.)

7.6.4.4 When the GC pattern of the residue resembles that of technical chlordane, quantitate chlordane residues by comparing the total area of the chlordane chromatogram from peaks A through F (Figure 9a) in the sample versus the same part of the standard chromatogram. Peak G may be obscured in a sample by the presence of other pesticides. If G is not obscured, include it in the measurement for both standard and sample. If the heptachlor epoxide peak is relatively small, include it as part of the total chlordane area for calculation of the residue. If heptachlor and/or heptachlor epoxide are much out of proportion as in Figure 6j, calculate these separately and subtract their areas from total area to give a corrected chlordane area. (Note that octachlor epoxide, a metabolite of chlordane, can easily be mistaken for heptachlor epoxide on a nonpolar GC column.)

7.6.4.5 To measure the total area of the chlordane chromatogram, proceed as in Sec. 7.6.2 on toxaphene. Inject an amount of technical chlordane standard which will produce a chromatogram in which peaks E and F are approximately the same size as those in the sample chromatograms. Construct the baseline beneath the standard from the beginning of peak A to the end of peak F as shown in Figure 9a. Use the distance from the trough between peaks E and F to the baseline in the chromatogram of the standard to construct the baseline in the chromatogram of the sample. Figure 9b shows how the presence of toxaphene causes the baseline under chlordane to take an upward angle. When the size of peaks E and F in standard and sample chromatograms are the same, the distance from the trough of the peaks to the baselines should be the same. Measurement of chlordane area should be done by total peak area if possible.

NOTE: A comparison has been made of the total peak area integration method and the addition of peak heights method for several samples containing chlordane. The peak heights A, B, C, D, E, and F were measured in millimeters from peak maximum of each to the baseline constructed under the total chlordane area and were then added together. These results obtained by the two techniques are too close to ignore this method of "peak height addition" as a means of calculating chlordane. The technique has inherent difficulties because not all the peaks are symmetrical and not all are present in the same ratio in standard and in sample. This method does offer a means of calculating results if no means of measuring total area is practical.

7.6.5 Polychlorinated biphenyls (PCBs): Quantitation of residues of PCB involves problems similar to those encountered in the quantitation of toxaphene, Strobane, and chlordane. In each case, the chemical is made up of numerous compounds. So the chromatograms are multi-peak. Also in each case, the chromatogram of the residue may not match that of the standard.

7.6.5.1 Mixtures of PCBs of various chlorine contents were sold for many years in the U.S. by the Monsanto Co. under the

tradename Aroclor (1200 series and 1016). Though these Aroclors are no longer marketed, the PCBs remain in the environment and are sometimes found as residues in foods, especially fish.

7.6.5.2 PCB residues are quantitated by comparison to one or more of the Aroclor materials, depending on the chromatographic pattern of the residue. A choice must be made as to which Aroclor or mixture of Aroclors will produce a chromatogram most similar to that of the residue. This may also involve a judgment about what proportion of the different Aroclors to combine to produce the appropriate reference material.

7.6.5.3 Quantitate PCB residues by comparing total area or height of residue peaks to total area of height of peaks from appropriate Aroclor(s) reference materials. Measure total area or height response from common baseline under all peaks. Use only those peaks from the sample that can be attributed to chlorobiphenyls. These peaks must also be present in the chromatogram of the reference materials. Mixtures of Aroclors may be required to provide the best match of GC patterns of sample and reference.

7.6.6 DDT: DDT found in samples often consists of both o,p'- and p,p'-DDT. Residues of DDE and DDD are also frequently present. Each isomer of DDT and its metabolites should be quantitated using the pure standard of that compound and reported as such.

7.6.7 Hexachlorocyclohexane (BHC, from the former name, benzene hexachloride): Technical grade BHC is a cream-colored amorphous solid with a very characteristic musty odor; it consists of a mixture of six chemically distinct isomers and one or more heptachloro-cyclohexanes and octachloro-cyclohexanes.

7.6.7.1 Commercial BHC preparations may show a wide variance in the percentage of individual isomers present. The elimination rate of the isomers fed to rats was 3 weeks for the α-, γ-, and δ-isomers and 14 weeks for the β-isomer. Thus it may be possible to have any combination of the various isomers in different food commodities. BHC found in dairy products usually has a large percentage of β-isomer.

7.6.7.2 Individual isomers (α, β, γ, and δ) were injected into gas chromatographs equipped with flame ionization, microcoulometric, and electron capture detectors. Response for the four isomers is very nearly the same whether flame ionization or microcoulometric GLC is used. The α-, γ-, and δ-isomers show equal electron affinity. β-BHC shows a much weaker electron affinity compared to the other isomers.

7.6.7.3 Quantitate each isomer (α, β, γ, and δ) separately against a standard of the respective pure isomer, using a GC column which separates all the isomers from one another.

8.0 QUALITY CONTROL

8.1 Refer to Chapter One for specific quality control procedures. Quality control to validate sample extraction is covered in Method 3500 and in the extraction method utilized. If extract cleanup was performed, follow the QC in Method 3600 and in the specific cleanup method.

8.2 Quality control required to evaluate the GC system operation is found in Method 8000.

8.2.1 The quality control check sample concentrate (Method 8000) should contain each single-component parameter of interest at the following concentrations in acetone or other water miscible solvent: 4,4'-DDD, 10 mg/L; 4,4'-DDT, 10 mg/L; endosulfan II, 10 mg/L; endosulfan sulfate, 10 mg/L; endrin, 10 mg/L; and any other single-component pesticide, 2 mg/L. If this method is only to be used to analyze for PCBs, chlordane, or toxaphene, the QC check sample concentrate should contain the most representative multi-component parameter at a concentration of 50 mg/L in acetone.

8.2.2 Table 3 indicates the QC acceptance criteria for this method. Table 4 gives method accuracy and precision as functions of concentration for the analytes of interest. The contents of both Tables should be used to evaluate a laboratory's ability to perform and generate acceptable data by this method.

8.3 Calculate surrogate standard recovery on all samples, blanks, and spikes. Determine if the recovery is within limits (limits established by performing QC procedures outlined in Method 8000).

8.3.1 If recovery is not within limits, the following is required.

- Check to be sure there are no errors in calculations, surrogate solutions and internal standards. Also, check instrument performance.
- Recalculate the data and/or reanalyze the extract if any of the above checks reveal a problem.
- Reextract and reanalyze the sample if none of the above are a problem or flag the data as "estimated concentration".

8.4 GC/MS confirmation: Any compounds confirmed by two columns may also be confirmed by GC/MS if the concentration is sufficient for detection by GC/MS as determined by the laboratory generated detection limits.

8.4.1 The GC/MS would normally require a minimum concentration of 10 ng/μL in the final extract, for each single-component compound.

8.4.2 The pesticide extract and associated blank should be analyzed by GC/MS as per Sec. 7.0 of Method 8270.

8.4.3 The confirmation may be from the GC/MS analysis of the base/neutral-acid extractables extracts (sample and blank). However, if the compounds are not detected in the base/neutral-acid extract even though the concentration is high enough, a GC/MS analysis of the pesticide extract should be performed.

8.4.4 A reference standard of the compound must also be analyzed by GC/MS. The concentration of the reference standard must be at a level that would demonstrate the ability to confirm the pesticides/PCBs identified by GC/ECD.

9.0 METHOD PERFORMANCE

9.1 The method was tested by 20 laboratories using organic-free reagent water, drinking water, surface water, and three industrial wastewaters spiked at six concentrations. Concentrations used in the study ranged from 0.5 to 30 µg/L for single-component pesticides and from 8.5 to 400 µg/L for multi-component parameters. Single operator precision, overall precision, and method accuracy were found to be directly related to the concentration of the parameter and essentially independent of the sample matrix. Linear equations to describe these relationships for an electron capture detector are presented in Table 4.

9.2 The accuracy and precision obtained will be determined by the sample matrix, sample-preparation technique, optional cleanup techniques, and calibration procedures used.

10.0 REFERENCES

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TABLE 1.
GAS CHROMATOGRAPHY OF PESTICIDES AND PCBs*

Analyte	Retention time (min) Col. 1	Col. 2	Method Detection limit (µg/L)
Aldrin	2.40	4.10	0.004
α-BHC	1.35	1.82	0.003
β-BHC	1.90	1.97	0.006
δ-BHC	2.15	2.20	0.009
γ-BHC (Lindane)	1.70	2.13	0.004
Chlordane (technical)	e	e	0.014
4,4'-DDD	7.83	9.08	0.011
4,4'-DDE	5.13	7.15	0.004
4,4'-DDT	9.40	11.75	0.012
Dieldrin	5.45	7.23	0.002
Endosulfan I	4.50	6.20	0.014
Endosulfan II	8.00	8.28	0.004
Endosulfan sulfate	14.22	10.70	0.066
Endrin	6.55	8.10	0.006
Endrin aldehyde	11.82	9.30	0.023
Heptachlor	2.00	3.35	0.003
Heptachlor epoxide	3.50	5.00	0.083
Methoxychlor	18.20	26.60	0.176
Toxaphene	e	e	0.24
PCB-1016	e	e	nd
PCB-1221	e	e	nd
PCB-1232	e	e	nd
PCB-1242	e	e	0.065
PCB-1248	e	e	nd
PCB-1254	e	e	nd
PCB-1260	e	e	nd

*U.S. EPA. Method 617. Organochlorine Pesticides and PCBs. Environmental Monitoring and Support Laboratory, Cincinnati, Ohio 45268.

e = Multiple peak response.

nd = not determined.

TABLE 2.
DETERMINATION OF ESTIMATED QUANTITATION LIMITS (EQLs) FOR VARIOUS MATRICES[a]

Matrix	Factor
Ground water	10
Low-concentration soil by sonication with GPC cleanup	670
High-concentration soil and sludges by sonication	10,000
Non-water miscible waste	100,000

a EQL = [Method detection limit (see Table 1)] X [Factor found in this table]. For non-aqueous samples, the factor is on a wet-weight basis. Sample EQLs are highly matrix-dependent. The EQLs listed herein are provided for guidance and may not always be achievable.

TABLE 3.
QC ACCEPTANCE CRITERIA[a]

Analyte	Test conc. (μg/L)	Limit for s (μg/L)	Range for $\bar{x}$ (μg/L)	Range P, P_s (%)
Aldrin	2.0	0.42	1.08-2.24	42-122
α-BHC	2.0	0.48	0.98-2.44	37-134
β-BHC	2.0	0.64	0.78-2.60	17-147
δ-BHC	2.0	0.72	1.01-2.37	19-140
γ-BHC	2.0	0.46	0.86-2.32	32-127
Chlordane	50	10.0	27.6-54.3	45-119
4,4'-DDD	10	2.8	4.8-12.6	31-141
4,4'-DDE	2.0	0.55	1.08-2.60	30-145
4,4'-DDT	10	3.6	4.6-13.7	25-160
Dieldrin	2.0	0.76	1.15-2.49	36-146
Endosulfan I	2.0	0.49	1.14-2.82	45-153
Endosulfan II	10	6.1	2.2-17.1	D-202
Endosulfan Sulfate	10	2.7	3.8-13.2	26-144
Endrin	10	3.7	5.1-12.6	30-147
Heptachlor	2.0	0.40	0.86-2.00	34-111
Heptachlor epoxide	2.0	0.41	1.13-2.63	37-142
Toxaphene	50	12.7	27.8-55.6	41-126
PCB-1016	50	10.0	30.5-51.5	50-114
PCB-1221	50	24.4	22.1-75.2	15-178
PCB-1232	50	17.9	14.0-98.5	10-215
PCB-1242	50	12.2	24.8-69.6	39-150
PCB-1248	50	15.9	29.0-70.2	38-158
PCB-1254	50	13.8	22.2-57.9	29-131
PCB-1260	50	10.4	18.7-54.9	8-127

s = Standard deviation of four recovery measurements, in μg/L.

$\bar{x}$ = Average recovery for four recovery measurements, in μg/L.

P, P_s = Percent recovery measured.

D = Detected; result must be greater than zero.

[a]Criteria from 40 CFR Part 136 for Method 608. These criteria are based directly upon the method performance data in Table 4. Where necessary, the limits for recovery have been broadened to assure applicability of the limits to concentrations below those used to develop Table 4.

TABLE 4.
METHOD ACCURACY AND PRECISION AS FUNCTIONS OF CONCENTRATION*

Analyte	Accuracy, as recovery, x' ($\mu g/L$)	Single analyst precision, s_r' ($\mu g/L$)	Overall precision, S' ($\mu g/L$)
Aldrin	0.81C+0.04	0.16$\bar{x}$-0.04	0.20$\bar{x}$-0.01
α-BHC	0.84C+0.03	0.13$\bar{x}$+0.04	0.23$\bar{x}$-0.00
β-BHC	0.81C+0.07	0.22$\bar{x}$+0.02	0.33$\bar{x}$-0.95
δ-BHC	0.81C+0.07	0.18$\bar{x}$+0.09	0.25$\bar{x}$+0.03
γ-BHC	0.82C-0.05	0.12$\bar{x}$+0.06	0.22$\bar{x}$+0.04
Chlordane	0.82C-0.04	0.13$\bar{x}$+0.13	0.18$\bar{x}$+0.18
4,4'-DDD	0.84C+0.30	0.20$\bar{x}$-0.18	0.27$\bar{x}$-0.14
4,4'-DDE	0.85C+0.14	0.13$\bar{x}$+0.06	0.28$\bar{x}$-0.09
4,4'-DDT	0.93C-0.13	0.17$\bar{x}$+0.39	0.31$\bar{x}$-0.21
Dieldrin	0.90C+0.02	0.12$\bar{x}$+0.19	0.16$\bar{x}$+0.16
Endosulfan I	0.97C+0.04	0.10$\bar{x}$+0.07	0.18$\bar{x}$+0.08
Endosulfan II	0.93C+0.34	0.41$\bar{x}$-0.65	0.47$\bar{x}$-0.20
Endosulfan Sulfate	0.89C-0.37	0.13$\bar{x}$+0.33	0.24$\bar{x}$+0.35
Endrin	0.89C-0.04	0.20$\bar{x}$+0.25	0.24$\bar{x}$+0.25
Heptachlor	0.69C+0.04	0.06$\bar{x}$+0.13	0.16$\bar{x}$+0.08
Heptachlor epoxide	0.89C+0.10	0.18$\bar{x}$-0.11	0.25$\bar{x}$-0.08
Toxaphene	0.80C+1.74	0.09$\bar{x}$+3.20	0.20$\bar{x}$+0.22
PCB-1016	0.81C+0.50	0.13$\bar{x}$+0.15	0.15$\bar{x}$+0.45
PCB-1221	0.96C+0.65	0.29$\bar{x}$-0.76	0.35$\bar{x}$-0.62
PCB-1232	0.91C+10.79	0.21$\bar{x}$-1.93	0.31$\bar{x}$+3.50
PCB-1242	0.91C+10.79	0.21$\bar{x}$-1.93	0.31$\bar{x}$+3.50
PCB-1248	0.91C+10.79	0.21$\bar{x}$-1.93	0.31$\bar{x}$+3.50
PCB-1254	0.91C+10.79	0.21$\bar{x}$-1.93	0.31$\bar{x}$+3.50
PCB-1260	0.91C+10.79	0.21$\bar{x}$-1.93	0.31$\bar{x}$+3.50

x' = Expected recovery for one or more measurements of a sample containing concentration C, in $\mu g/L$.

s_r' = Expected single analyst standard deviation of measurements at an average concentration of $\bar{x}$, in $\mu g/L$.

S' = Expected interlaboratory standard deviation of measurements at an average concentration found of $\bar{x}$, in $\mu g/L$.

C = True value for the concentration, in $\mu g/L$.

$\bar{x}$ = Average recovery found for measurements of samples containing a concentration of C, in $\mu g/L$.

Figure 1
Gas Chromatogram of Pesticides

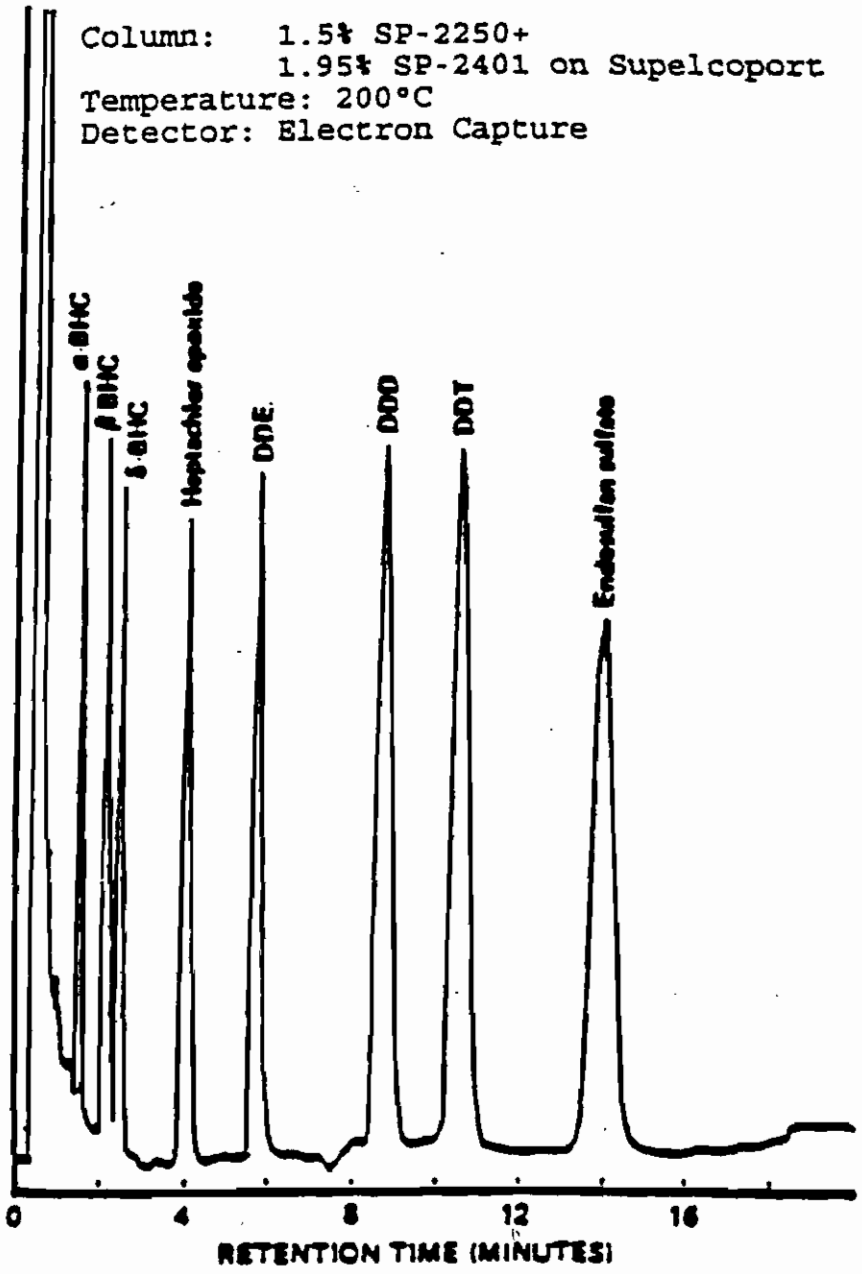


Figure 2
Gas Chromatogram of Chlordane

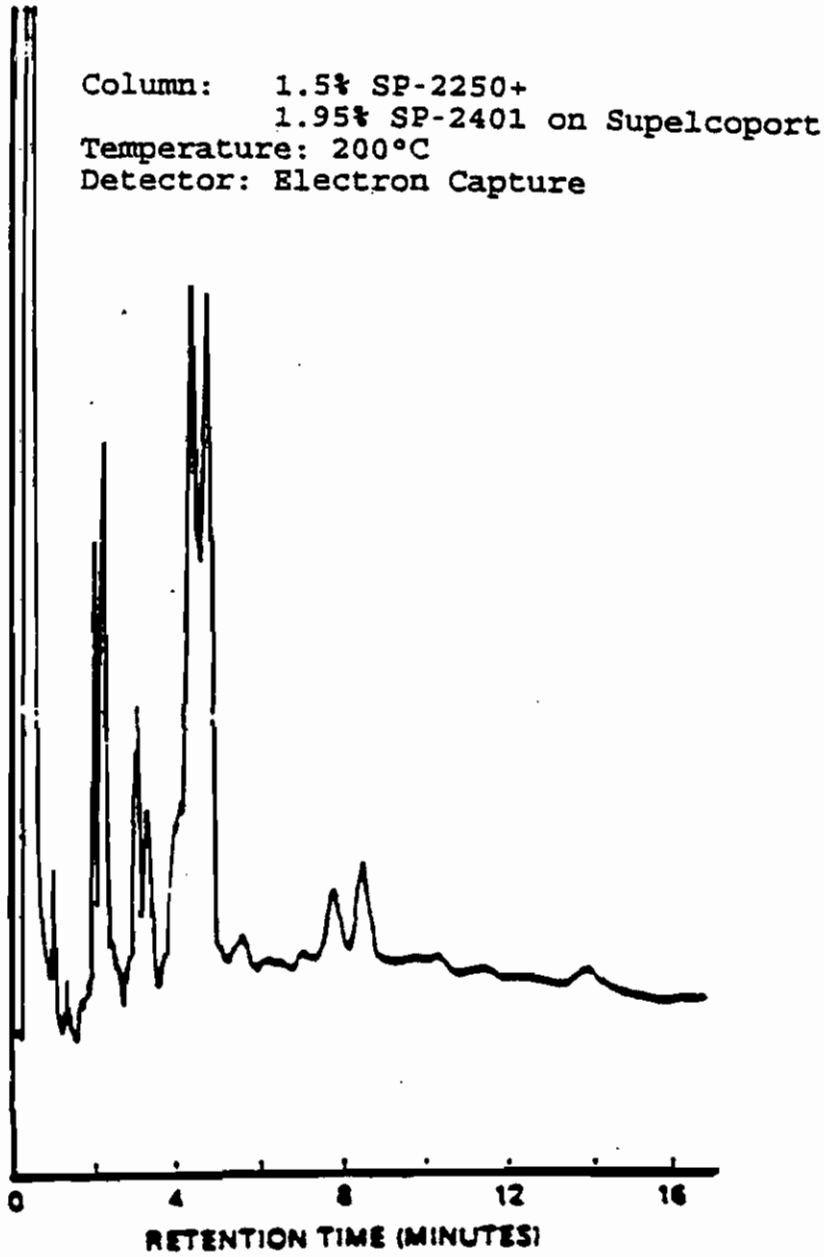


Figure 3
Gas Chromatogram of Toxaphene

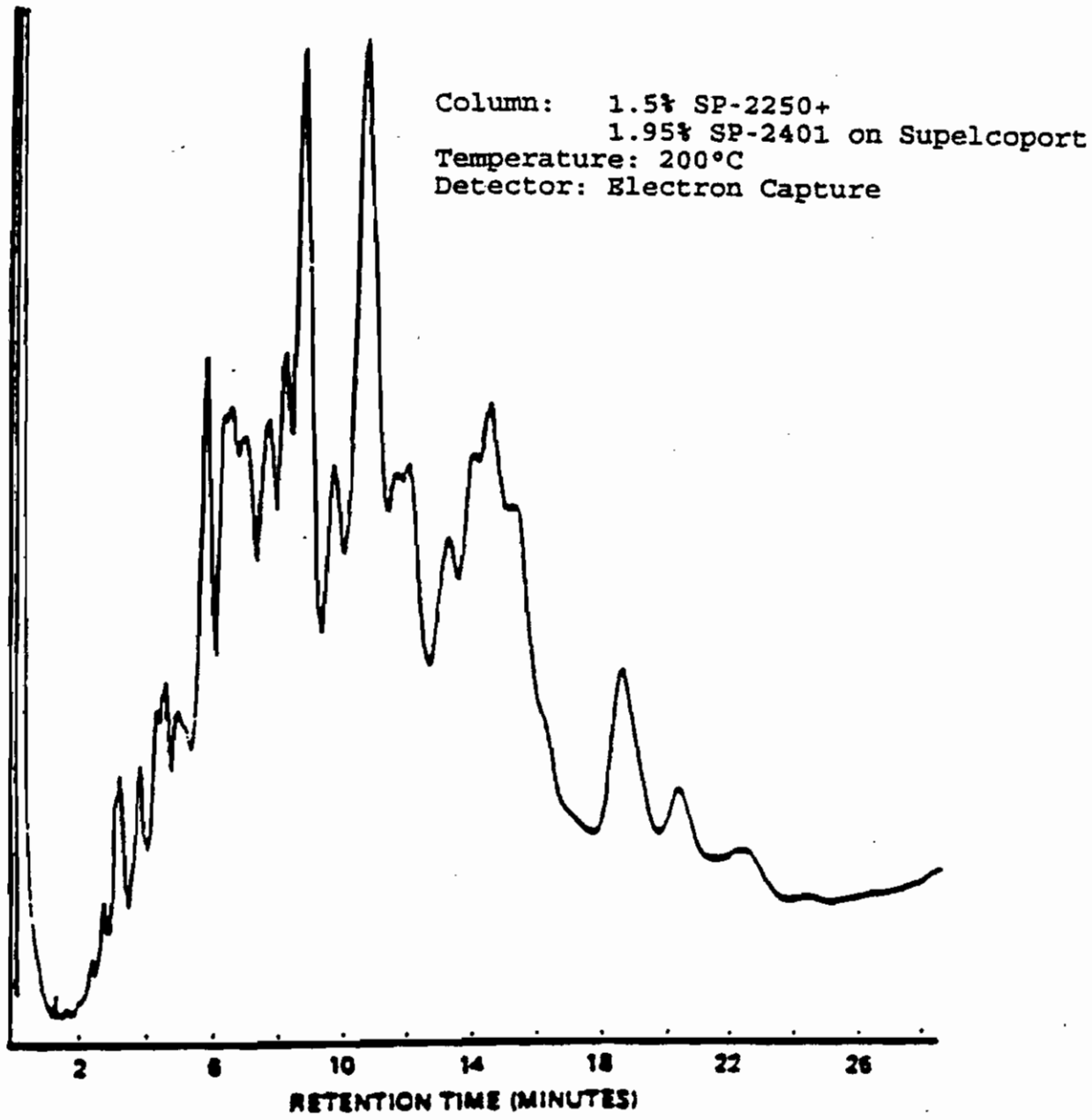


Figure 4
Gas Chromatogram of Aroclor 1254

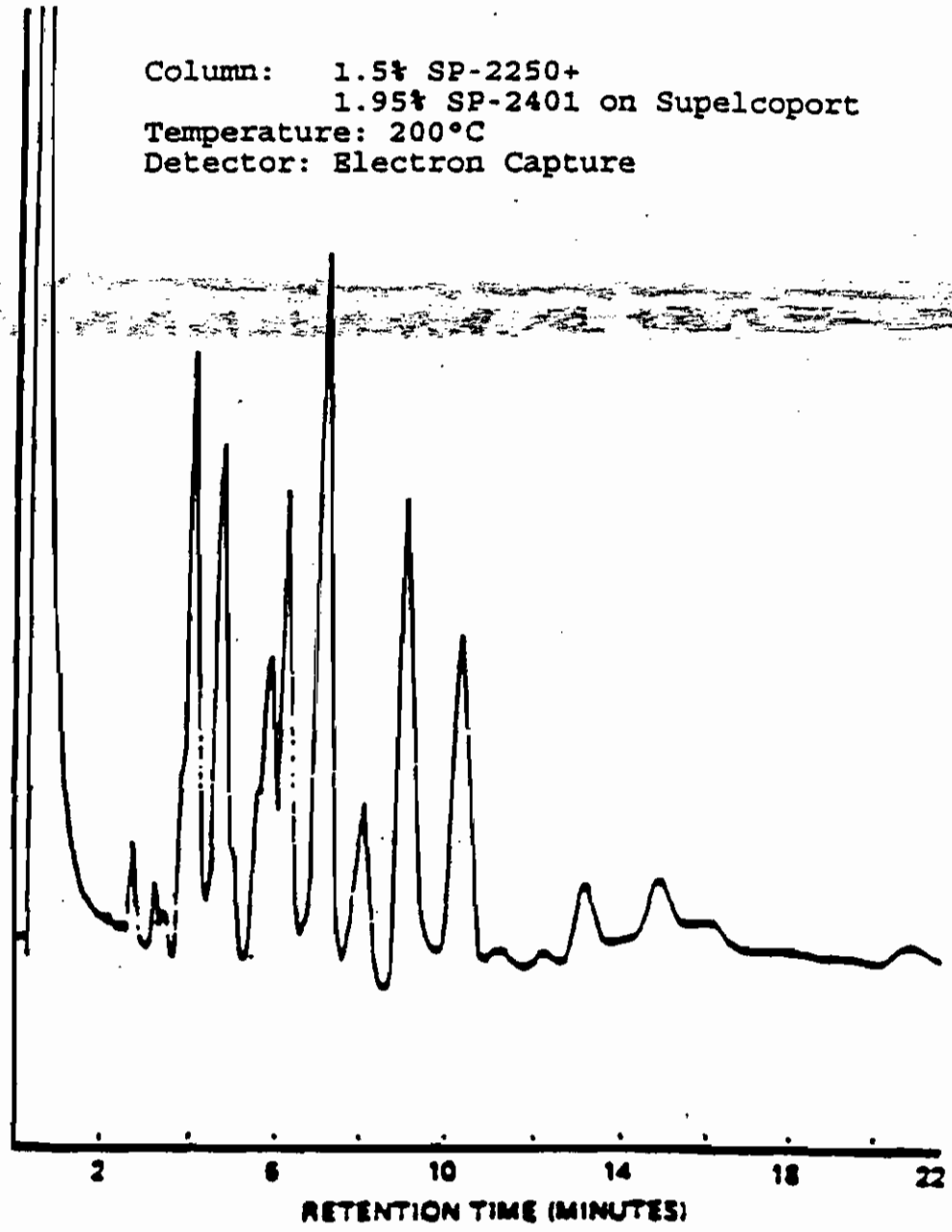


Figure 5
Gas Chromatogram of Aroclor 1260

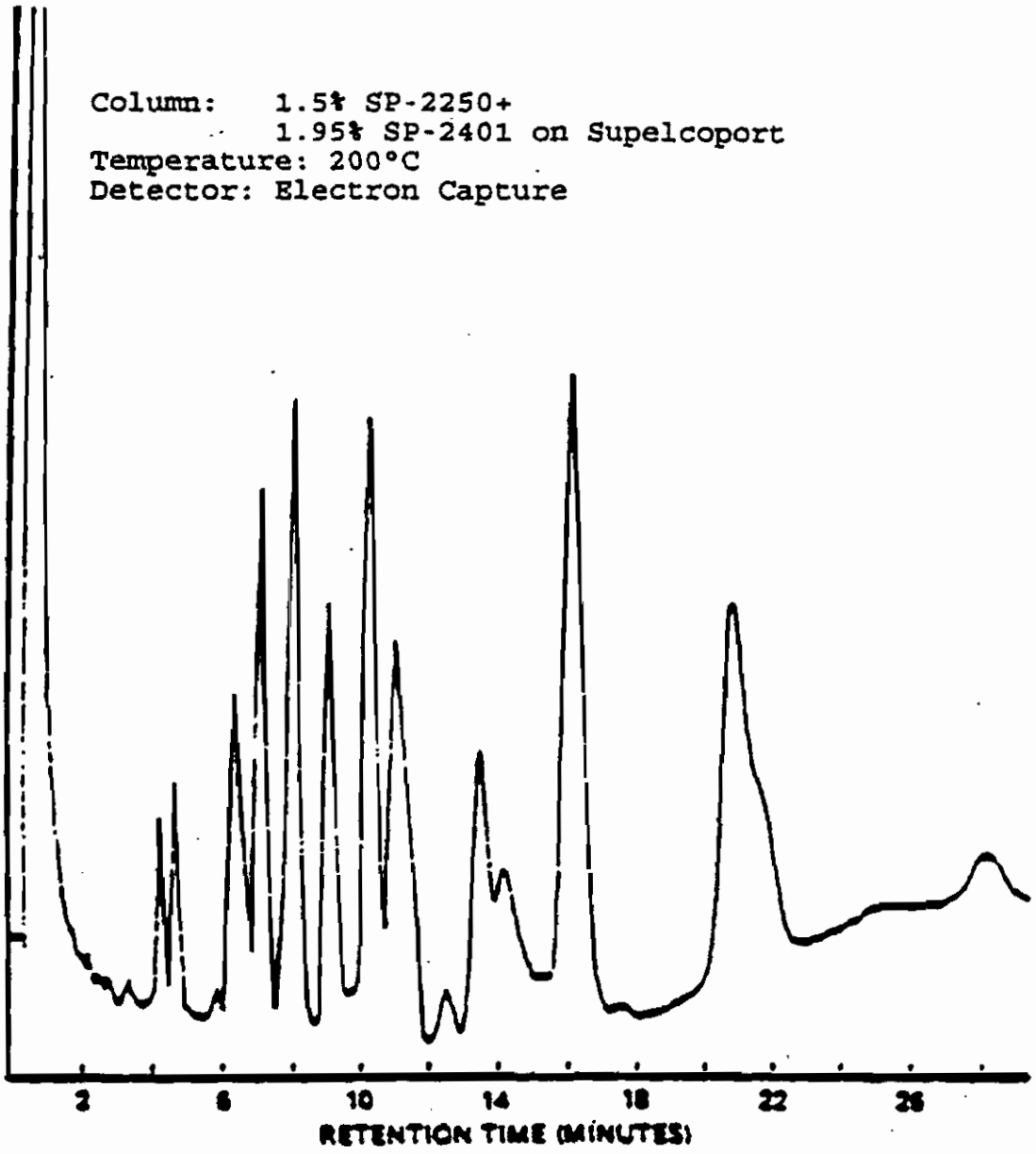


Figure 6

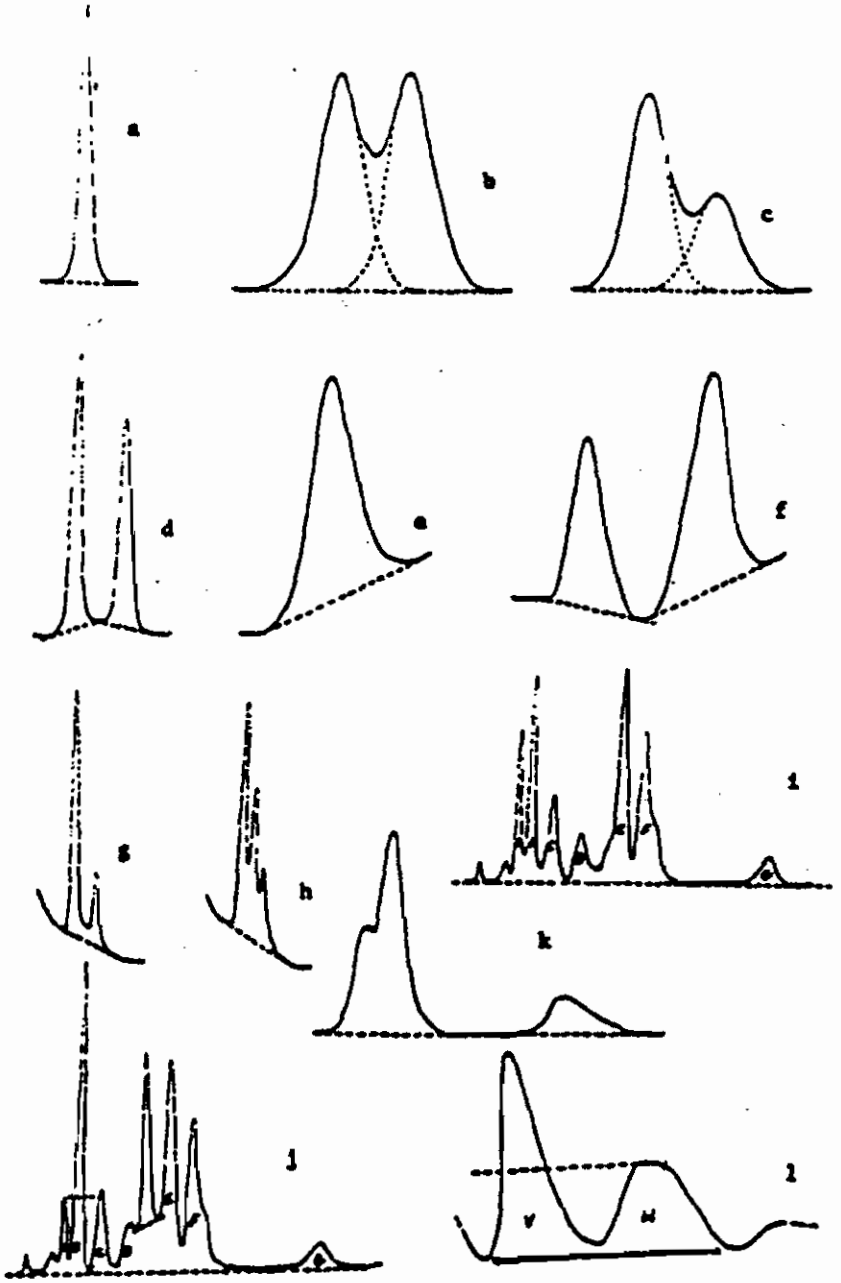


Fig.6--Baseline construction for some typical gas chromotagraphic peaks. a: symmetrical separated flat baseline; b and c: overlapp flat baseline; d: separated (pen does not return to baseline between peaks); e: separated sloping baseline; f: separated (pen goes below baseline between peaks); g: α- and γ-BHC sloping baseline; h: α-,β- and γ-BHC sloping baseline; i: chlordane flat baseline; j: heptachlor and heptachlor epoxide superimposed on chlordane; k: chair-shaped peaks, unsymmetrical peak; l: p,p'-DDT superimposed on toxaphene.

Figure 7

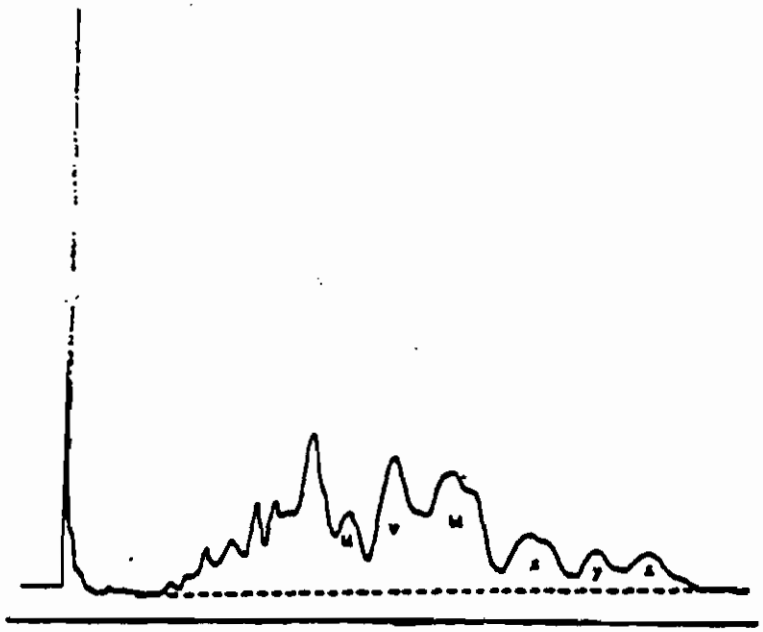


Fig.- 7a -- Baseline construction for multiple residues with standard toxaphene.

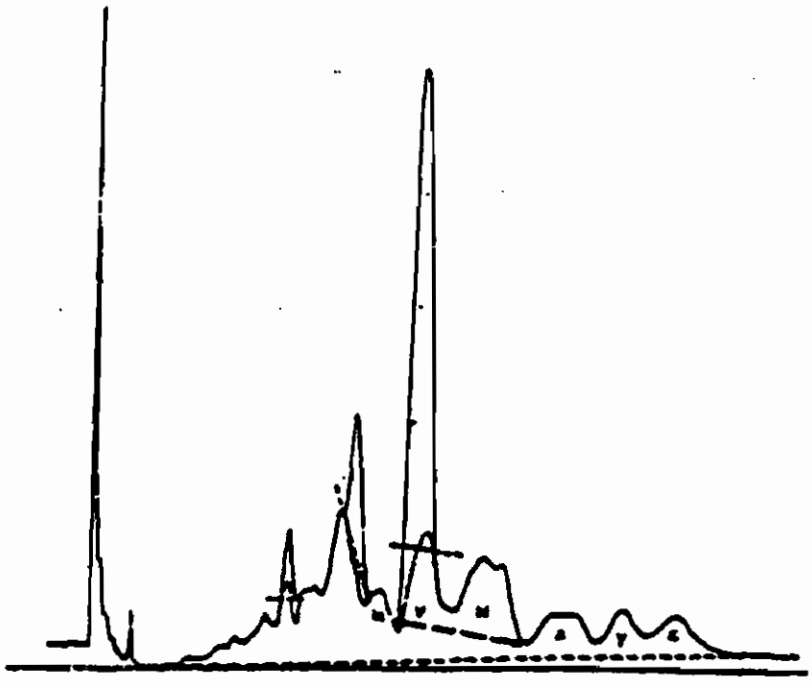


Fig.- 7b -- Baseline construction for multiple residues with toxaphene, DDE and o,p'-, and p,p'-DDT

Figure 8

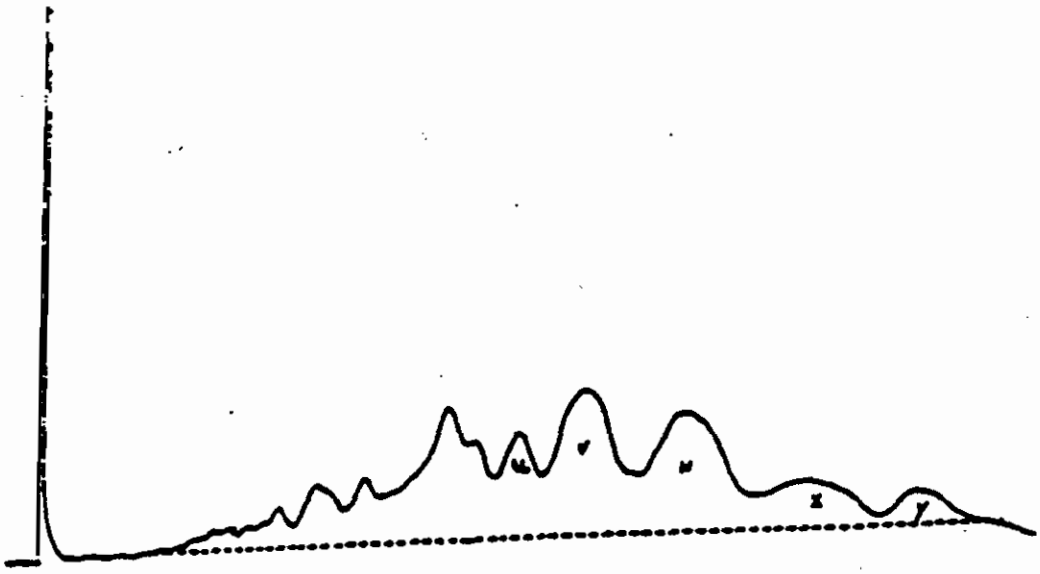


Fig.- 8a -- Baseline construction for multiple residues: standard toxaphene.

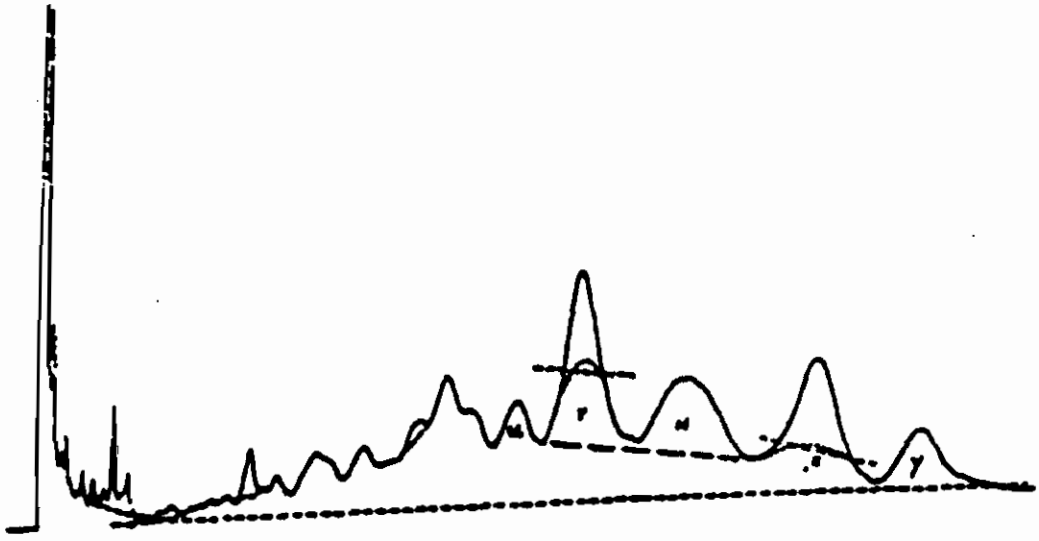


Fig.- 8b -- Baseline construction for multiple residues: rice bran with BHC, toxaphene, DDT, and methoxychlor.

Figure 9

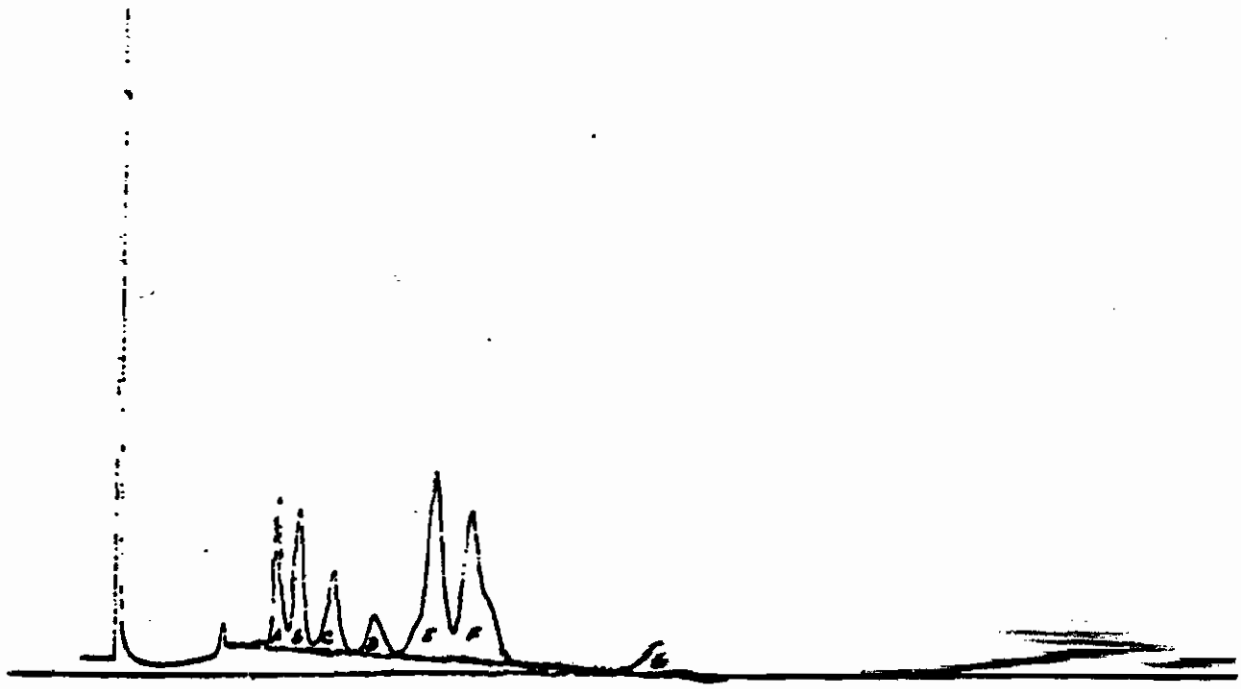


Fig.- 9a -- Baseline construction for multiple

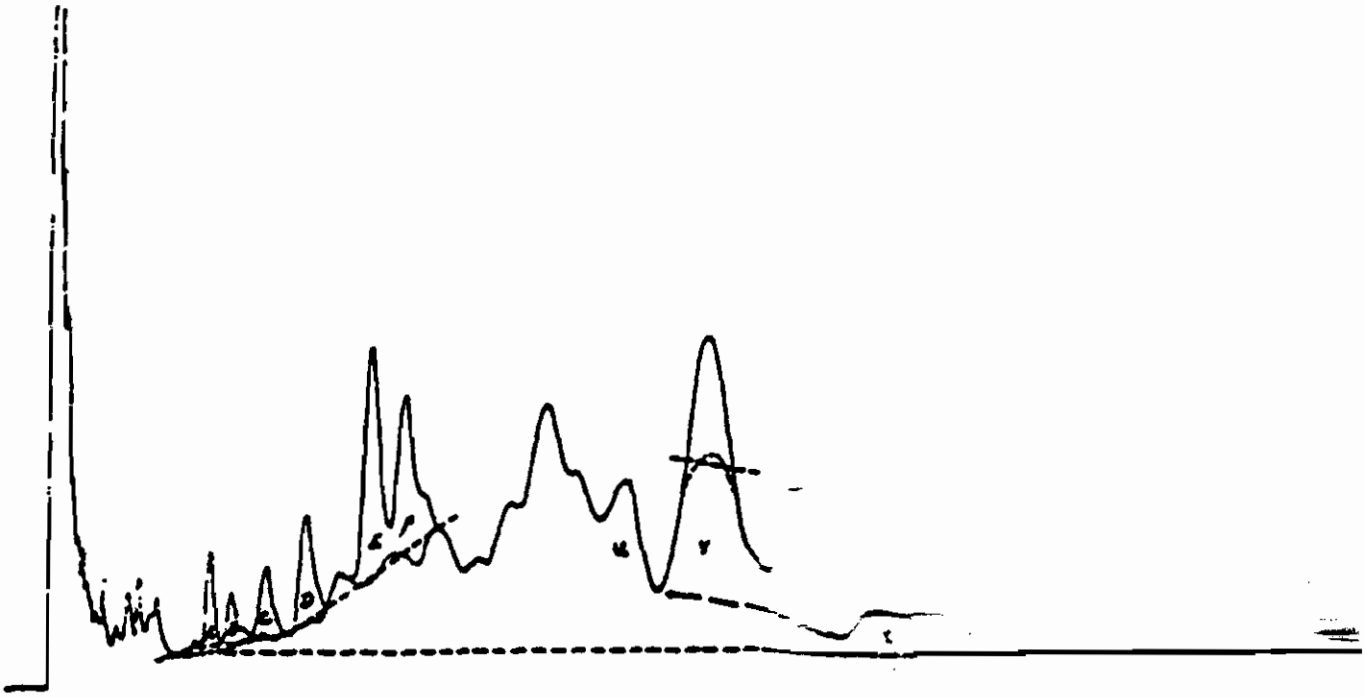
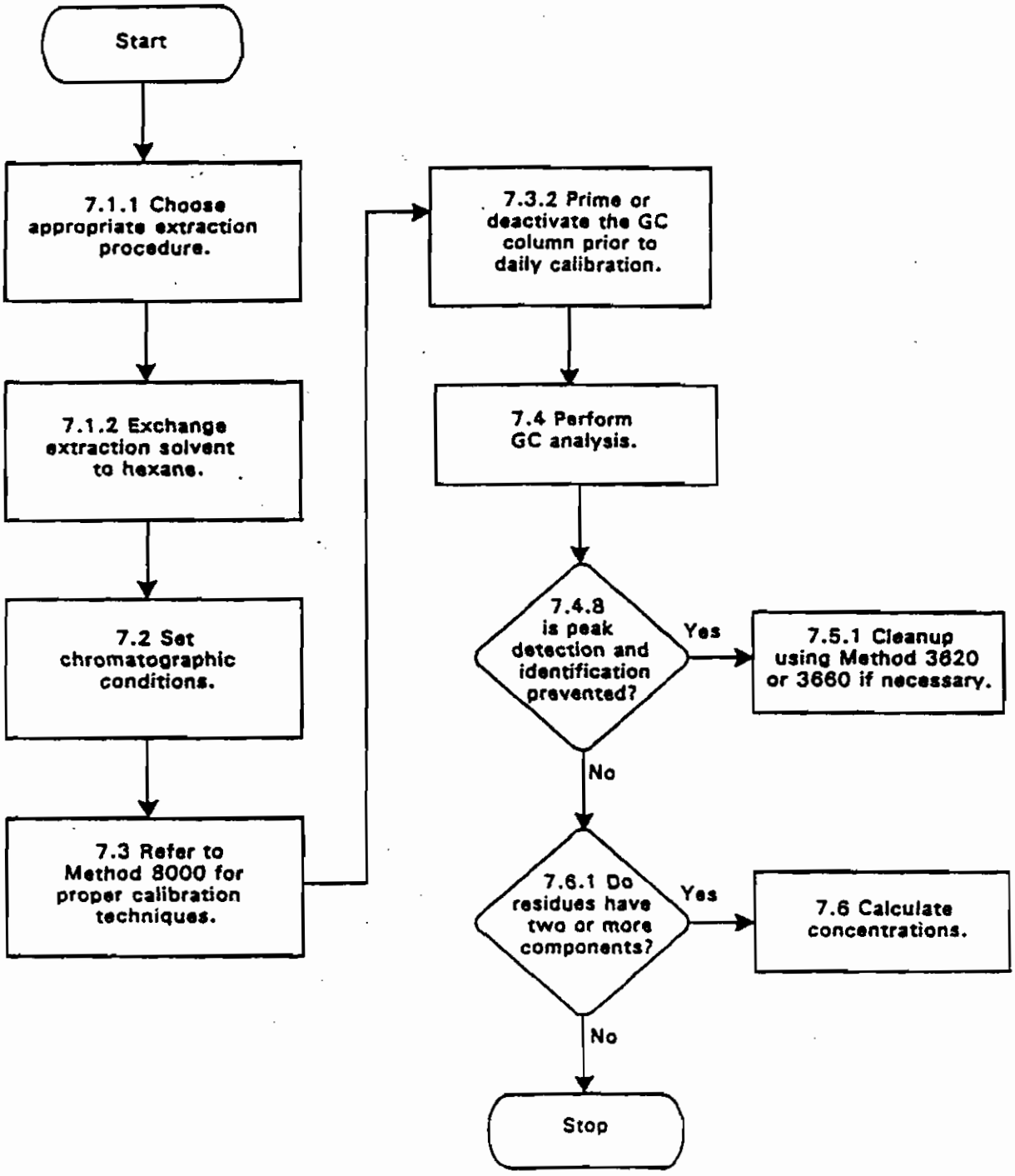


Fig.- 9b -- Baseline construction for multiple
chlordane, toxaphene, and


Start
7.1.1 Choose appropriate extraction procedure.
7.1.2 Exchange extraction solvent to hexane.
7.2 Set chromatographic conditions.
7.3 Refer to Method 8000 for proper calibration techniques.
7.3.2 Prime or deactivate the GC column prior to daily calibration.
7.4 Perform GC analysis.
7.4.8 is peak detection and identification prevented?
Yes
7.5.1 Cleanup using Method 3620 or 3660 if necessary.
No
7.6.1 Do residues have two or more components?
Yes
7.6 Calculate concentrations.
No
Stop

APPENDIX C

NEW YORK STATE DEPARTMENT OF HEALTH

METHOD 311-1

NEW YORK STATE DEPARTMENT OF HEALTH
WADSWORTH CENTER FOR LABORATORIES AND RESEARCH
ALBANY, N.Y. 12201

10/6/81

POLYCHLORINATED BIPHENYLS IN AMBIENT AIR

1. Scope and Application

1.1 This method covers the determination of polychlorinated biphenyls (PCBs) as Aroclors 1016/1242, 1221, 1254, 1260 in ambient air.

1.2 A 24 hour sample containing Aroclor 1016 collected at approximately 1.0 liter per minute is efficiently trapped in one cartridge, even at 95% relative humidity.

1.3 The minimum detectable concentration is 0.033 ug/1.5 ml of extract which is approximately 0.02 ug/cubic meter when sampled for 24 hours at 1 to 2 liters per minute.

2. Summary of Method

2.1 Polychlorinated biphenyls are quantitatively trapped on florisil in a cartridge. The PCB is desorbed by extraction with hexane. Sample clean-up is performed, if necessary. Analysis for PCBs is carried out on a gas chromatograph with a 3% SE-30 on GAS-CHROM Q column and electron capture detector. Results are reported in ug/cubic meter .

3. Interferences

3.1 Solvents, glassware, and other sample processing hardware may yield discrete artifacts and/or elevated base lines, causing misinterpretation of gas chromatograms. All of these materials must be demonstrated to be free from interferences under the conditions of the analysis. Specific selection of reagents and purification of solvents by distillation in all-glass systems may be required. Refer to Section 510-1, Analytical Handbook.

3.2 Organophosphorus pesticides - a number of organophosphorus pesticides, such as those containing a nitro group, e.g., parathion, also respond to the electron capture detector and may interfere with the determination of the organochlorine pesticides. Such compounds can be identified by their response to the flame photometric detector.

3.3 Interferences from chlorinated pesticides can be removed, if necessary, according to the method of Trotter.

4. Apparatus and Materials

4.1 Pasteur pipets - no. P5180-2, 5-3/4 inch length

4.2 Pasteur pipet bulb

4.3 Air or nitrogen - for evaporating sample

4.4 Vacuum pump - capable of maintaining an air pressure differential greater than 0.8 atmosphere at the desired flow rate

4.5 Muffle furnace - equipped with temperature control and indicator

4.6 Gas chromatograph:

4.6.1 Electron capture detector

4.6.2 Column dimensions - 6' X 1/4"

4.6.3 3% SE-30 on GAS-CHROM Q, 80/100 mesh, 2 mm I.D., Conditioned at 220 C.

4.6.4 Alternate column - 1.5% OV17/1.95% QF-1, on GAS-CHROM Q,80/100 mesh

4.7 Test tubes - graduated, ground-glass stoppered, 5 or 10 ml size

4.8 Tygon tubing - 1/4" I.D., 3/8" O.D. for cartridge connections

4.9 Teflon tubing - 7 mm I.D.

4.10 Caplugs - size 9/32 in SC (Caplugs Division, Protective Closures Co. Inc., 2150 Elmwood Ave., Buffalo, NY)

4.11 Wrist Action Shaker, Burrell.

5. Reagents, Solvents and Standards

5.1 Florisil - 60-100 mesh, Fisher Scientific Co., #F-100. Wash Florisil with several portions of hexane and dry on a steambath to remove hexane residue. Place Florisil in a muffle furnace at 320 C overnight. Deactivate the Florisil by shaking 100 grams with 2 ml distilled water for 15 minutes on a mechanical wrist action shaker. Store Florisil in a screw-cap jar.

5.2 Glass wool - hexane rinsed

5.3 Hexanes - nanograde, Mallinckrodt, no. 4159

5.4 PCB standards - (Monsanto) Aroclor 1016, 1221, 1254,1260. Other Aroclors may be of interest, these are the most common. For each Aroclor of interest prepare the following dilute standards:

5.4.1 1 mg/ml stock PCB solution - weigh 100 mg of PCB and place in a 100 ml volumetric flask. Dilute to the mark with hexane.

5.4.2 10 ug/ml PCB solution - dilute 1.0 ml of stock PCB solution to 100 ml with hexane in a volumetric flask.

5.4.3 2.0 ug/ml PCB solution - dilute 1.0 ml of 10 ug/ml PCB solution to 5 ml with hexane in a volumetric flask.

5.4.4 1.0 ug/ml PCB solution - dilute 1.0 ml of 10 ug/ml PCB solution to 10 ml with hexane in a volumetric flask.

5.4.5 0.4 ug/ml PCB solution - dilute 1.0 ml of 10 ng/ml PCB solution to 25 ml with hexane in a volumetric flask.

5.4.6 4.0 ug/ml PCB solution - dilute 2 ml of 10 ug/ml PCB solution to 5 ml with hexane in a volumetric flask.

5.4.7 0.04 ug/ml PCB solution - dilute 0.1 ml of 10 ug/ml PCB solution to 25 ml with hexane in a volumetric flask.

6. Quality Control

6.1 All reagents must be checked for contamination by GLC prior to use.

6.2 At least one blank is analyzed with each batch of cartridges or samples.

7. Procedure

7.1 Preparation of Sampling Cartridges

NOTE: If a large number of cartridges are to be prepared, it is recommended that a particulate mask and disposable plastic gloves be used.

7.1.1 All glassware must be free of PCB. Rinse all glassware with at least 3 separate portions of hexane. Bake glassware for two hours at 350 C. Collect a final hexane rinse and concentrate to 0.5 ml for analysis to verify that all interfering chromatogram peaks are removed.

7.1.2 Cut 1" to 1 1/4" from the finely drawn end of a Pasteur pipet using a triangular file. Fire polish the cut end.

7.1.3 Place a piece of hexane-rinsed glass wool into a Pasteur pipet. Push it down to the pointed end of the pipet. To the pipet, add about 0.4 grams of florisil. Place a small plug of hexane-rinsed glass wool at the top of the Florisil. Prepare several sampling cartridges in this manner, including a sufficient number to be used as blank cartridges.

7.1.4 After construction of the cartridge, attach the cartridge outlet to a vacuum hose, keeping the cartridge vertical. Turn on the vacuum pump and tap the side of the cartridge lightly for about five seconds to settle and pack the Florisil. Push the glass wool plug down to the surface of the Florisil to insure close contact. This packing procedure eliminates the need to keep the cartridge vertical during sampling. Store the cartridges in an air-tight container.

7.1.5 Assemble cartridge into a sampling train as shown in Figure 1. One cartridge is usually sufficient to form an efficient sampling train. If a train containing two cartridges in series is desired, connections should be made with a short piece of Teflon tubing, butting the glass tubes, and using Tygon sleeving to prevent leaks. Measure and record the flow rate. Seal the ends of the cartridge train with aluminum foil caps or Caplugs to prevent contamination.

7.1.6 If at all possible, an inlet probe should not be used. If this cannot be avoided, a glass probe should be used. Teflon and Tygon have been found to adsorb PCBs and therefore are not recommended.

7.1.7 The sample should be collected for an appropriate period of time, usually 24 hours. The cartridge train should be resealed with aluminum foil or Caplugs before transporting back to the laboratory.

7.2 Analysis of Florisil Cartridges

7.2.1 After flow rate measurement, disassemble the sampling train. Label the cartridges.

7.2.2 Elute the PCBs into a graduated 5 or 10 ml test tube with a ground glass stopper using about 5 ml of nanograde hexane, adding the hexane to the inlet of the cartridge. Collect the eluate in the graduated test tube.

7.2.3 Concentrate the extract by nitrogen (HP) stream evaporation to bring the PCB concentration of the extract into working range, usually 1 to 5 ml volume. Place extract into 2 ml capacity vials with injectable caps for use with automated gas chromatograph sampler.

7.2.4 Run samples and standards on the gas chromatograph under the following conditions:

7.2.4.1 Column temperature - 180 C isothermal for Aroclors 1016/1242,1221,1254 and 1260.

7.2.4.2 Detector temperature - electron capture detector, 275 C

7.2.4.3 Inlet temperature - 200 C

7.2.4.4 Carrier gas - nitrogen, 40 ml/min

7.2.4.5 Carrier flow rate - 60 cc/min

7.2.4.6 Injection size - 5 ul for manual injection; 1 ul for automated injection.

7.2.5 Blank cartridge content should be subtracted from sample cartridges to determine net Aroclor content. Amounts greater than 0.033 ug Aroclor in the blank cartridge extracts may indicate contamination of the cartridge train either before of after sampling (i.e., glassware contamination).

7.2.6 Peaks should be quantitated by area comparisons under specific Aroclor peaks of standards of known amounts.

8. Calculations

8.1 Air Volume Sampled

$$V = RT/1000$$

where, V = volume of air sampled in cubic meters
R = air flow rate in liters per minute
T = time sampled in minutes

8.2 ug PCB Collected: Read from standard curve or by proportion to standard on day of run.

$$\text{ug PCB} = (A-B)\ C$$

where, A = ug/ml PCB in sample extract
B = ug/ml PCB in blank extract (usually is zero)
C = ml of hexane extract

8.3 Report results as ug/cubic meter in air sampled. Report as either total PCB or specific Aroclor.

9. References

9.1 Dell'Acqua, B. NYSDH, In-house experiment.

9.2 Giam, C.S., Chan, H.S., Neff, G.S. 1975. Rapid and Inexpensive Method for Detection of Polychlorinated Biphenyls and Phthalates in Air. Analytical Chemistry 47(13): 2319-20.

9.3 Trotter, W.J. 1973. Removing the Interference of DDT and its Analogs in the Analysis for Residues of Polychlorinated Biphenyls. Journal of the Association of Official Analytical Chemists 58(3): 461-5.

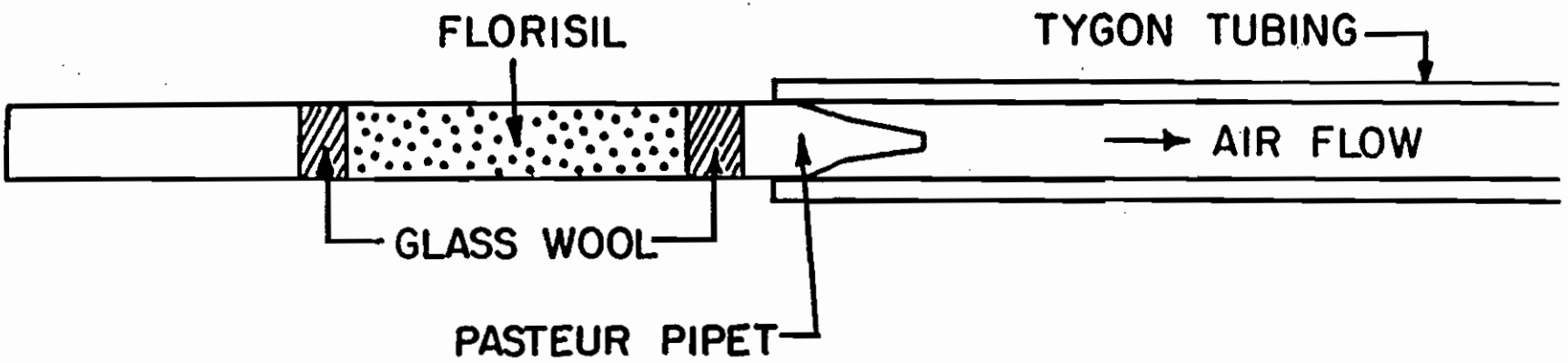
9.4 Determination of Polychlorinated Biphenyls in Ambient Air, NYSDH, Division of Laboratories and Research, Environmental Health Institute, APC method 26., June 1979.

9.5 NIOSH Health and Hazard Evaluation Report No. 86-472; p. 18, 19; HETA86-472-1832; Commercial Office Buildings, Boston, MA.

HANDBOOK5(311-1)
July 1980; Corrected January 14, 1981; Corrected March 12, 1981

FIGURE 1

FLORISIL CARTRIDGE



SUNY New Paltz
Wipe Sample Locatio

Bliss Hall, Building #22
Basement
B4 - Floor Area
First Floor
103 - Floor Area
119 - Desk
Second Floor
205 - Desk
216 - Floor Area
Third Floor
326 - Floor Area
309 - Floor Area

Gage Hall, Building #21
Basement East
B8- Floor Area
First Floor East
105 - Desk
RD's Apartment - Floor Area
First Floor West
117 - Floor Area
133 Overflow - Desk
Second Floor East
218 - Desk
221 - Floor Area
Third Floor East
302 - Floor Area
Third Floor West
343 - Desk
328 - Desk

Scudder Hall, Building #20
First Floor
115 - Desk
116 - Desk
112 - Desk
Second Floor
205 - Floor Area
216 - Desk
230 - Desk
Third Floor
306 - Desk
309 - Floor Area

Capen Hall, Building #9
First Floor
110 - Floor Area
Second Floor
220 - Floor Area
Third Floor
323 - Desk
318 - Floor Area

SUNY New Paltz
PCB Air Sample Locations

Bliss Hall, Building #22
First Floor
112 Overflow
114
Second Floor
214
218
230
203
223

Gage Hall, Building #21
First Floor East
112
Second Floor East
223
203
Second Floor West
227
231
225
Third Floor East
302
305
Third Floor West
325

Scudder Hall, Building #20
Basement
B3
First Floor
107
Second Floor
209
222
Third Floor
320
322
328

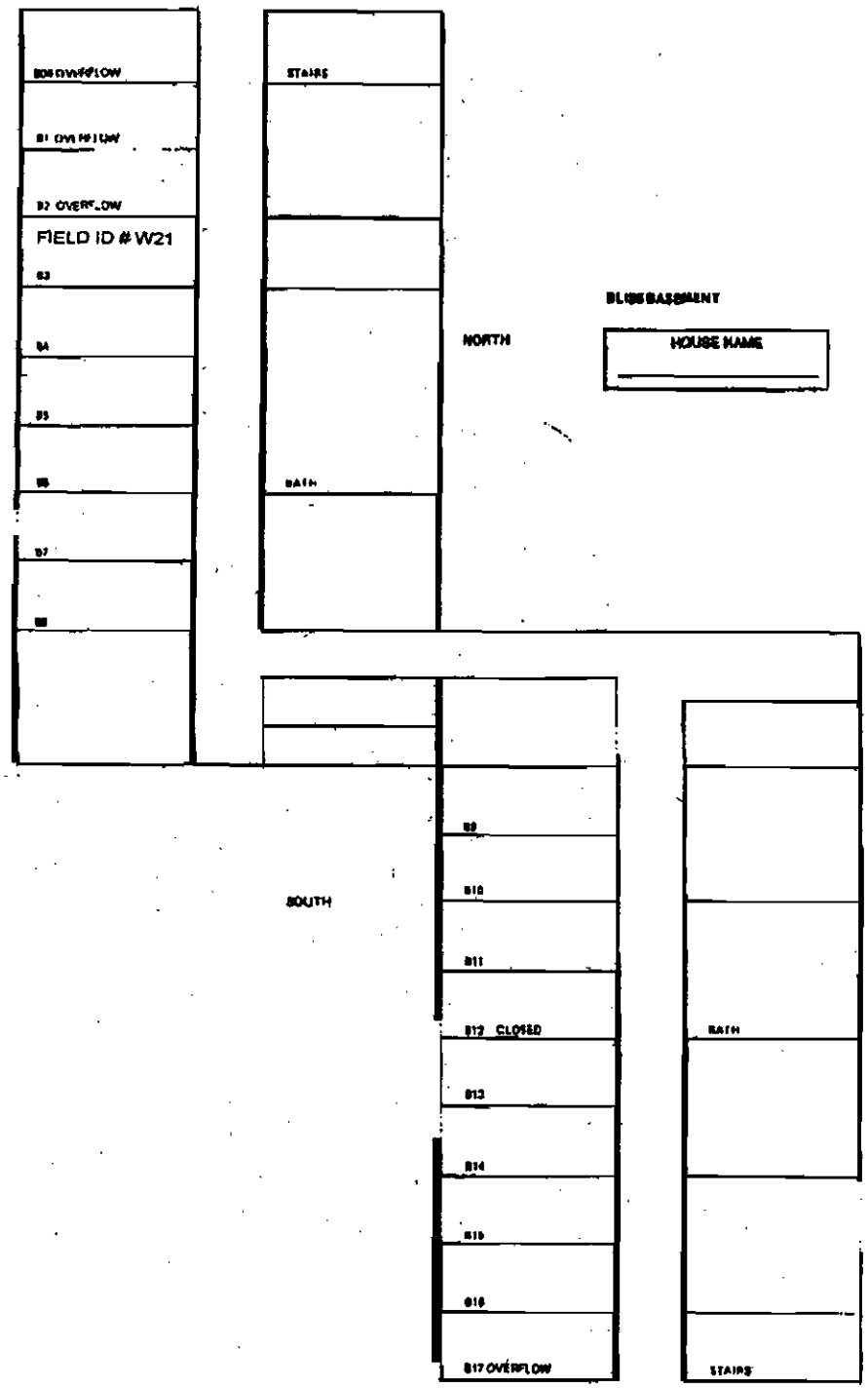
Capen Hall, Building #9
First Floor
121
Second Floor
217
226
Third Floor
302
311
315

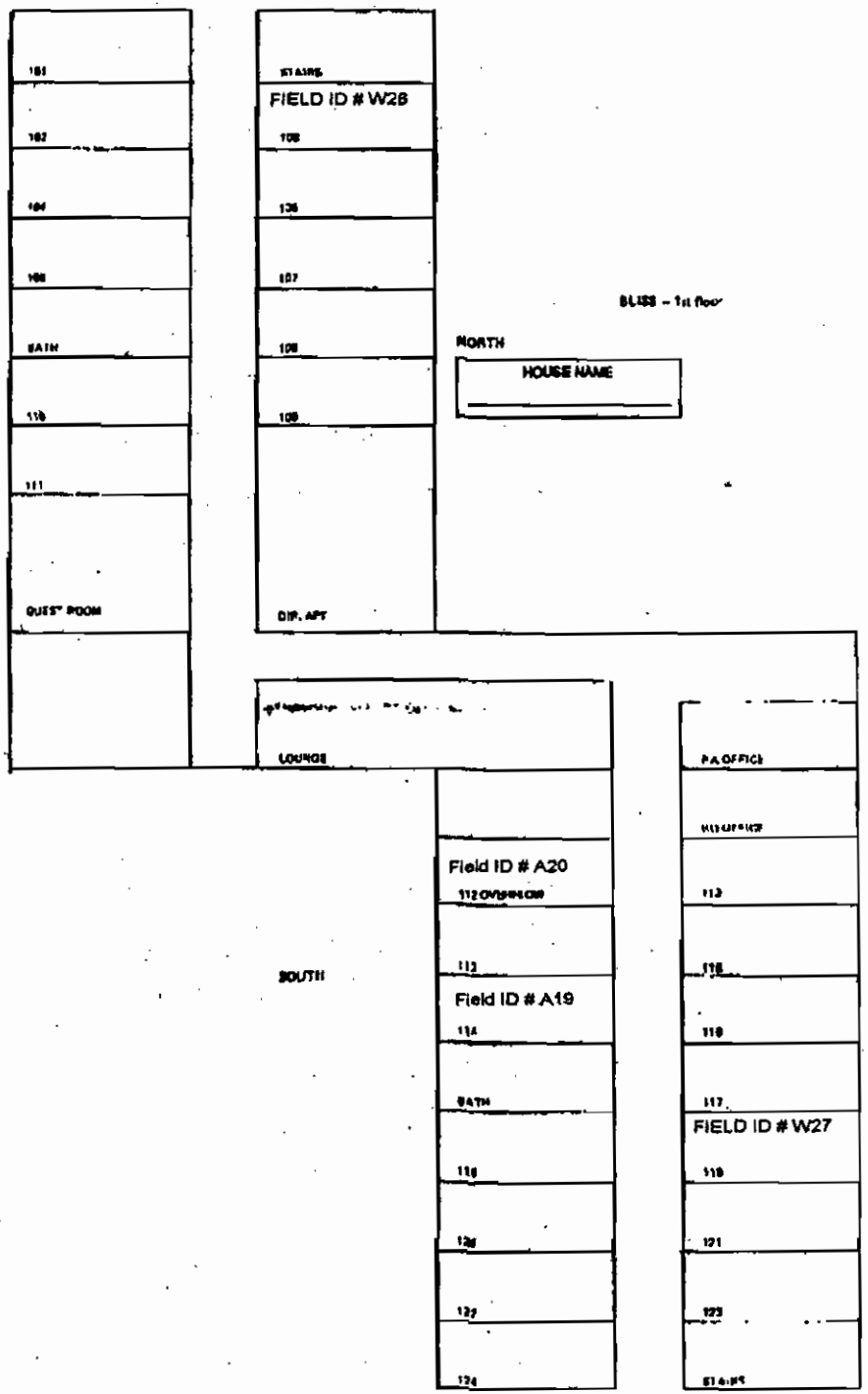
Table 1
SUNY New Paltz
Wipe Sample Locations

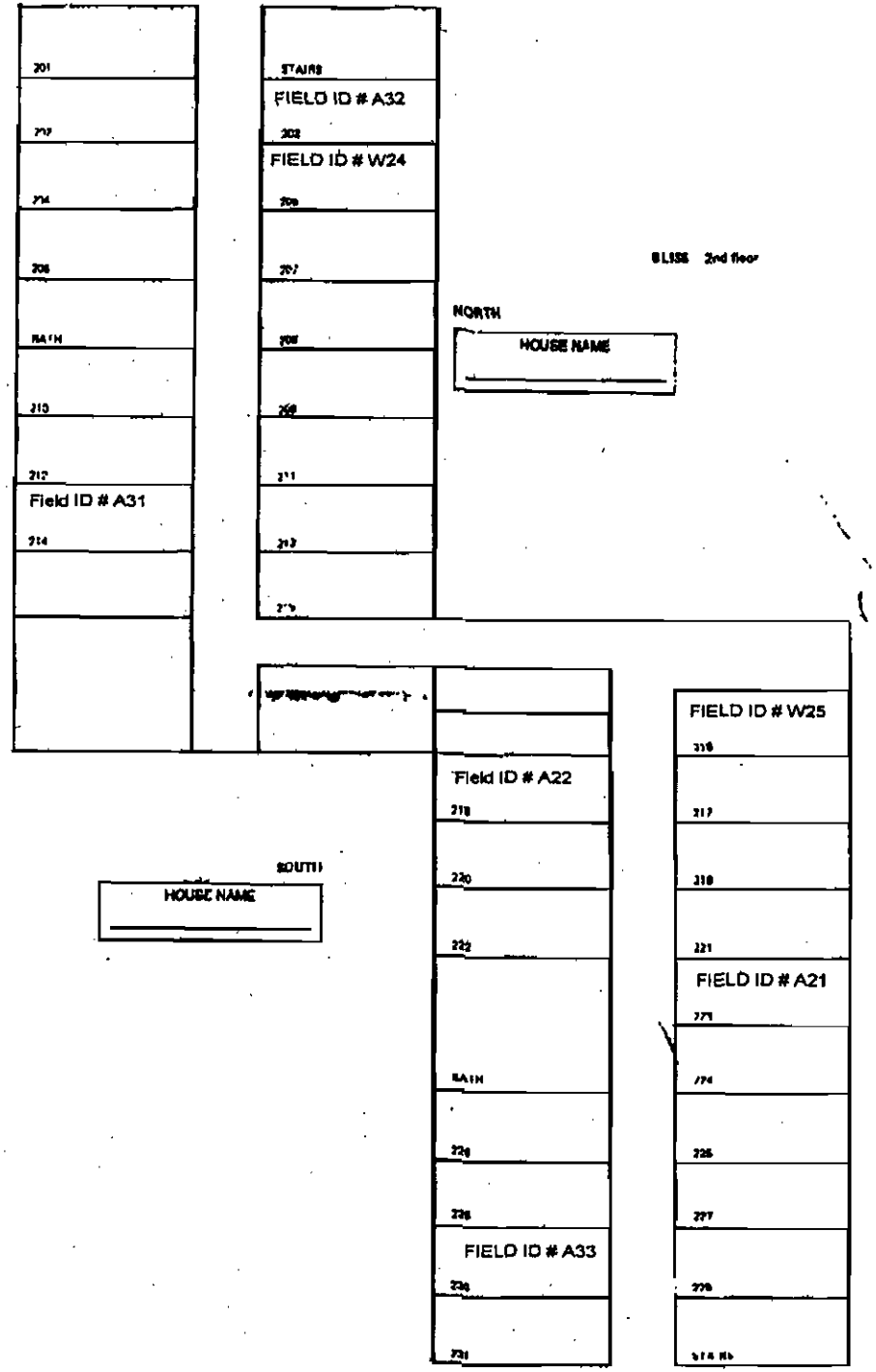
Sample Location		Field ID Number
Bliss Hall, Building #22		
Basement:	B4 – Floor Area	W21
First Floor:	103 – Floor Area	W26
	119 – Desk	W27
Second Floor:	205 – Desk	W24
	216 – Floor Area	W25
Third Floor:	326 – Floor Area	W23
	309 – Floor Area	W22
Gage Hall, Building #21		
Basement East:	B8 – Floor Area	W19
First Floor East:	105 – Desk	W17
	RD's Apartment – Floor Area	W17
First Floor West:	117 – Floor Area	W16
	133 Overflow – Desk	W15
Second Floor East:	218 – Desk	W14
	221 – Floor Area	W13
Third Floor East	302 – Floor Area	W20
Third Floor West	343 – Desk	W12
	328 – Desk	W11
Scudder Hall, Building #20		
First Floor:	115 – Desk	W10
	116 – Desk	W9
	112 - Desk	W8
Second Floor:	205 – Floor Area	W7
	216 – Desk	W6
	230 – Desk	W5
Third Floor:	306 – Desk	W3
	309 – Floor Area	W4
Capen Hall, Building #9		
First Floor:	110 – Floor Area	W32
Second Floor:	220 – Floor Area	W31
Third Floor:	323 – Desk	W29
	318 – Floor Area	W28
Field Blank		W1
Field Blank		W2
Field Blank		W30

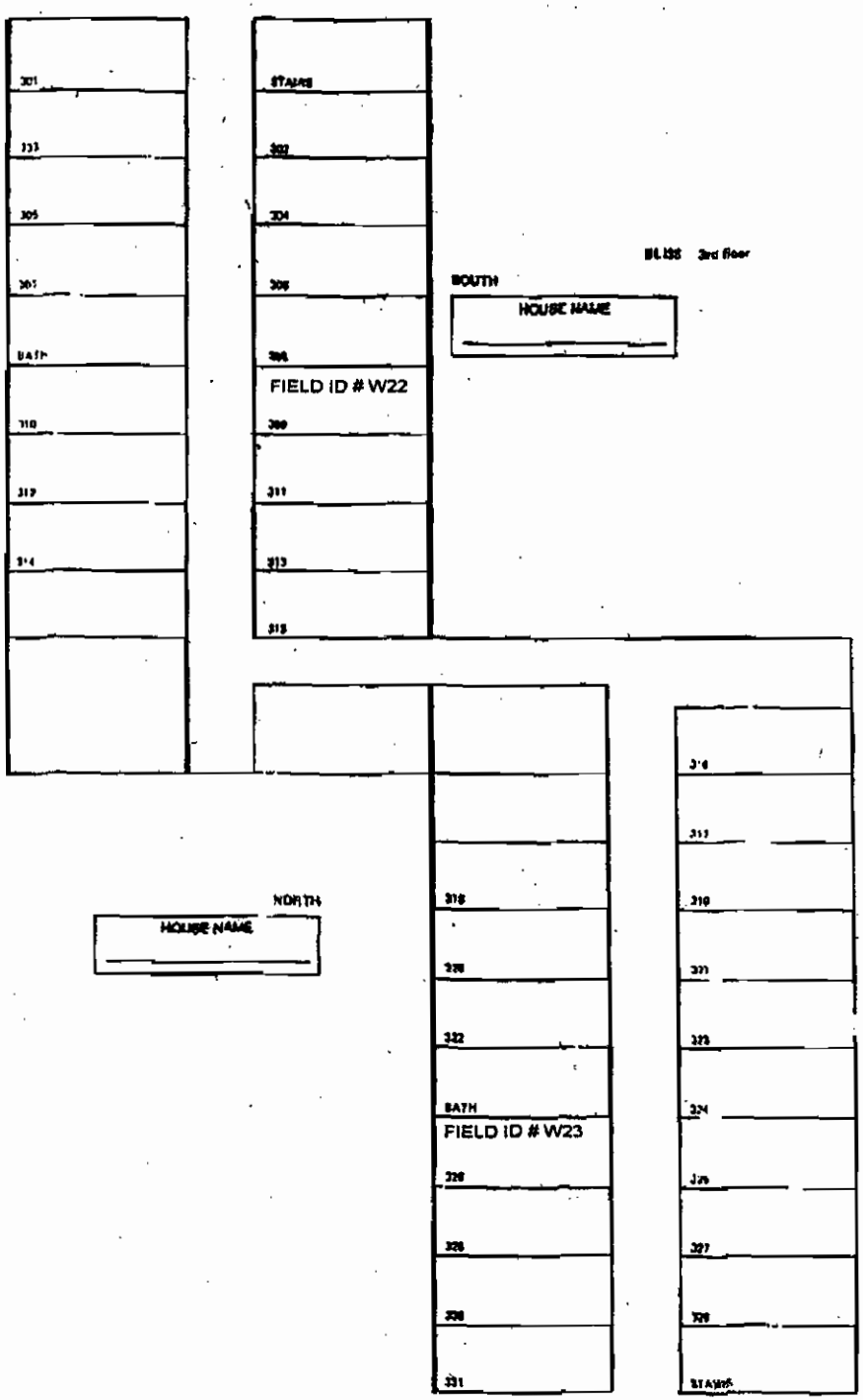
Table 2
SUNY New Paltz
PCB Air Sample Locations

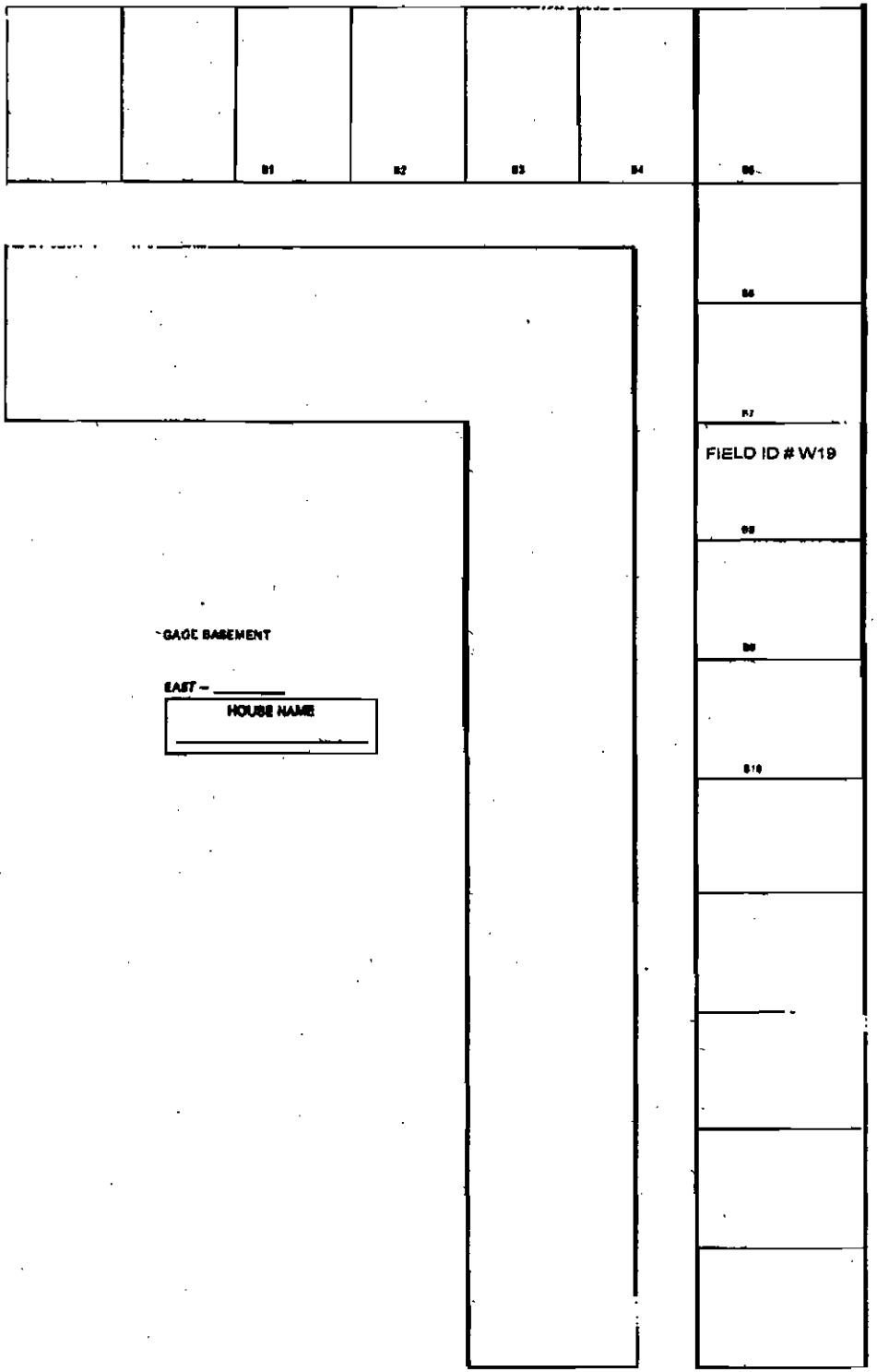
Sample Location		Field ID Number
Bliss Hall, Building #22		
First Floor:	112 Overflow	A20
	114	A19
Second Floor:	214	A31A
	218	A22
	230	A33
	203	A30
	223	A21
Gage Hall, Building #21		
First Floor East:	112	A30
Second Floor East:	223	A26
	203	A10
Second Floor West:	227	A27
	231	A29
	225	A28
Third Floor East:	302	A12
	305	A25
Third Floor West:	325	All
Scudder Hall, Building #20		
Basement:	*B3*	A3
First Floor:	*107*	A13
Second Floor:	*209*	A14
	222	A15
Third Floor:	*320*	A18
	322	A17
	328	A16
Capen Hall, Building #9		
First Floor:	121	A4
Second Floor:	217	A6
	226	A5
Third Floor:	302	A7
	311	A8
	315	A9
Field Blank		A1
Field Blank		A2
Field Blank		A23
Outside Gage Hall		A24


B08 OVERFLOW
B1 OVERFLOW
B2 OVERFLOW
FIELD ID # W21
B3
B4
B5
B6
B7
B8
STAIRS
BATH
NORTH
BLDG BASEMENT
HOUSE NAME
SOUTH
B9
B10
B11
B12 CLOSED
B13
B14
B15
B16
B17 OVERFLOW
BATH
STAIRS


FIELD ID # W28
NORTH
HOUSE NAME
BATH
QUES" ROOM
DIR. APT
LOUNGE
P.A.OFFICE
Field ID # A20
Field ID # A19
BATH
FIELD ID # W27
SOUTH


STAIRS
FIELD ID # A32
FIELD ID # W24
BATH
Field ID # A31
NORTH
HOUSE NAME
Field ID # A22
SOUTH
HOUSE NAME
FIELD ID # W25
FIELD ID # A21
BATH
FIELD ID # A33


BLISS 3rd floor
SOUTH
HOUSE NAME
FIELD ID # W22
NORTH
HOUSE NAME
BATH
FIELD ID # W23

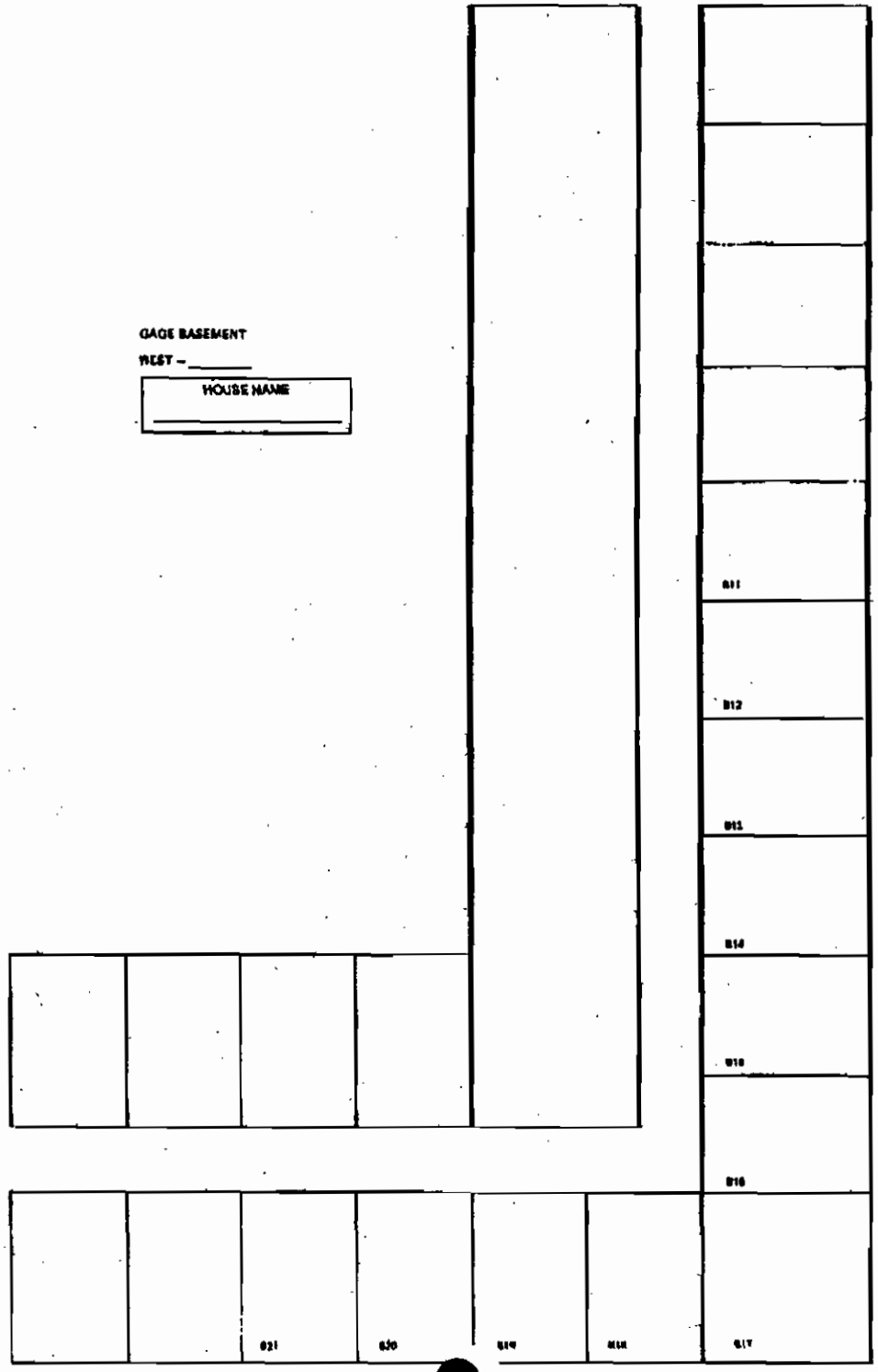

B1
B2
B3
B4
B5
B6
B7
FIELD ID # W19
B8
B9
B10
GAGE BASEMENT
EAST –
HOUSE NAME

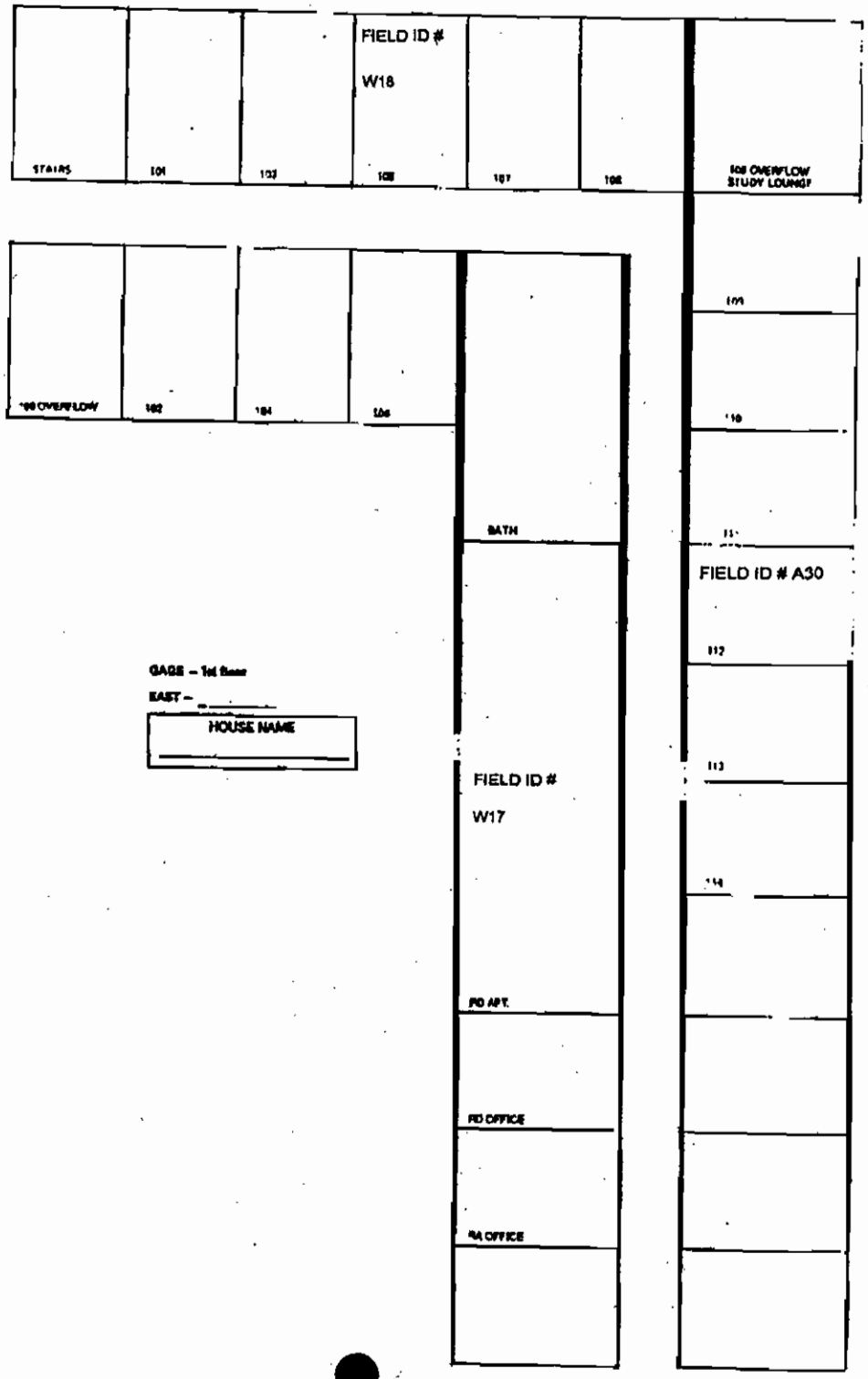
GAGE BASEMENT

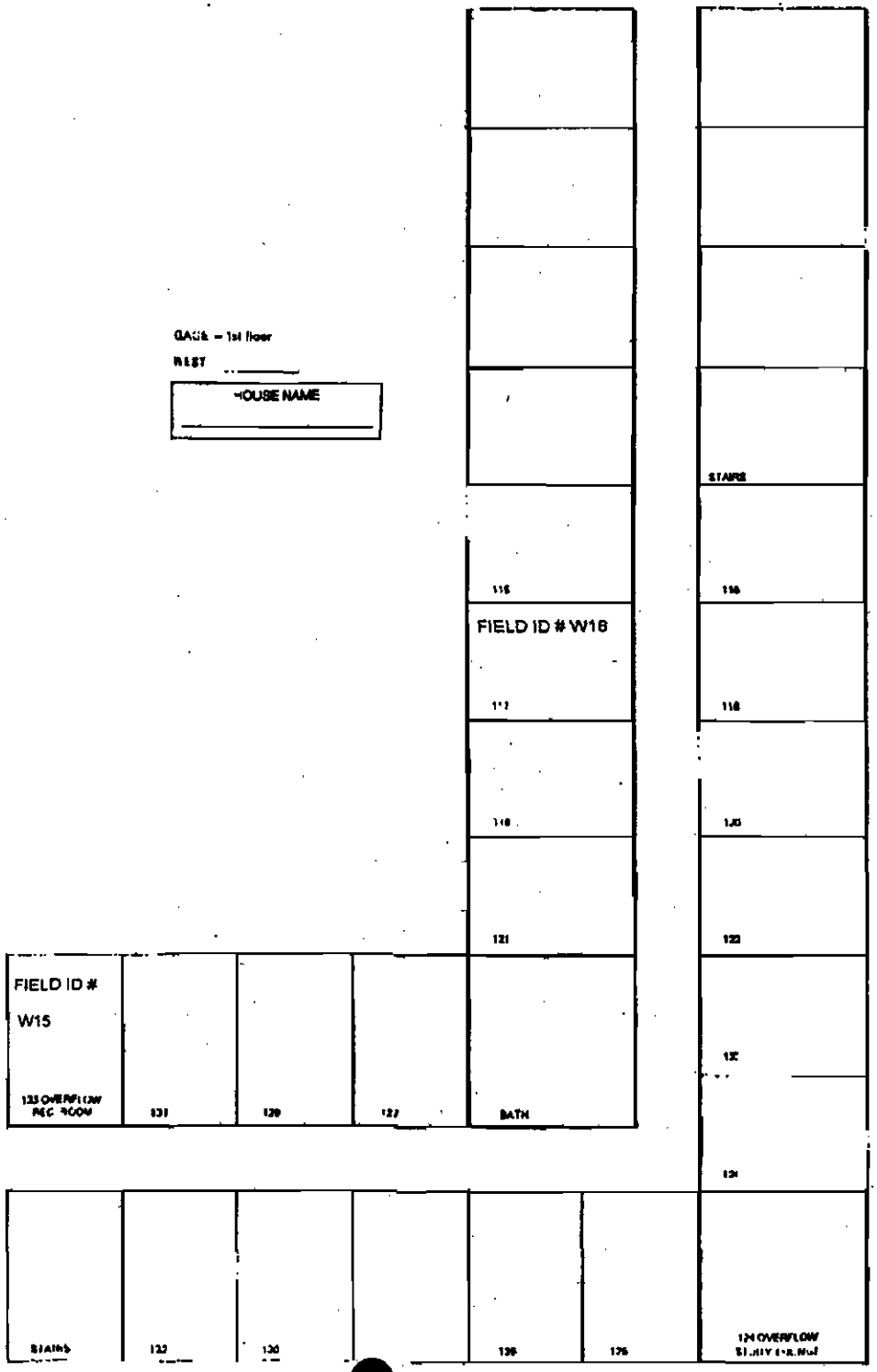
WEST – _____

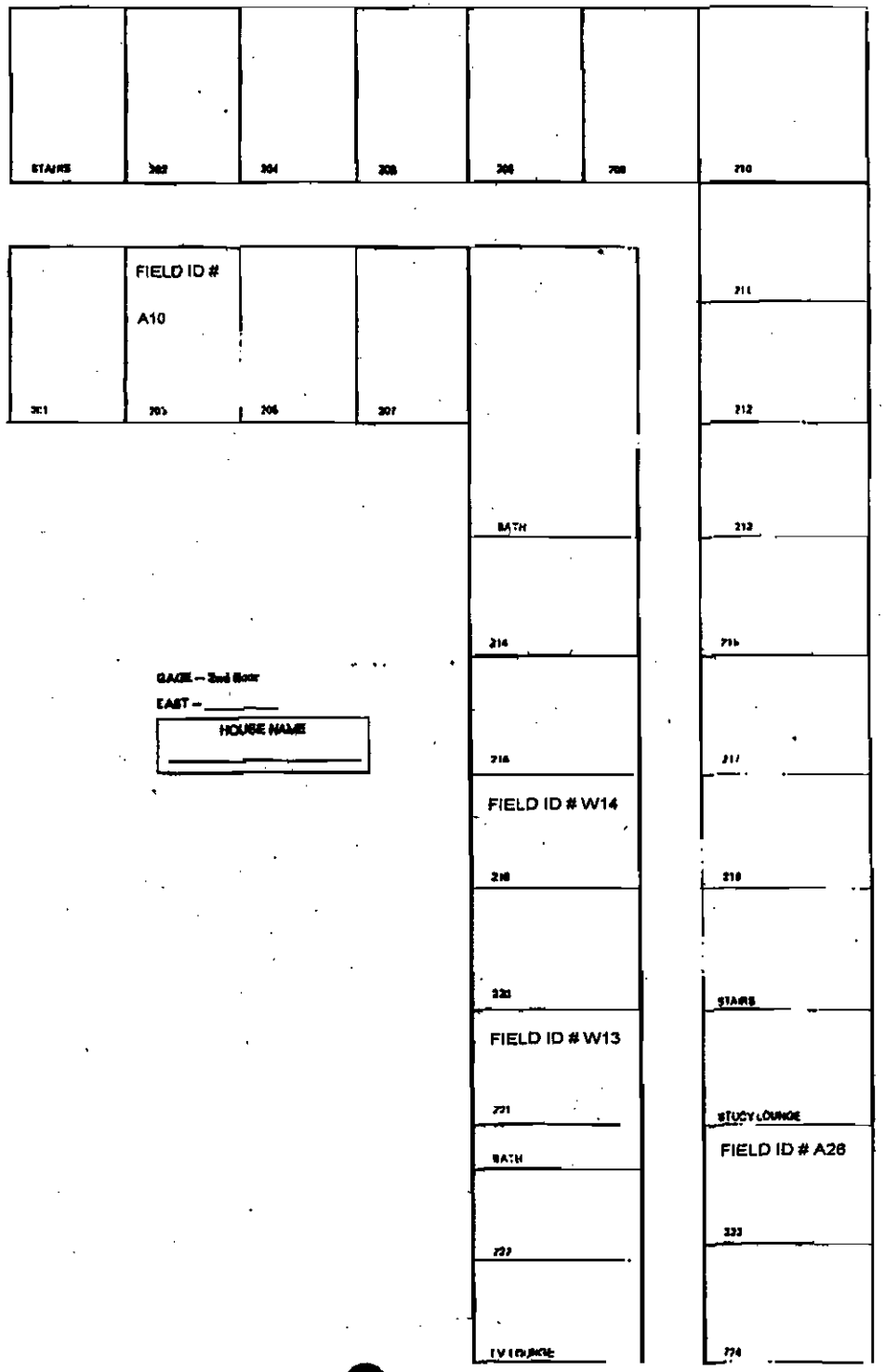
HOUSE NAME

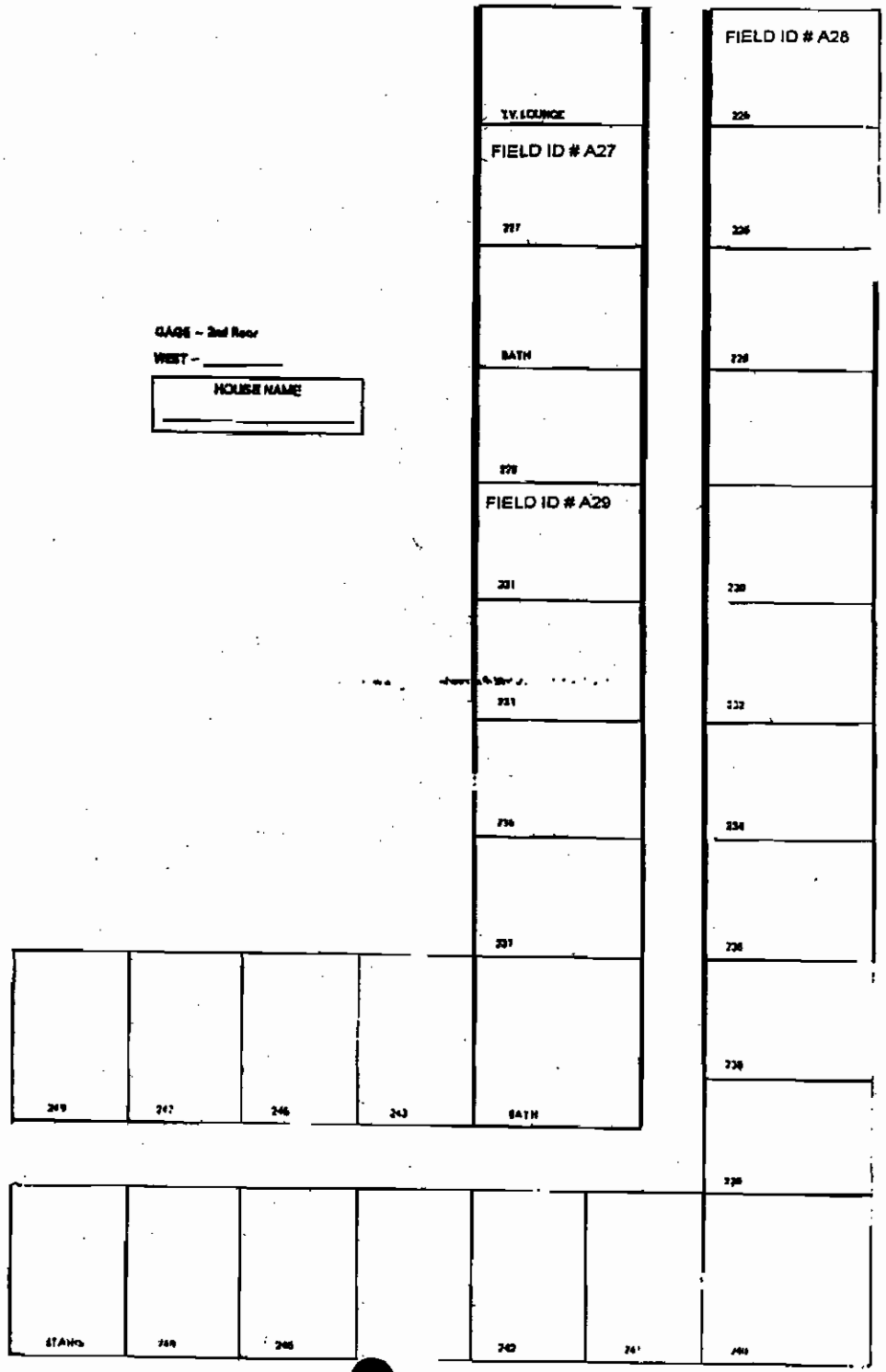
_____

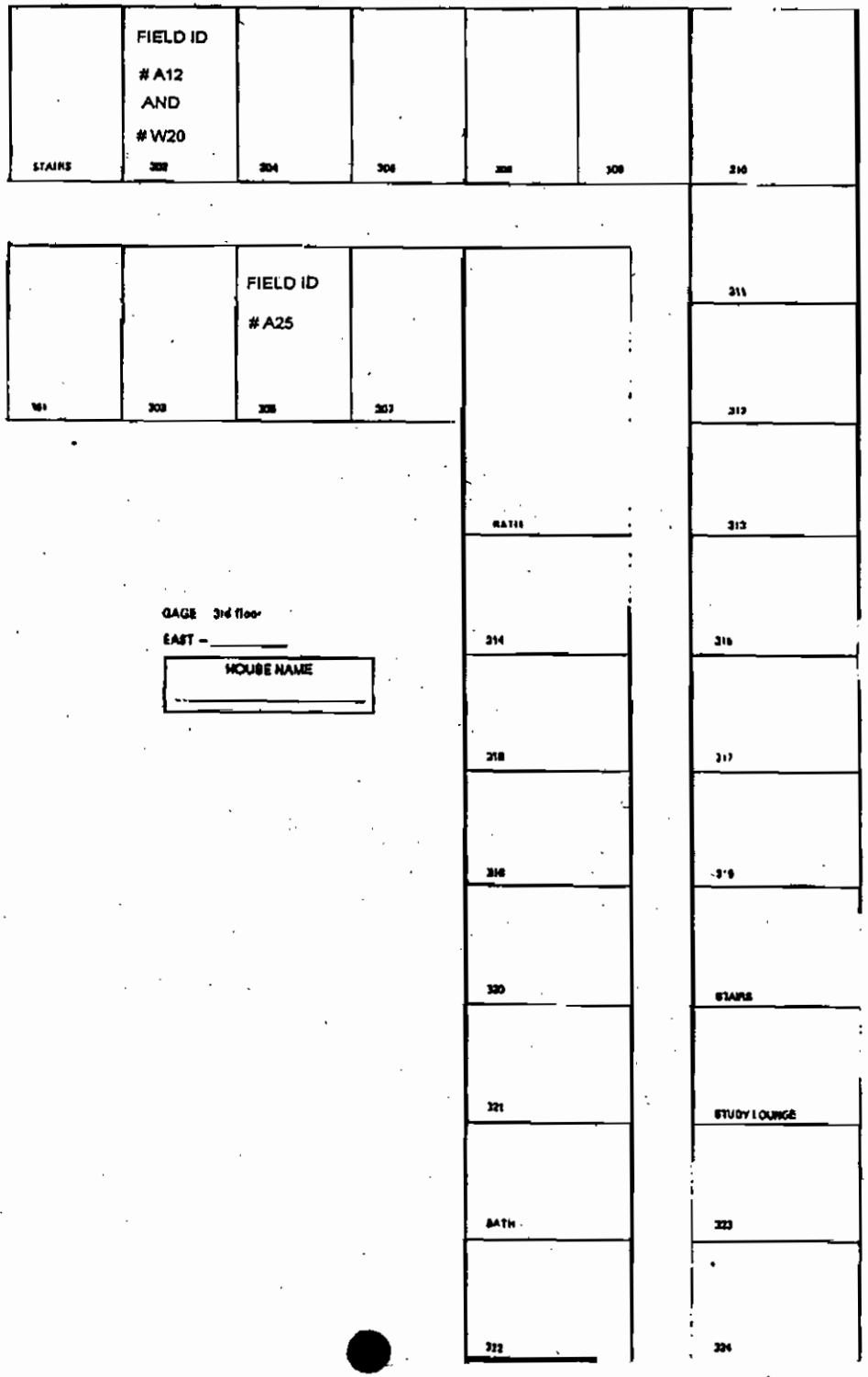


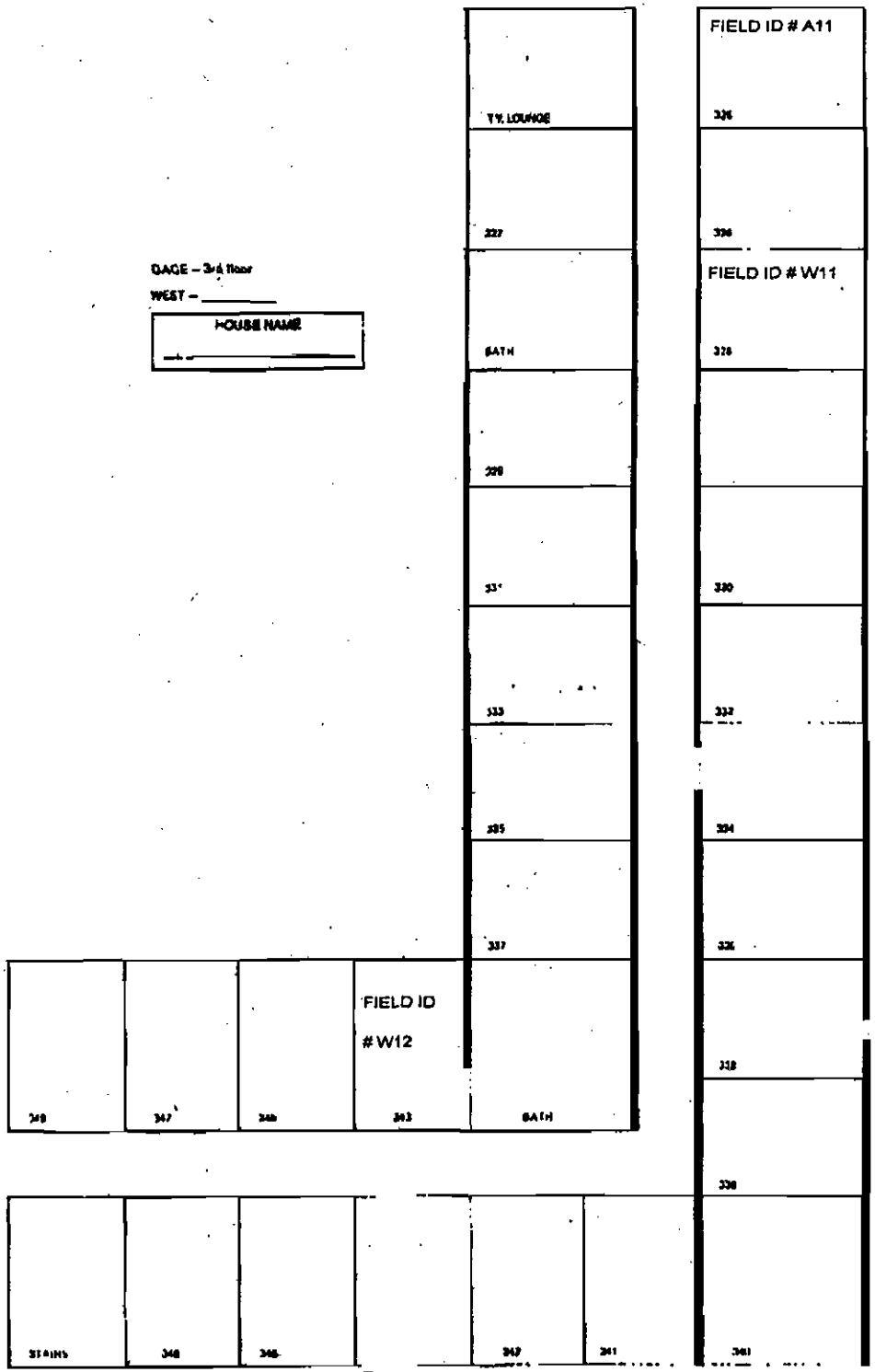

FIELD ID #
W18
STAIRS
101
103
108
107
108
108 OVERFLOW
STUDY LOUNGE
109
110
111
100 OVERFLOW
102
104
106
BATH
FIELD ID # A30
112
113
114
GAGE – 1st floor
EAST –
HOUSE NAME
FIELD ID #
W17
RD APT.
RD OFFICE
RA OFFICE

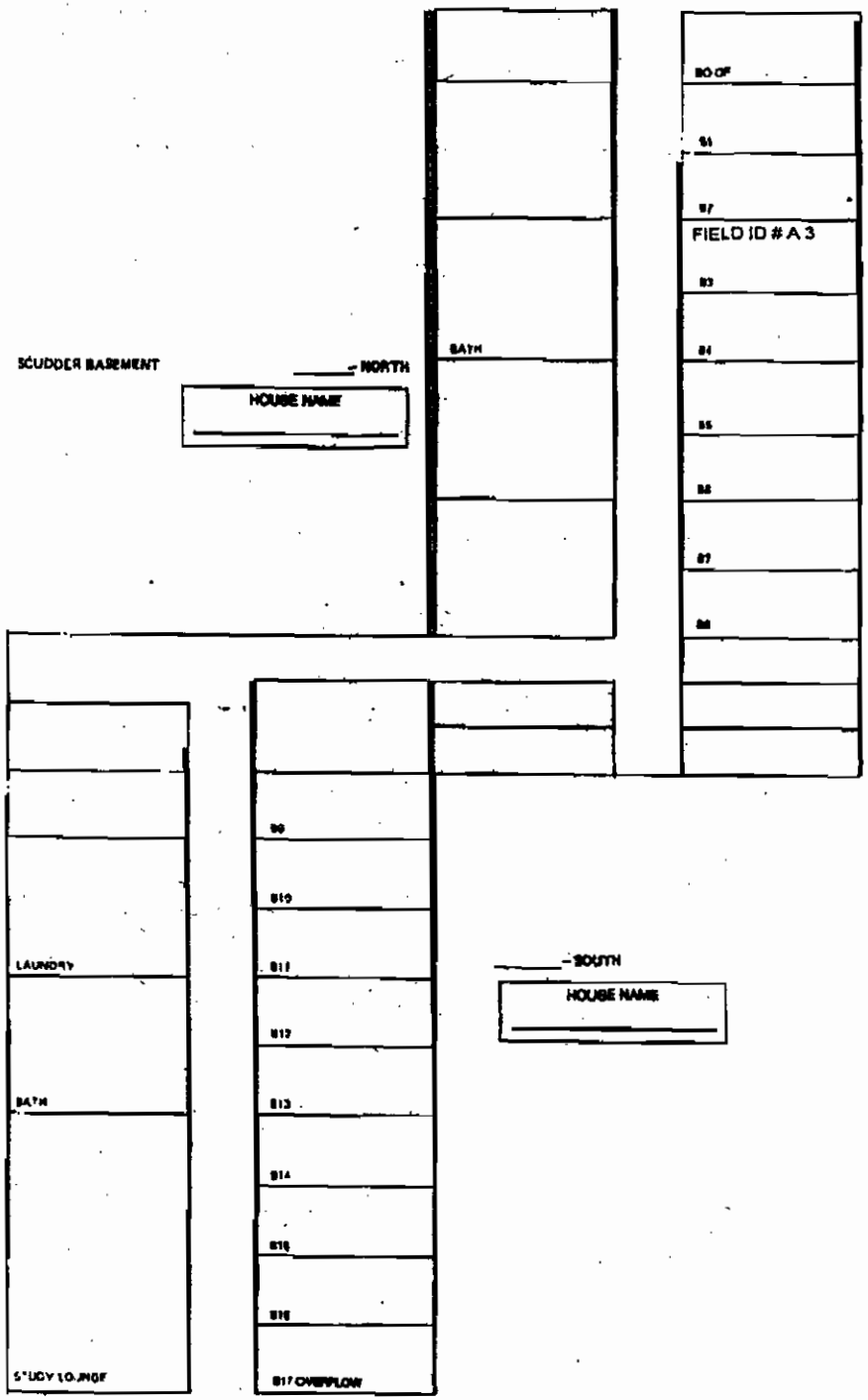

GAOL – 1st floor
WEST
HOUSE NAME
STAIRS
116
FIELD ID # W16
112
118
120
121
122
FIELD ID #
W15
123 OVERFLOW REC ROOM
131
129
127
BATH
124
STAIRS
132
130
126
124 OVERFLOW

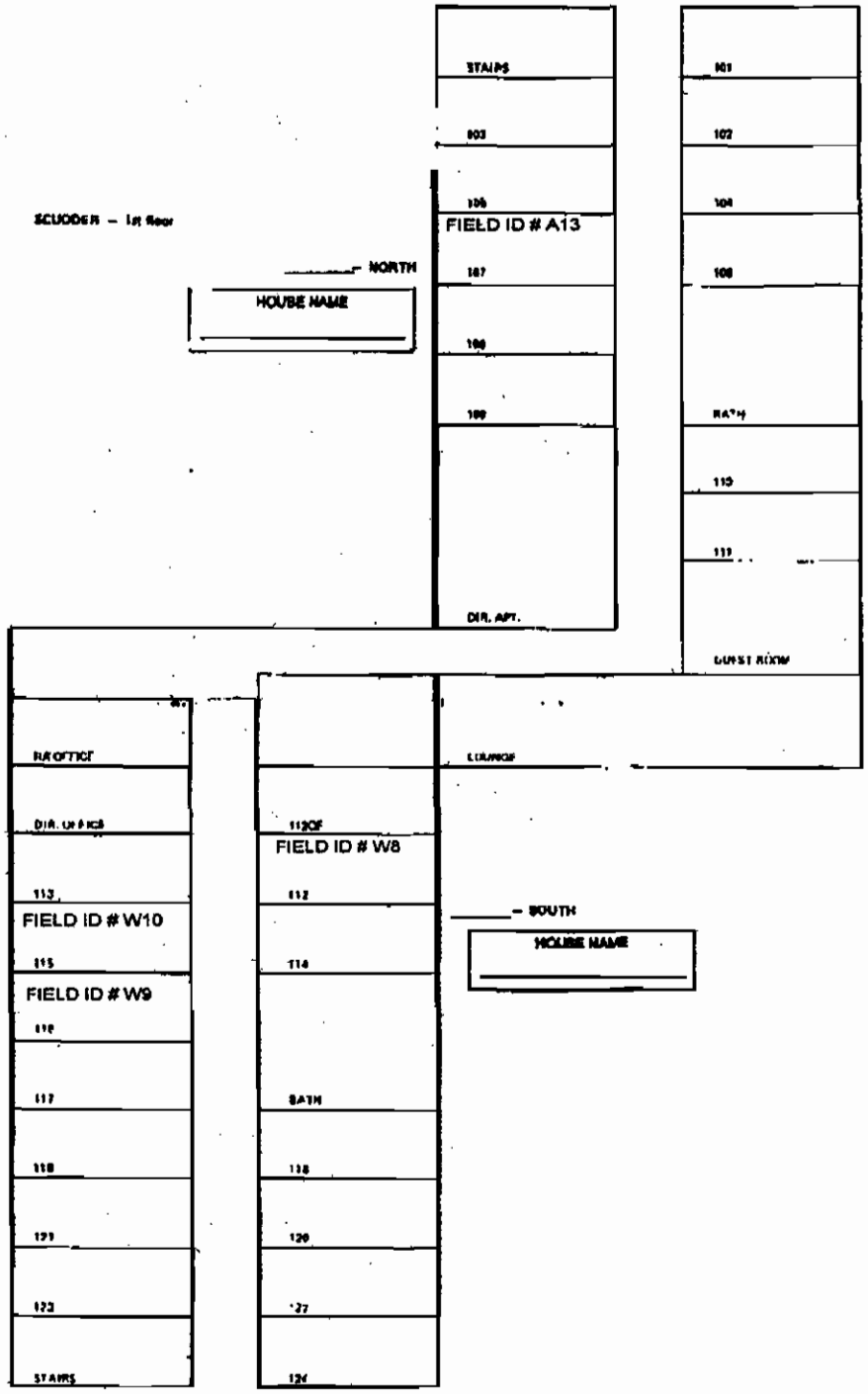

STAIRS
FIELD ID #
A10
BATH
FIELD ID # W14
FIELD ID # W13
STUDY LOUNGE
FIELD ID # A26
HOUSE NAME

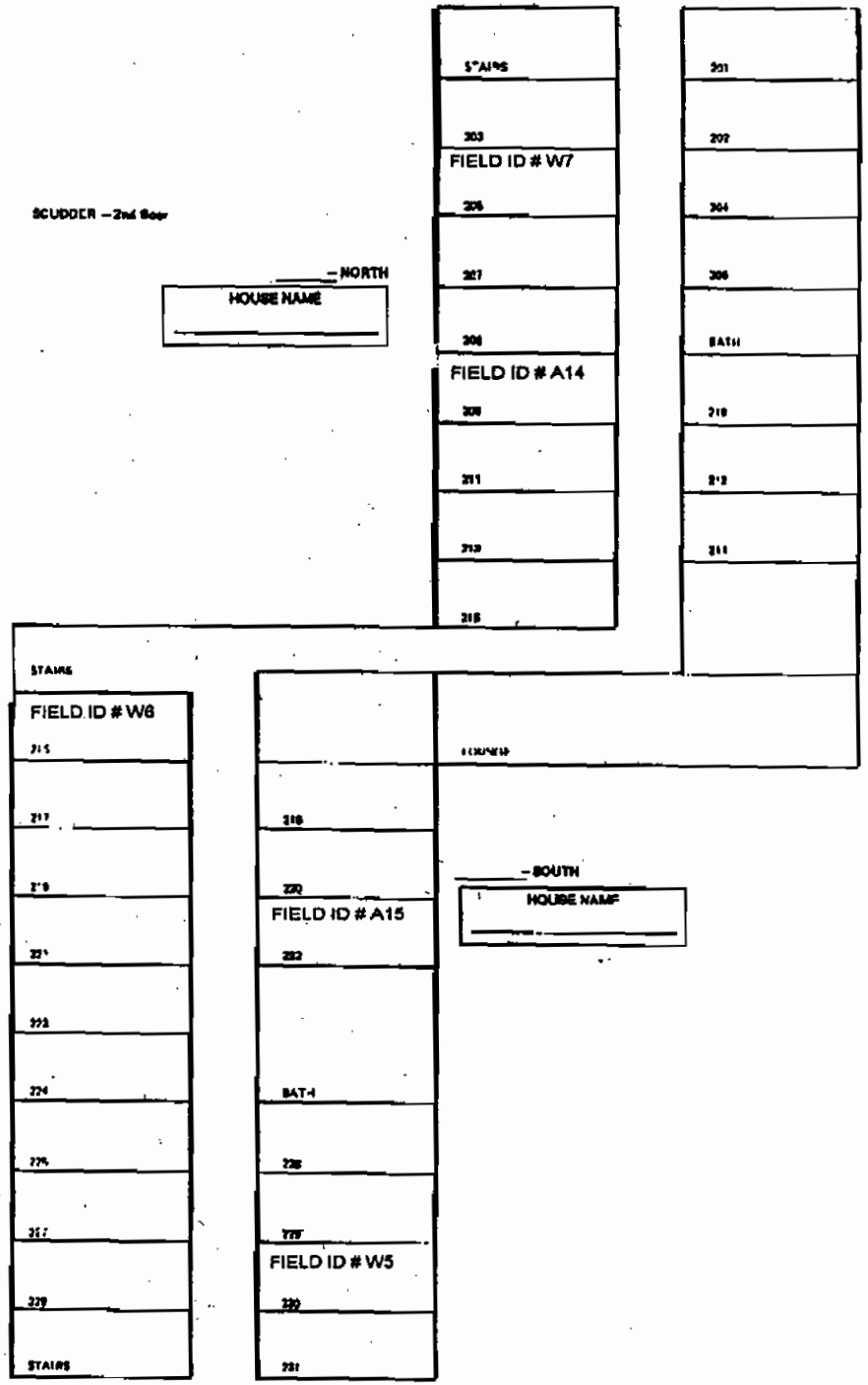

FIELD ID # A28
T.V. LOUNGE
FIELD ID # A27
BATH
FIELD ID # A29
BATH

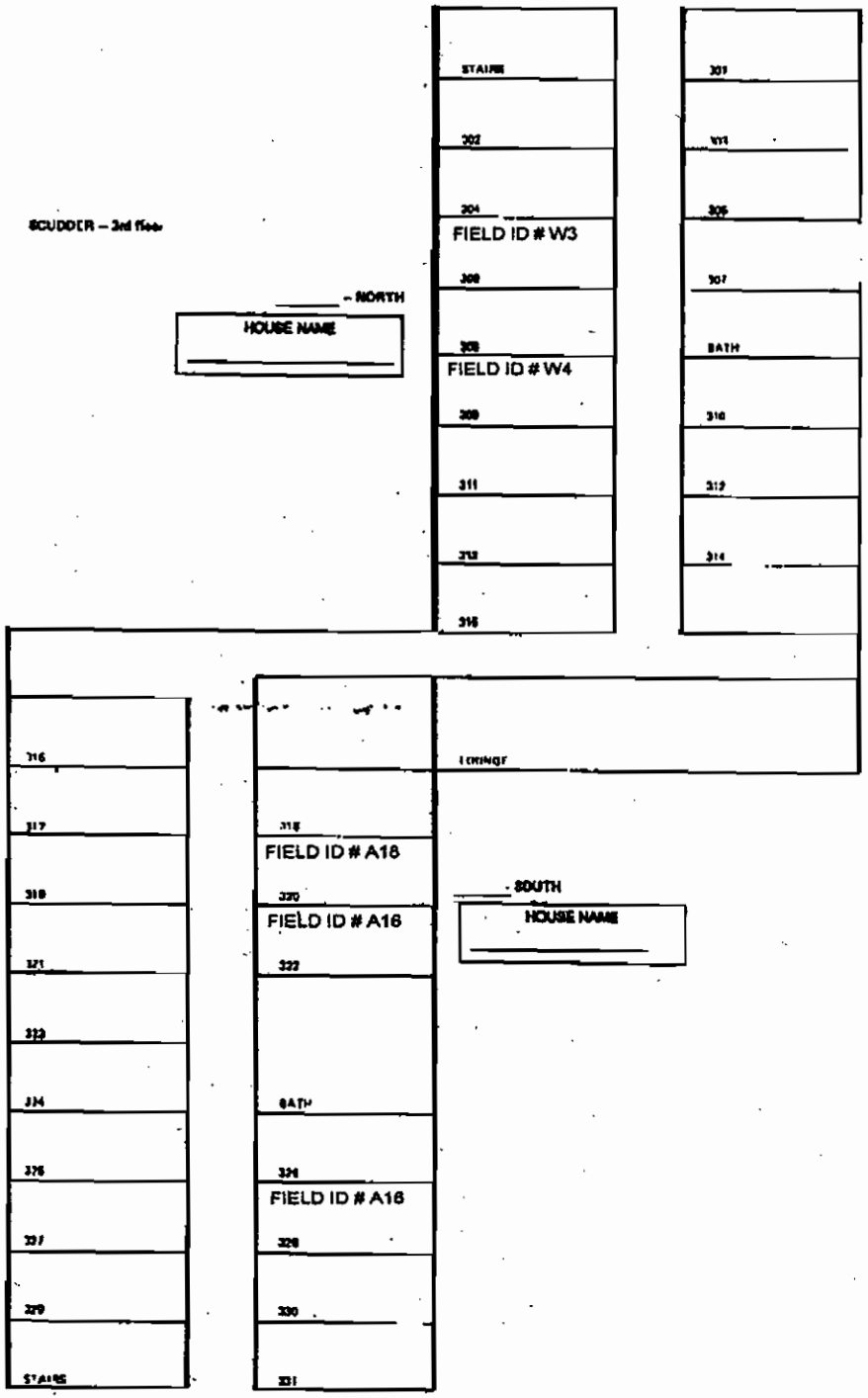

FIELD ID
A12
AND
W20
STAIRS
FIELD ID
A25
BATH
BATH
STAIRS
STUDY LOUNGE
GAGE 3rd floor
EAST –
HOUSE NAME

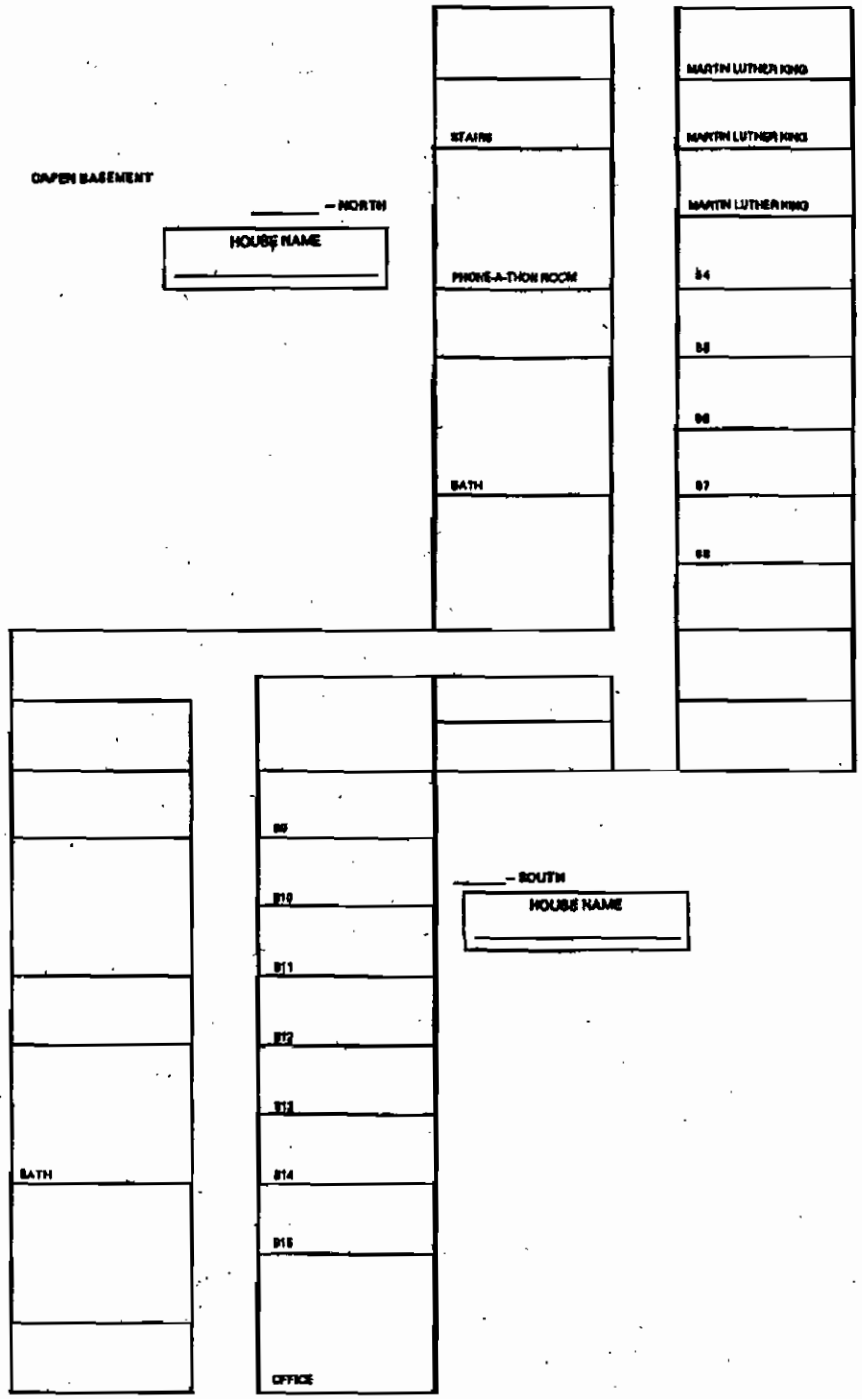

CAGE – 3rd floor
WEST –
HOUSE NAME
TV. LOUNGE
FIELD ID # A11
FIELD ID # W11
BATH
FIELD ID
W12
BATH
STAIRS

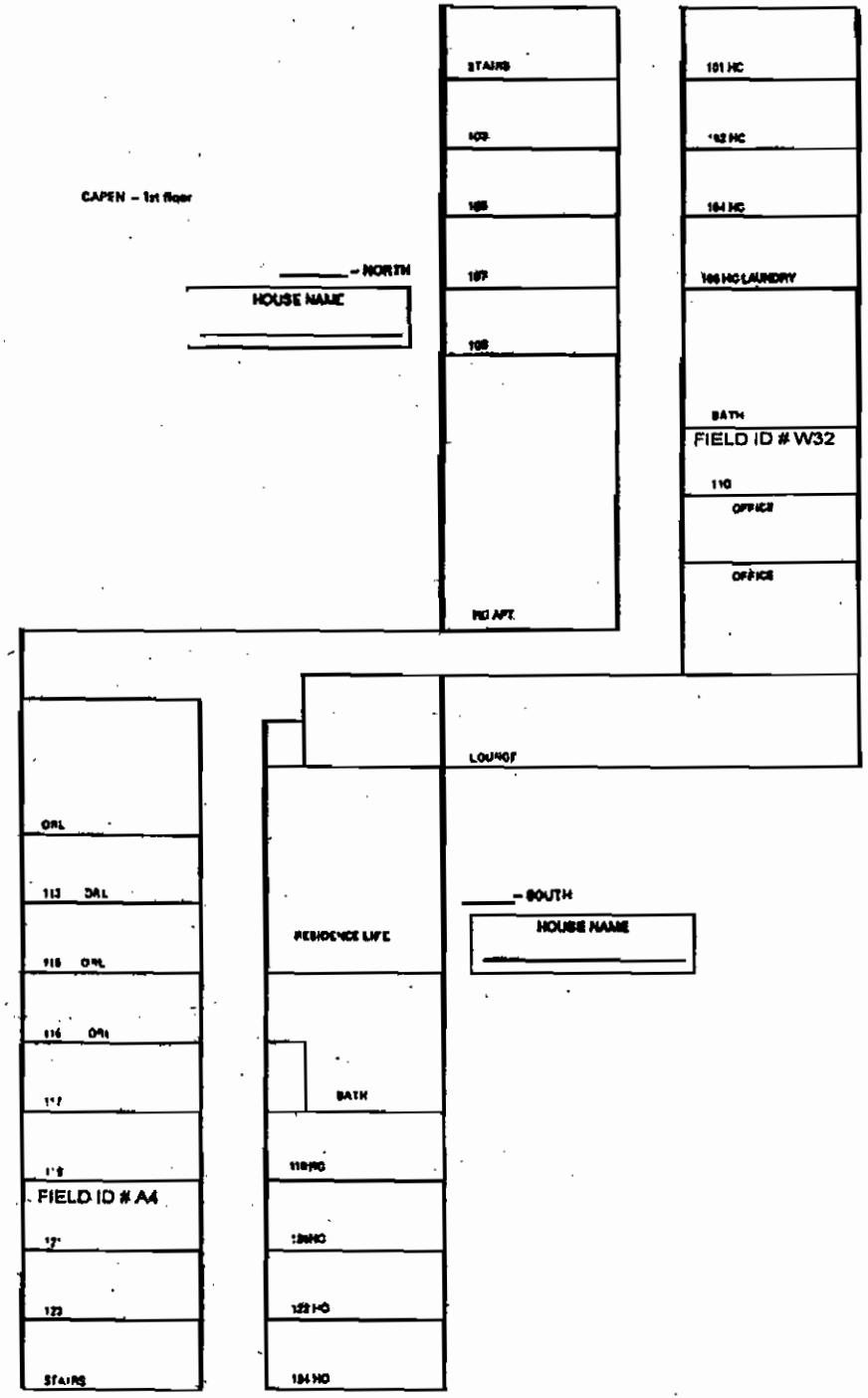

SCUDDER BASEMENT
NORTH
HOUSE NAME
BATH
ROOF
FIELD ID # A 3
LAUNDRY
BATH
STUDY LOUNGE
B17 OVERFLOW
SOUTH
HOUSE NAME

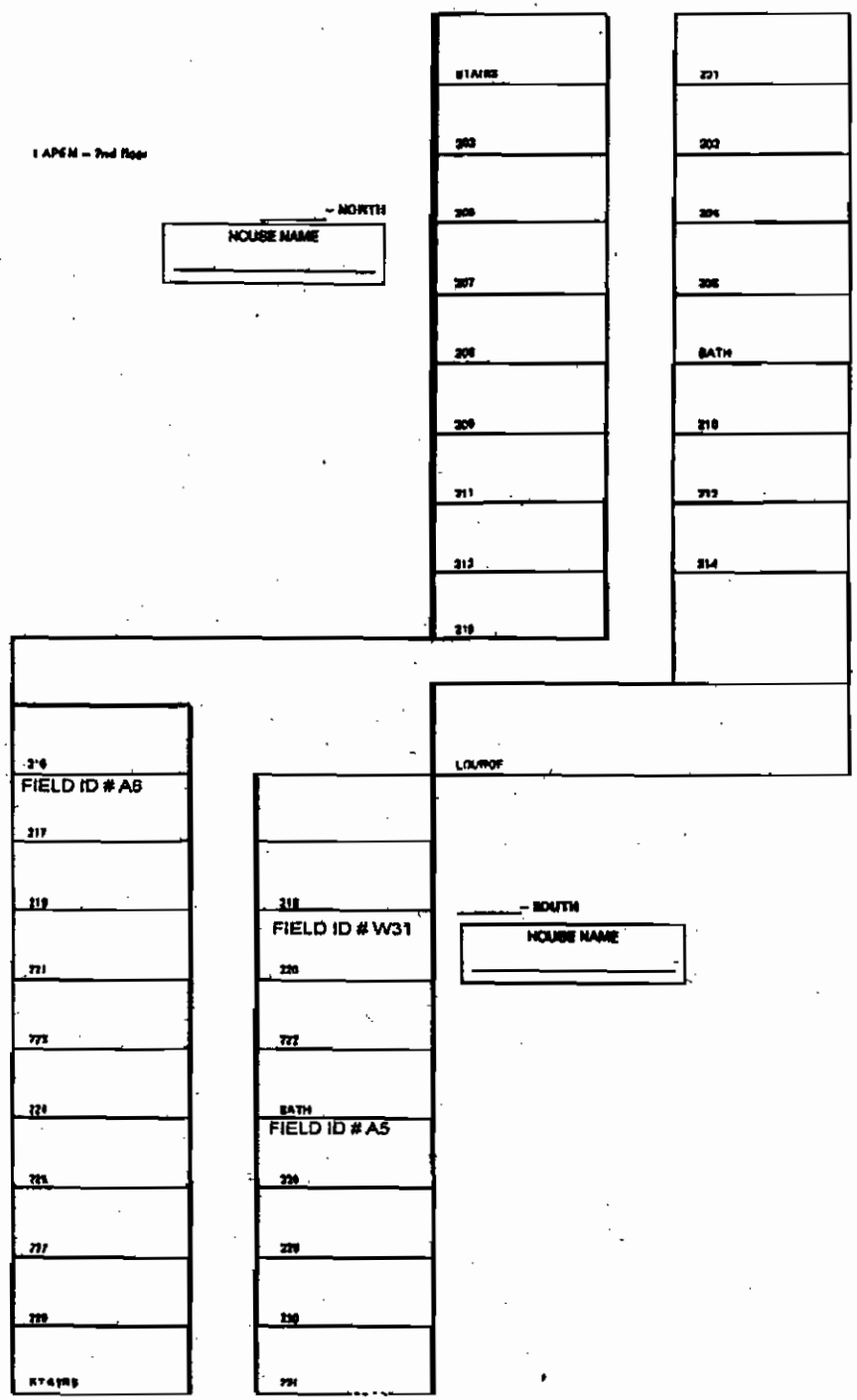

SCUDDER – 1st floor
NORTH
HOUSE NAME
STAIRS
FIELD ID # A13
DIR. APT.
RATH
GUEST ROOM
LIBRARY
FIELD ID # W8
FIELD ID # W10
FIELD ID # W9
BATH
SOUTH
HOUSE NAME

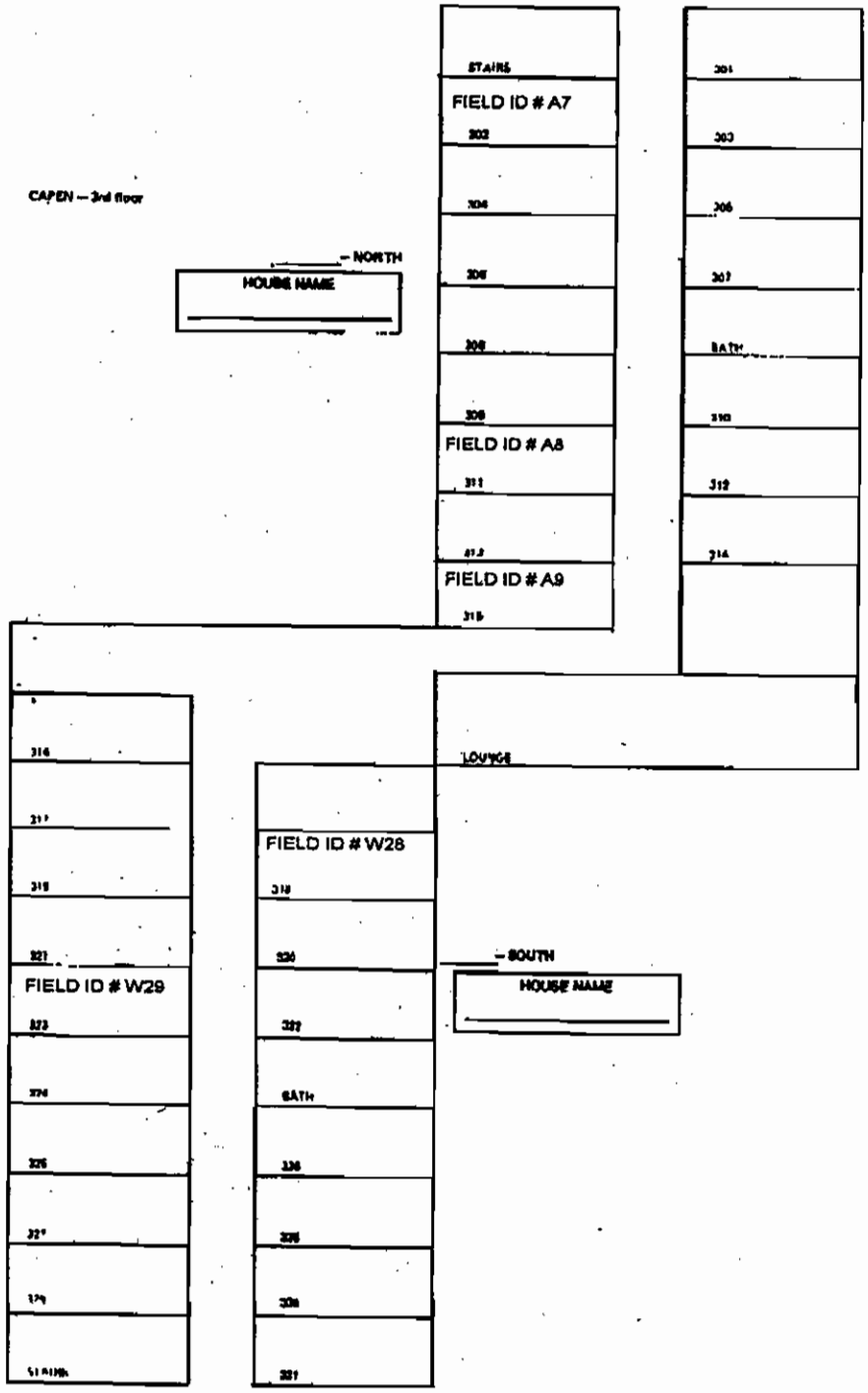

SCUDDER – 2nd floor
NORTH
HOUSE NAME
STAIRS
303
FIELD ID # W7
305
307
309
FIELD ID # A14
311
313
315
STAIRS
FIELD ID # W6
315
317
319
321
323
324
325
327
329
STAIRS
301
302
304
306
BATH
310
312
314
316
318
320
FIELD ID # A15
322
BATH
326
328
FIELD ID # W5
330
331
SOUTH
HOUSE NAME


SCUDDER – 3rd floor
– NORTH
HOUSE NAME
STAIRS
302
304
FIELD ID # W3
308
309
FIELD ID # W4
311
313
316
307
BATH
310
312
314
316
317
318
321
323
324
326
327
329
STAIRS
FIELD ID # A16
320
FIELD ID # A16
322
BATH
324
FIELD ID # A16
328
330
331
– SOUTH
HOUSE NAME


OPEN BASEMENT
NORTH
HOUSE NAME
STAIRS
PHONE-A-THON ROOM
BATH
MARTIN LUTHER KING
MARTIN LUTHER KING
MARTIN LUTHER KING
84
88
96
97
98
BATH
86
B10
B11
B12
B13
B14
B15
OFFICE
SOUTH
HOUSE NAME


CAPEN – 1st floor
– NORTH
HOUSE NAME
STAIRS
101 HC
102 HC
104 HC
106 HC LAUNDRY
BATH
FIELD ID # W32
110
OFFICE
OFFICE
RD APT.
LOUNGE
ORL
113 ORL
RESIDENCE LIFE
– SOUTH
HOUSE NAME
BATH
FIELD ID # A4
STAIRS
184 HD


NORTH
HOUSE NAME
STAIRS
203
205
207
209
211
213
215
221
223
225
BATH
227
229
231
LOUNGE
FIELD ID # A6
217
219
FIELD ID # W31
SOUTH
HOUSE NAME
FIELD ID # A5
BATH


CAPEN — 3rd floor
— NORTH
HOUSE NAME
STAIRS
FIELD ID # A7
FIELD ID # A8
FIELD ID # A9
LOUNGE
FIELD ID # W28
FIELD ID # W29
— SOUTH
HOUSE NAME

APPENDIX C
WIPE SAMPLING LABORATORY REPORT AND CHAIN OF CUSTODY DOCUMENTATION

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LABORATORY REPORT

for

SUNY-NEW PALTZ
ENGINEERING DEPARTMENT
****
**********, ***** ****

Attention: PETER BENTLEY

Purchase Order #: 970815EA

Report date: 08/21/97
Number of samples analyzed: 32
AES Project ID: 970818ED
Invoice #: 178962

ELAP ID#: 10709

AIHA ID#: 7866

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W1
AES sample #: 970818ED01
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W2
AES sample #: 970818ED02
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W3
AES sample #: 970818ED03
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W4
AES sample #: 970818ED04
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT	UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W5
AES sample #: 970818ED05

Date Sampled: 08/18/97
Date sample received: 08/19/97
Samples taken by: Frisone/Watson Location: Suny New Paltz
MATRIX: Wipe grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W6
AES sample #: 970818ED06
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT	UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/ 900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W7
AES sample #: 970818ED07
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W8
AES sample #: 970818ED08
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W9
AES sample #: 970818ED09
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W10
AES sample #: 970818ED10
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W11
AES sample #: 970818ED11
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W12
AES sample #: 970818ED12
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT	UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1 ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W13
AES sample #: 970818ED13
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W14
AES sample #: 970818ED14

Date Sampled: 08/18/97
Date sample received: 08/19/97
Samples taken by: Frisone/Watson Location: Suny New Paltz
MATRIX: Wipe grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W15
AES sample #: 970818ED15
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W16
AES sample #: 970818ED16
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W17
AES sample #: 970818ED17
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W18
AES sample #: 970818ED18
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W19
AES sample #: 970818ED19
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8030	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W20
AES sample #: 970818ED20

Date Sampled: 08/18/97
Date sample received: 08/19/97
Samples taken by: Frisone/Watson Location: Suny New Paltz
MATRIX: Wipe grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W21
AES sample #: 970818ED21
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W22
AES sample #: 970818ED22
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W23
AES sample #: 970818ED23
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W24
AES sample #: 970818ED24
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W25
AES sample #: 970818ED25
Date Sampled: 08/18/97
Date sample received: 08/19/97
Samples taken by: Frisone/Watson Location: Suny New Paltz
MATRIX: Wipe grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W26
AES sample #: 970818ED26
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W27
AES sample #: 970818ED27

Date Sampled: 08/18/97
Date sample received: 08/19/97
Samples taken by: Frisone/Watson Location: Suny New Paltz
MATRIX: Wipe grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W28
AES sample #: 970818ED28
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER	PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260		EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W29
AES sample #: 970818ED29
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

Adirondack Environmental Services, Inc.
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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W30
AES sample #: 970818ED30
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W31
AES sample #: 970818ED31
Samples taken by: Frisone/Watson
MATRIX: Wipe
Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

Adirondack Environmental Services Inc.
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CLIENT: SUNY-NEW PALTZ
CLIENT'S SAMPLE ID: W32
AES sample #: 970818ED32
Samples taken by: Frisone/Watson
MATRIX: Wipe

Date Sampled: 08/18/97
Date sample received: 08/19/97
Location: Suny New Paltz
grab

PARAMETER PERFORMED	METHOD	RESULT		UNITS	NOTEBK REF	TEST DATE
PCB-1016	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1221	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1232	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1242	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1248	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1254	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97
PCB-1260	EPA-8080	<1	ug/	900cm2	KF-PCB-W-7	08/19/97

APPROVED BY:
Report date: 08/21/97

Adirondack Environmental Services, Inc.
314 North Pearl Street
Albany, New York 12207
518-434-4546/434-0891 FAX

REQUEST FOR ANALYSI

CLIENT NAME	PROJECT NAME (Location)	SAMPLERS' (Names)
S.U.N.Y-NEW PALTZ	SUNY New Paltz PCB	J. FRISONE/P. WATSO
ADDRESS	PO NUMBER	SAMPLERS' (Signatures)
	970815EA	

AES SAMPLE NUMBER	SAMPLE IDENTIFICATION	DATE SAMPLED	TIME A = A.M. P = P.M.	MEDIA TYPE/ MATRIX	NO. OF CONT'S	AIR SAMPLE VOLUME (LITERS)	TOTAL SAMPLING TIME (MIN.)	ANALYSIS REQUESTED
970818 ED01	W1	8/18/97	833 A/P	W.Re	1			EPA 8080 - PC
ED02	W2		835 A/P					
ED03	W3		942 A/P					
ED04	W4		942 A/P					
ED05	W5		955 A/P					
ED06	W6		1015 A/P					
ED07	W7		1042 A/P					
ED08	W8		1035 A/P					
ED09	W9		1141 A/P					
ED10	W10		1040 A/P					
ED11	W11		1054 A/P					
ED12	W12		1050 A/P					
ED13	W13		1105 A/P					
ED14	W14	↓	1100 A/P	↓	↓			↓

SEND REPORT TO	SEND INVOICE TO	COMMENTS
PAUL WATSON	PAUL WATSON	

TURN-AROUND TIME — PLEASE CHECK ALL THAT APPLY

- ☐ *STANDARD SERVICE
- ☒ *RUSH SERVICE — Results requested by: 5 DAYS
- ☒ FAX RESULTS TO: PETER BETLEY FAX # 914 257-3388
- ☐ PHONE RESULTS TO: _____ PH # ( ) -

*Turn-around time varies by substance. For most substances, standard turn-around time is ten (10) working days. Please inquire for capacity of rush analysis.

LABORATORY APPROVAL	DATE	TIME	RECEIVED FOR LABORATORY BY	DATE	TIME
			M.	8/19/97	7:30

CHAIN OF CUSTODY

RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME
J. Frisone			
RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME

WHITE — Lab Copy YELLOW — Sampler Copy PINK — Generator Copy

Adirondack Environmental Services, Inc.
314 North Pearl Street
Albany, New York 12207
518-434-4546/434-0891 FAX

2 of 3

REQUEST FOR ANALYS

CLIENT NAME	PROJECT NAME (Location)	SAMPLERS' (Names)
SUNY - NEW PALTZ	SUNY NEW PALTZ-PCB	J. FRISONE / P. WATSON
ADDRESS	PO NUMBER	SAMPLERS' (Signatures)
	970815EA	

AES SAMPLE NUMBER	SAMPLE IDENTIFICATION	DATE SAMPLED	TIME A = A.M. P = P.M.	MEDIA TYPE/ MATRIX	NO. OF CONT'S	AIR SAMPLE VOLUME (LITERS)	TOTAL SAMPLING TIME (MIN.)	ANALYSIS REQUESTED
970818 ED15	W15	8/18/97	1115 A	WIPE	1			EPA 8080-PC
ED16	W16		1115 A					
ED17	W17		1132 A					
ED18	W18		1135 A					
ED19	W19		1140 A					
ED20	W20		1143 A					
ED21	W21		232 P					
ED22	W22		243 P					
ED23	W23		253 P					
ED24	W24		256 P					
ED25	W25		301 P					
ED26	W26		305 P					
ED27	W27		308 P					
ED28	W28	↓	324 P	↓	↓			↓

SEND REPORT TO	SEND INVOICE TO	COMMENTS
PAUL WATSON	PAUL WATSON	

TURN-AROUND TIME — PLEASE CHECK ALL THAT APPLY

- ☐ *STANDARD SERVICE
- ☒ *RUSH SERVICE — Results requested by: 5 DAYS
- ☒ FAX RESULTS TO: PETE BETLEY FAX # (914)257-3388
- ☐ PHONE RESULTS TO: _____ PH # (___) _____

*Turn-around time varies by substance. For most substances, standard turn-around time is ten (10) working days. Please inquire for capacity of rush analysis.

LABORATORY APPROVAL	DATE	TIME	RECEIVED FOR LABORATORY BY	DATE	TIME
			M.	8/19/97	730

CHAIN OF CUSTODY

RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME
RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME

WHITE — Lab Copy YELLOW — Sampler Copy PINK — Generator Copy

Adirondack Environmental Services, Inc.
314 North Pearl Street
Albany, New York 12207
518-434-4546/434-0891 FAX

3 of 3

REQUEST FOR ANALYSI

CLIENT NAME	PROJECT NAME (Location)	SAMPLERS' (Names)
S.U.N.Y. - NEW PALTZ	SUNY NEW PALTZ-PCB	J. FRISONE / P. WATSON
ADDRESS	PO NUMBER	SAMPLERS' (Signatures)
	970815EA	J. Frisone / Paul Watson

AES SAMPLE NUMBER	SAMPLE IDENTIFICATION	DATE SAMPLED	TIME A = A.M. P = P.M.	MEDIA TYPE/ MATRIX	NO. OF CONT'S	AIR SAMPLE VOLUME (LITERS)	TOTAL SAMPLING TIME (MIN.)	ANALYSIS REQUESTED
970819 ED29	W29	8/18/97	324 (A) (P)	WIPE	1			EPA 8080-PC
ED30	W30	↓	329 (A) (P)	↓	↓			↓
ED31	W31		331 (A) (P)					
ED32	W32		335 (A) (P)					
			A P					
			A P					
			A P					
			A P					
			A P					
			A P					
			A P					
			A P					
			A P					
			A P					

SEND REPORT TO	SEND INVOICE TO	COMMENTS
PAUL WATSON	PAUL WATSON	

TURN-AROUND TIME — PLEASE CHECK ALL THAT APPLY

- ☐ *STANDARD SERVICE
- ☒ *RUSH SERVICE — Results requested by: 5 DAYS
- ☒ FAX RESULTS TO: PETE BETLEY FAX # (914) 257-3388
- ☐ PHONE RESULTS TO: _____ PH # ( ) -

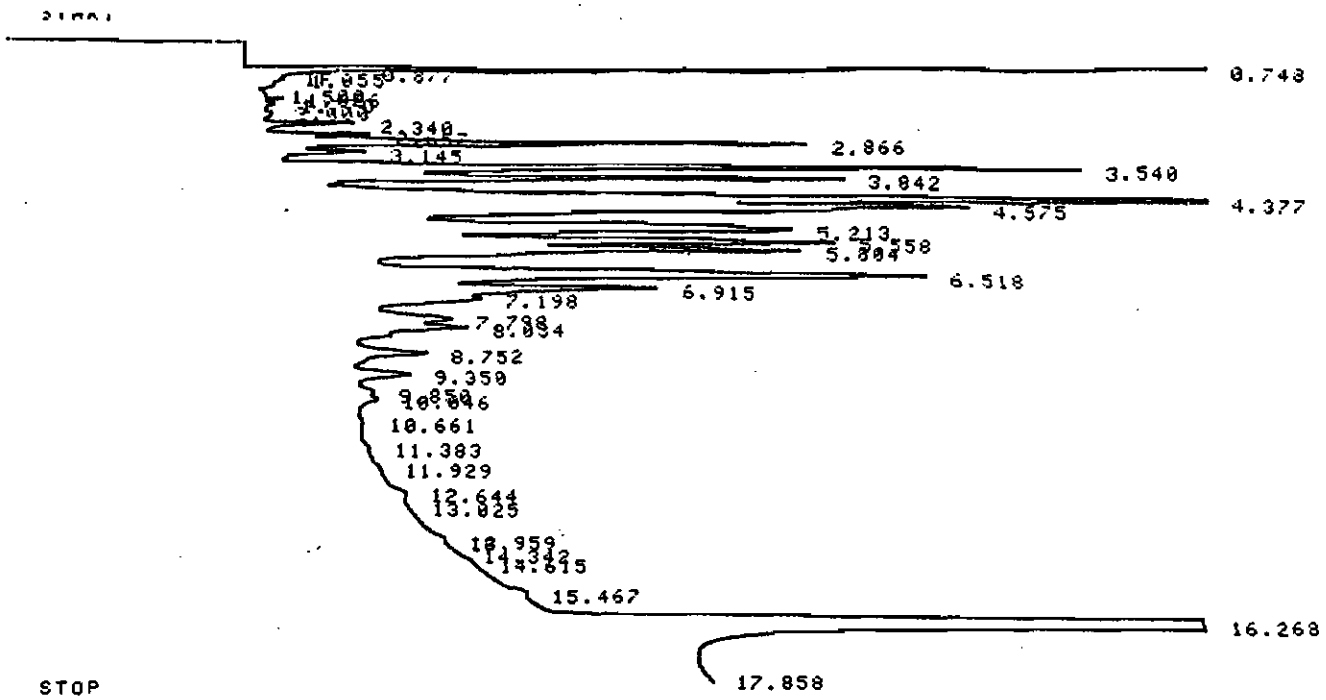
*Turn-around time varies by substance. For most substances, standard turn-around time is ten (10) working days. Please inquire for capacity of rush analysis.

LABORATORY APPROVAL	DATE	TIME	RECEIVED FOR LABORATORY BY	DATE	TIME
			M. L	8/19/97	730

CHAIN OF CUSTODY

RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME
J. Frisone			
RELINQUISHED BY (Signature)	RECEIVED BY (Signature)	DATE	TIME

WHITE — Lab Copy YELLOW — Sampler Copy PINK — Generator Copy



STOP

RUN# 13326 AUG 19, 1997 07:47:46

SAMPLE NAME: 1242 0.5 SAMPLE# 1
2

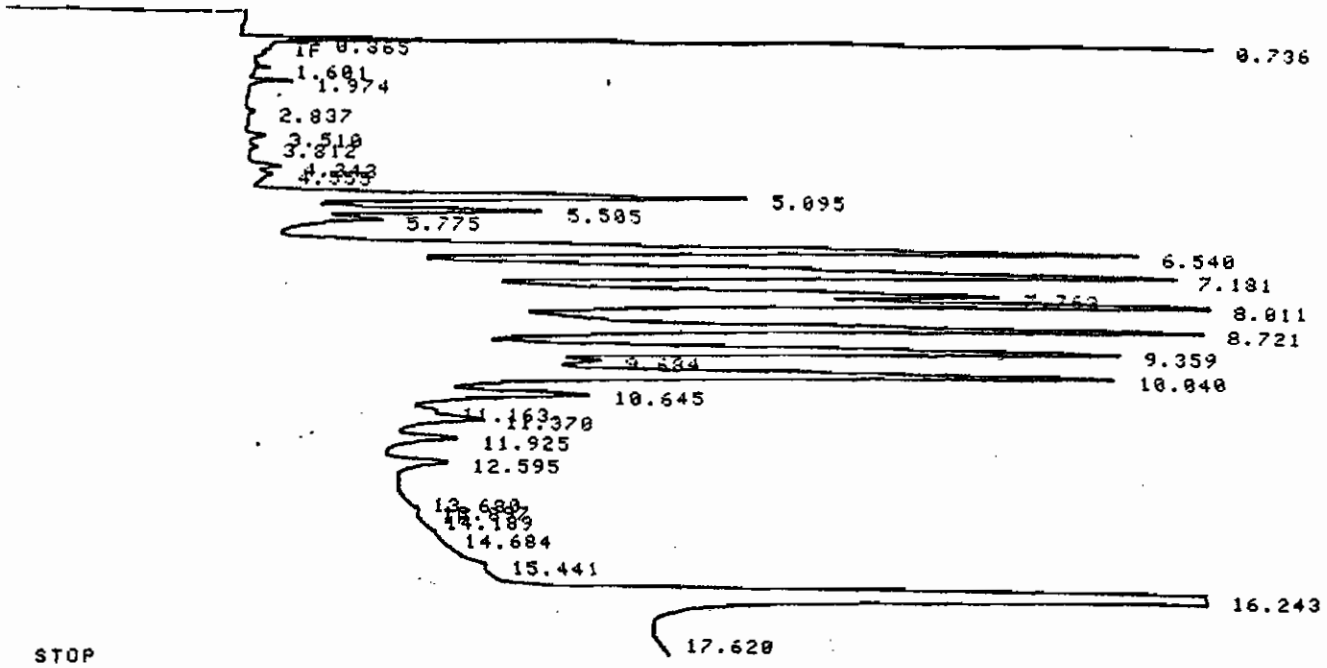
AREA%

RT	AREA	TYPE	WIDTH	AREA%
.748	78115	PV	.040	6.51715
.877	3623	VB	.045	.30227
1.055	129	BB	.039	.01076
1.500	251	PB	.697	.02094
1.626	722	BB	.051	.06024
1.770	719	BB	.075	.05999
2.000	61	BB	.038	.00509
2.340	7029	PB	.088	.58643
2.657	3941	BB	.062	.32880
2.866	49854	BB	.112	4.15933
3.145	5281	BB	.089	.44059
3.540	80763	BB	.123	6.73807
3.842	42811	BB	.105	3.57173
4.377	90103	BB	.115	7.51731
4.575	45808	BB	.178	3.82177
5.213	76417	BB	.244	6.37548
5.558	42507	BB	.143	3.54637
5.804	37098	BB	.140	3.09589
6.518	102766	PB	.226	8.57379
6.915	26246	BB	.149	2.18971
7.198	4446	BB	.174	.37093
7.798	6856	PB	.177	.57200
8.034	6694	BB	.123	.55848
8.752	13556	VB	.210	1.13098
9.350	9340	BB	.191	.77924
9.850	882	PB	.127	.07359
10.046	1237	BB	.134	.10320
10.661	136	PB	.087	.01135
14.615	138	BB	.460	.01151
16.268	461051	BB	.210	38.46558
17.858	27	PB	.056	.00225

PC BW7

TOTAL AREA=1198607
MUL FACTOR=1.0000E+00

RUN #13327 AUG 19, 1997 08:09:36
START



STOP

RUN# 13327 AUG 19, 1997 08:09:36

SAMPLE NAME: 1254 0.5 SAMPLE# 2

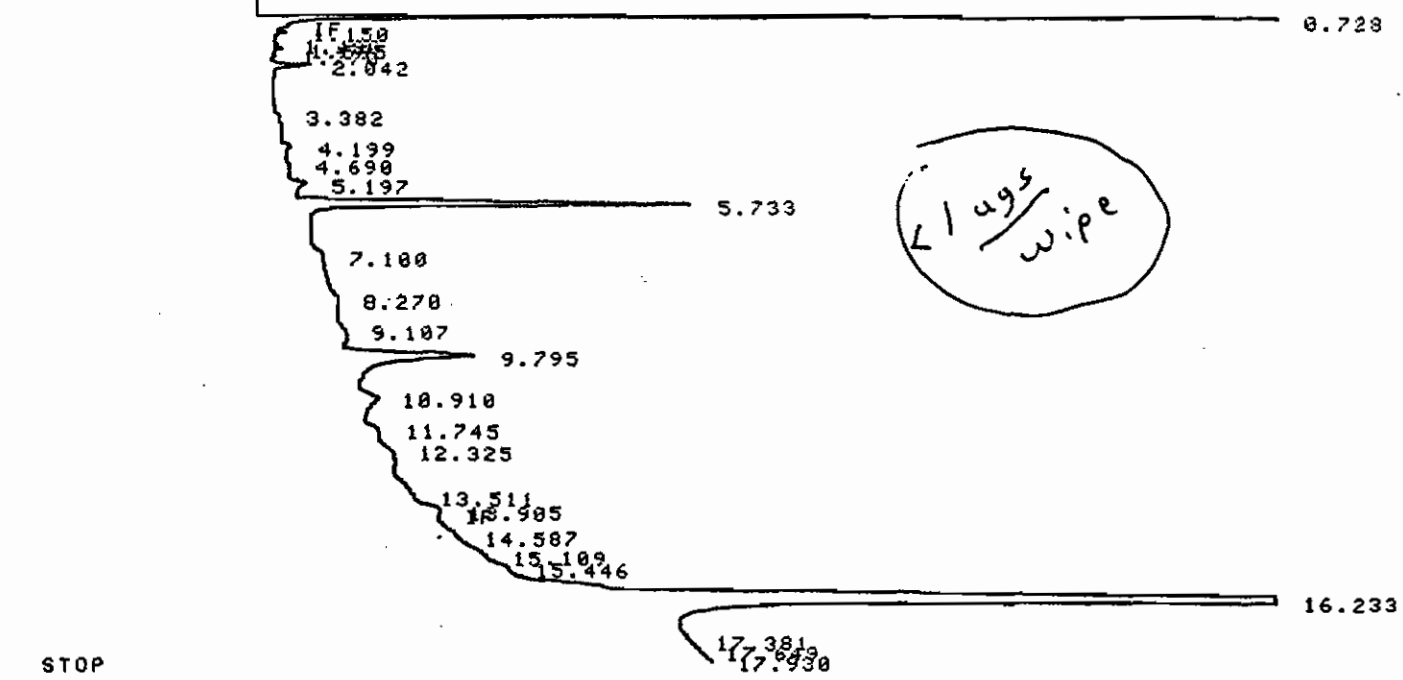
AREA%

RT	AREA	TYPE	WIDTH	AREA%
.736	38716	PB	.033	2.59745
.865	228	BP	.010	.01530
1.601	767	VB	.054	.05146
1.974	3080	BB	.080	.20664
2.837	872	VB	.105	.05850
3.510	1842	PB	.118	.12358
3.812	900	BB	.106	.06038
4.343	2453	PB	.112	.16457
4.555	1559	BB	.129	.10459
5.095	73570	PB	.177	4.93580
5.585	26012	BB	.134	1.74514
5.775	6406	BB	.119	.42978
6.540	152407	BB	.219	10.22495
7.181	176131	BB	.276	11.81659
7.763	35875	BB	.152	2.40685
8.011	74975	BB	.145	5.03005
8.721	142921	BB	.230	9.58854
9.359	110643	BB	.289	7.42301
9.684	3223	BB	.101	.21623
10.040	113226	BB	.214	7.59631
10.645	27551	BB	.205	1.84839
11.163	511	BB	.101	.03428
11.370	9647	BB	.190	.64722
11.925	10608	BB	.189	.71169
12.595	11506	PB	.217	.77194
13.897	218	BB	.151	.01463
14.684	427	PB	2.372	.02865
16.243	464222	BB	.213	31.14454
17.620	44	PB	.244	.00295

TOTAL AREA=1490540
MUL FACTOR=1.0000E+00

10690

RUN #13334 AUG 19, 1997 11:16:39
START



STOP

RUN# 13334 AUG 19, 1997 11:16:39

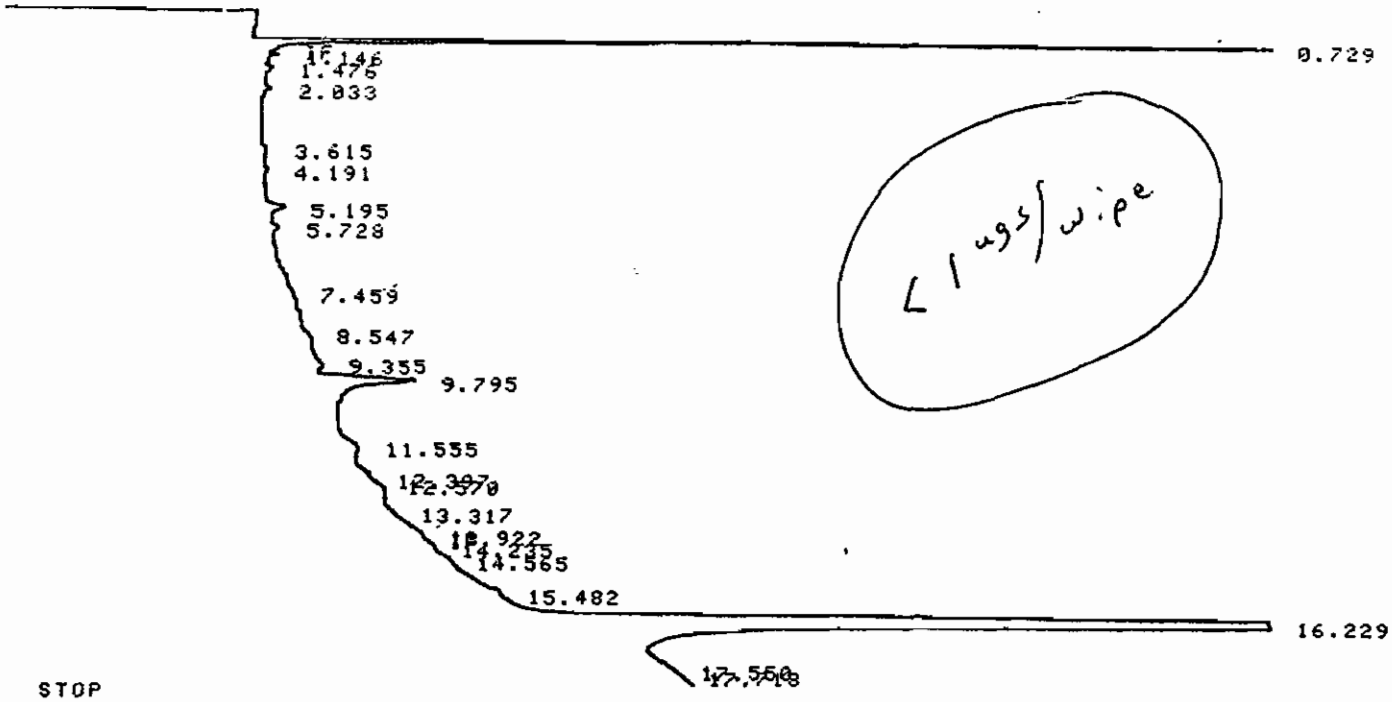
SAMPLE NAME: 818ED1 SAMPLE# 1
2

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.728	52132	PB	.040	13.80450
1.150	480	PB	.050	.12710
1.480	257	PB	.056	.06805
1.595	454	BB	.051	.12022
1.770	220	PB	.052	.05826
2.042	2656	BB	.094	.70331
3.382	260	PB	.083	.06885
4.199	752	PB	.117	.19913
5.197	1645	PB	.147	.43559
5.733	44269	PB	.134	11.72238
7.100	185	PB	.193	.04899
8.270	66	PB	.550	.01748
9.795	23672	PB	.224	6.26832
13.511	527	PB	2.928	.13955
13.905	1430	PB	.168	.37866
16.233	248460	PB	.228	65.79197
17.381	103	PB	.429	.02727
17.649	40	VB	.222	.01059
17.930	37	PB	.206	.00980

TOTAL AREA= 377645
MUL FACTOR=1.0000E+00

RUN #13335 AUG 19, 1997 11:38:27
START



STOP

RUN# 13335 AUG 19, 1997 11:38:27

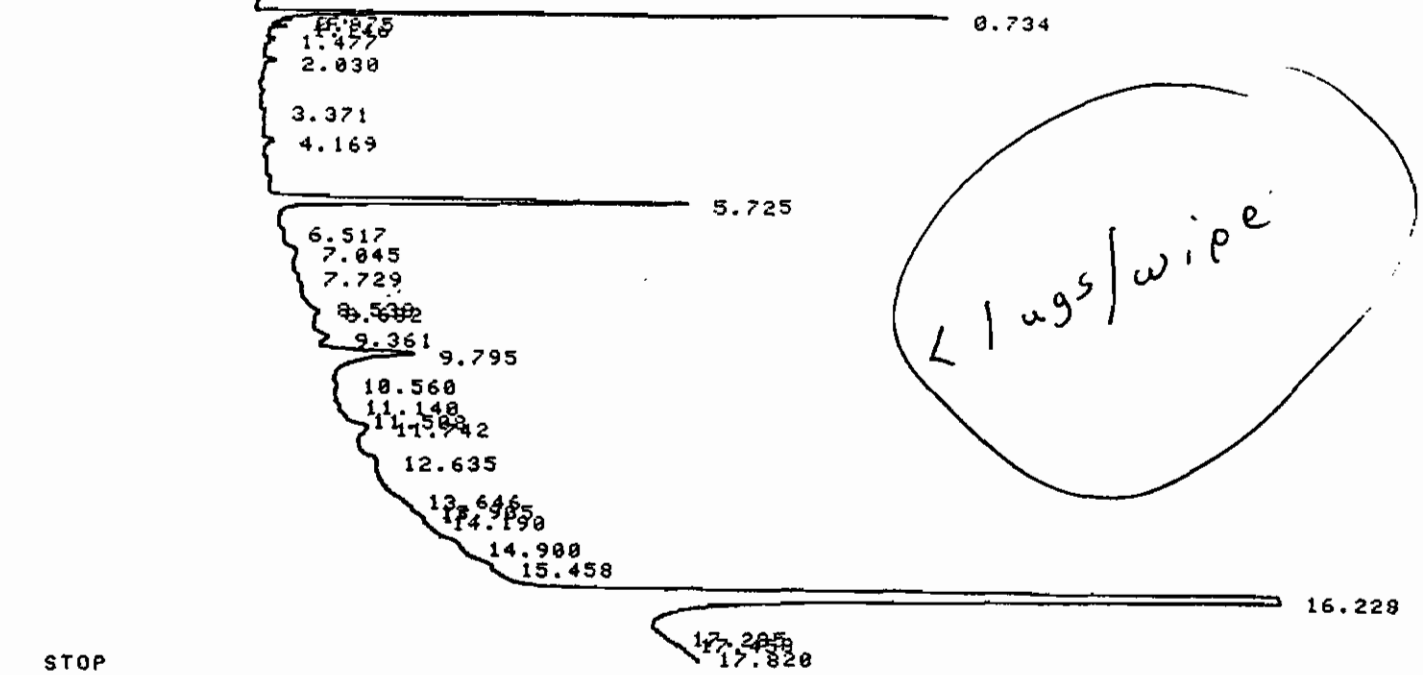
SAMPLE NAME: 818ED2 SAMPLE# 2

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.729	64158	PB	.036	16.01712
1.146	426	PB	.044	.10635
1.476	345	PB	.061	.08613
2.033	819	VB	.104	.20446
3.615	613	BB	.129	.15304
4.191	710	PB	.138	.17725
5.195	2219	PB	.153	.55398
5.728	881	I PB	.124	.21994
9.795	16279	PB	.217	4.06407
12.397	335	PB	1.396	.08363
13.922	716	PB	.628	.17875
14.235	257	PB	.428	.06416
15.482	276	PB	.460	.06890
16.229	312479	BB	.220	78.01072
17.718	46	BB	.192	.01148

TOTAL AREA= 400559
MUL FACTOR=1.0000E+00

RUN #13336 AUG 19, 1997 12:00:31
START



4 lugs/wipe

STOP

RUN# 13336 AUG 19, 1997 12:00:31

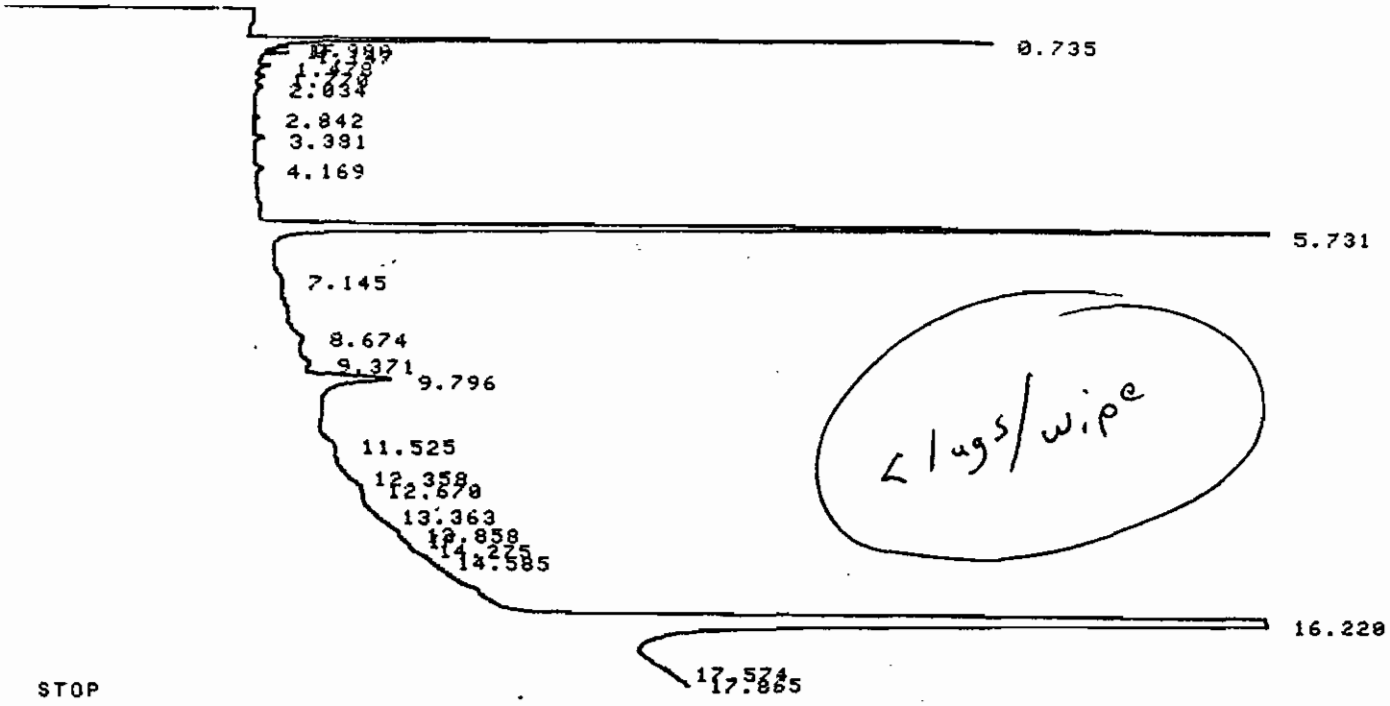
SAMPLE NAME: 818ED3 SAMPLE# 3
4

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.734	32979	PV	.056	9.00893
.975	1640	VB	.077	.44800
1.146	647	BB	.043	.17674
1.477	722	PB	.087	.19723
2.030	1087	PB	.096	.29694
3.371	319	PB	.087	.08714
5.725	46595	PB	.133	12.72844
6.517	287	PB	.126	.07840
7.045	127	PB	.068	.03469
8.538	16	PB	.089	.00437
8.682	979	BB	.147	.26744
9.361	2214	PB	.215	.60480
9.795	19895	BB	.258	5.43475
10.560	13	PB	.043	.00355
11.508	72	PB	.240	.01967
11.742	3586	BB	.249	.97959
13.905	109	BB	.908	.02978
14.190	270	PB	.409	.07376
16.228	254198	PB	.218	69.43754
17.820	323	BB	1.346	.08823

TOTAL AREA= 366070
MUL FACTOR=1.0000E+00

RUN #13337 AUG 19, 1997 12:22:18
START



4 wipes/wipe

STOP

RUN# 13337 AUG 19, 1997 12:22:18

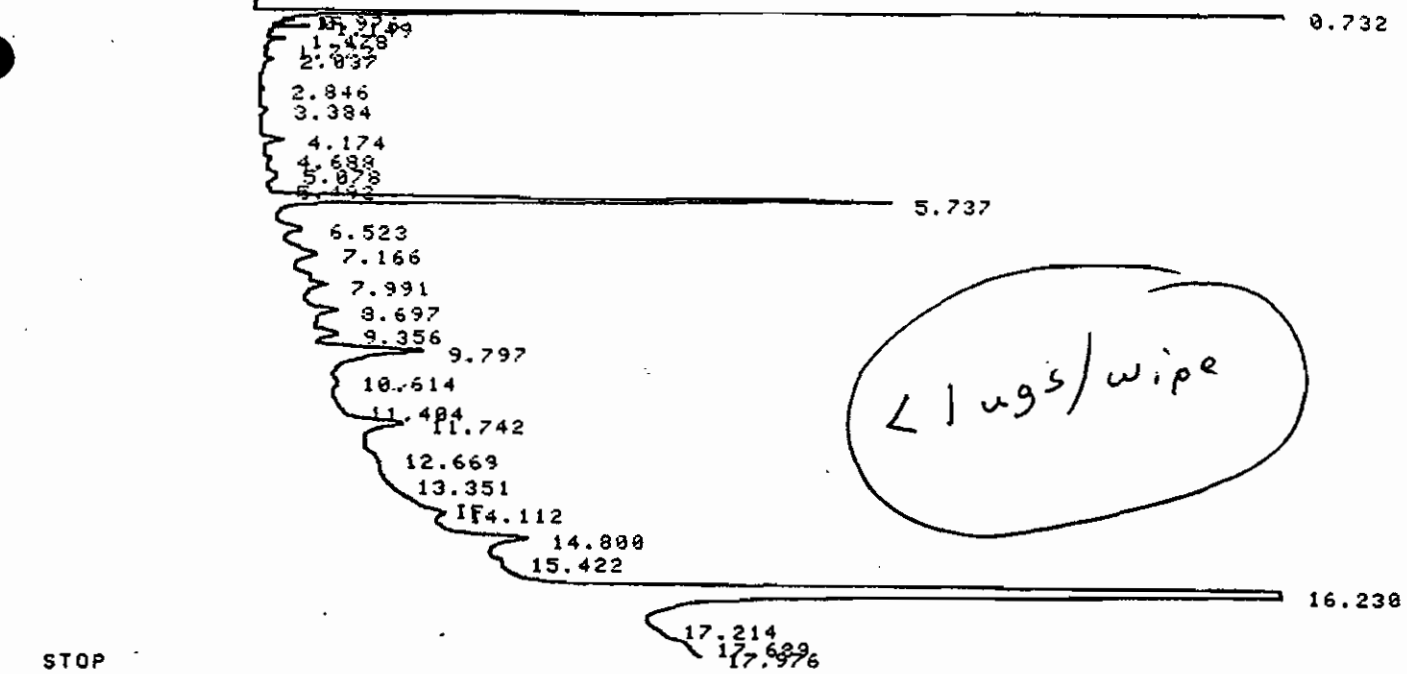
SAMPLE NAME: 818ED4 SAMPLE# 4
725LBLK

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.735	36548	PB	.058	8.31714
.980	601	BB	.043	.13677
1.147	861	PB	.037	.19594
1.478	699	PB	.064	.15907
1.770	552	PB	.069	.12562
2.034	603	BB	.091	.13722
2.842	420	BB	.076	.09558
3.381	870	BB	.093	.19798
4.169	958	PB	.143	.21801
5.731	118194	PB	.130	26.89713
7.145	633	PB	.251	.14405
9.371	1113	PB	.199	.25328
9.796	16311	PB	.239	3.71185
11.525	54	PB	.129	.01229
12.358	104	PB	.433	.02367
12.670	54	BB	.225	.01229
14.275	122	PB	.678	.02776
16.220	260673	BB	.195	59.32070
17.865	60	PB	.250	.01365

TOTAL AREA= 439430
MUL FACTOR=1.0000E+00

RUN #13338 AUG 19, 1997 12:44:12
START



STOP

RUN# 13338 AUG 19, 1997 12:44:12

SAMPLE NAME: 818ED5 SAMPLE# 5

AREA%

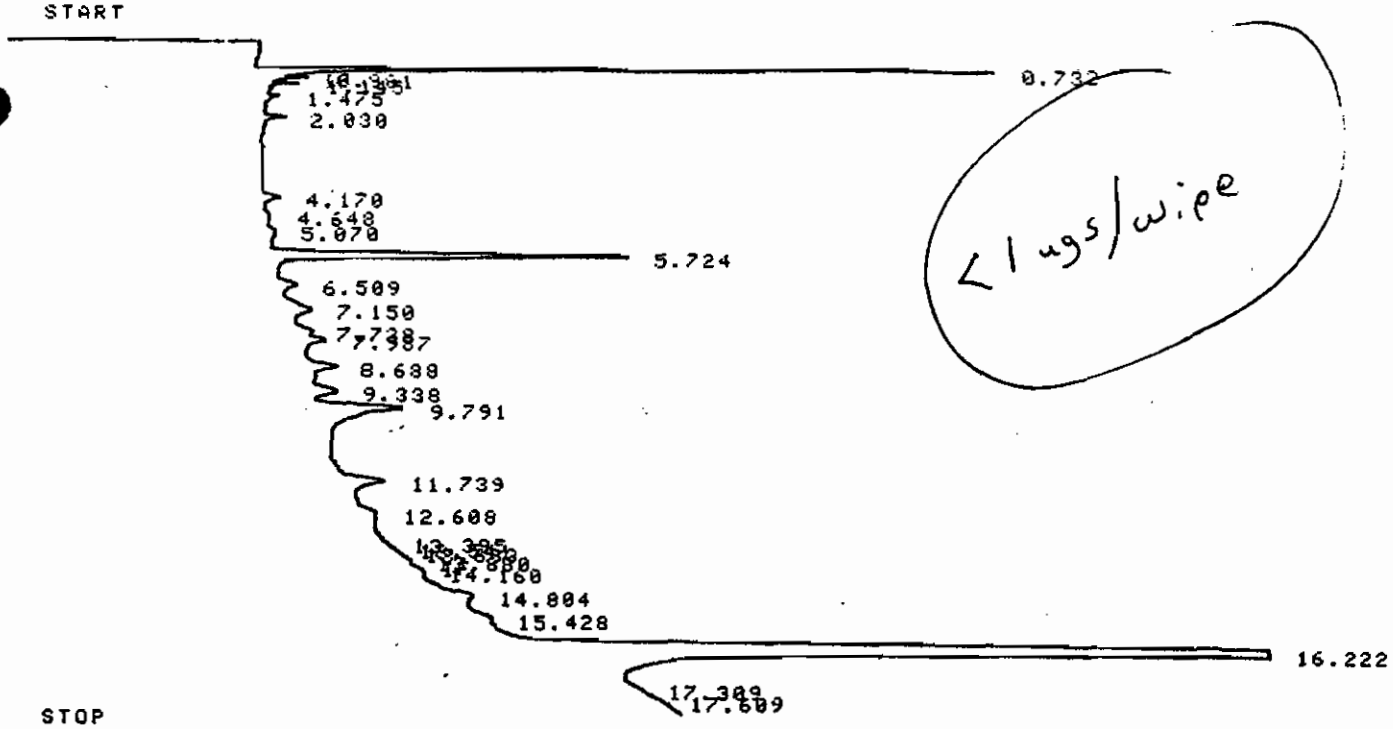
RT	AREA	TYPE	WIDTH	AREA%
.732	47961	PB	.848	10.15699
.976	731	BB	.057	.15481
1.149	1266	BB	.039	.26811
1.478	1159	PB	.073	.24545
1.773	302	PB	.064	.06396
2.037	818	I VB	.097	.17323
2.846	424	BB	.080	.08979
3.384	426	PB	.083	.09022
4.174	2265	BB	.135	.47967
4.688	717	PB	.124	.15184
5.078	407	PB	.092	.08619
5.492	166	BB	.084	.03515
5.737	67999	BB	.130	14.40056
6.523	3696	PB	.188	.78272
7.166	6350	BB	.279	1.34478
7.991	2315	BB	.151	.49026
8.697	4180	PB	.195	.88522
9.356	4182	PB	.211	.88565
9.797	21445	BB	.254	4.54154
10.614	349	PB	.135	.07391
11.404	127	BB	2.117	.02690
11.742	9802	BB	.230	2.07583
12.669	199	PB	.663	.04214
14.800	11187	PB	.220	2.36914
15.422	36	PB	.120	.00762
16.230	283688	PB	.215	60.07830

TOTAL AREA= 472197
MUL FACTOR=1.0000E+00

$\frac{20723}{45537} \times 0.025 \times 20$

$= 0.2$ ugs/wipe

RUN #13339 AUG 19, 1997 13:06:00
START



< 1 ugs/wipe

STOP

RUN# 13339 AUG 19, 1997 13:06:00

SAMPLE NAME: 818ED6 SAMPLE# 6
5

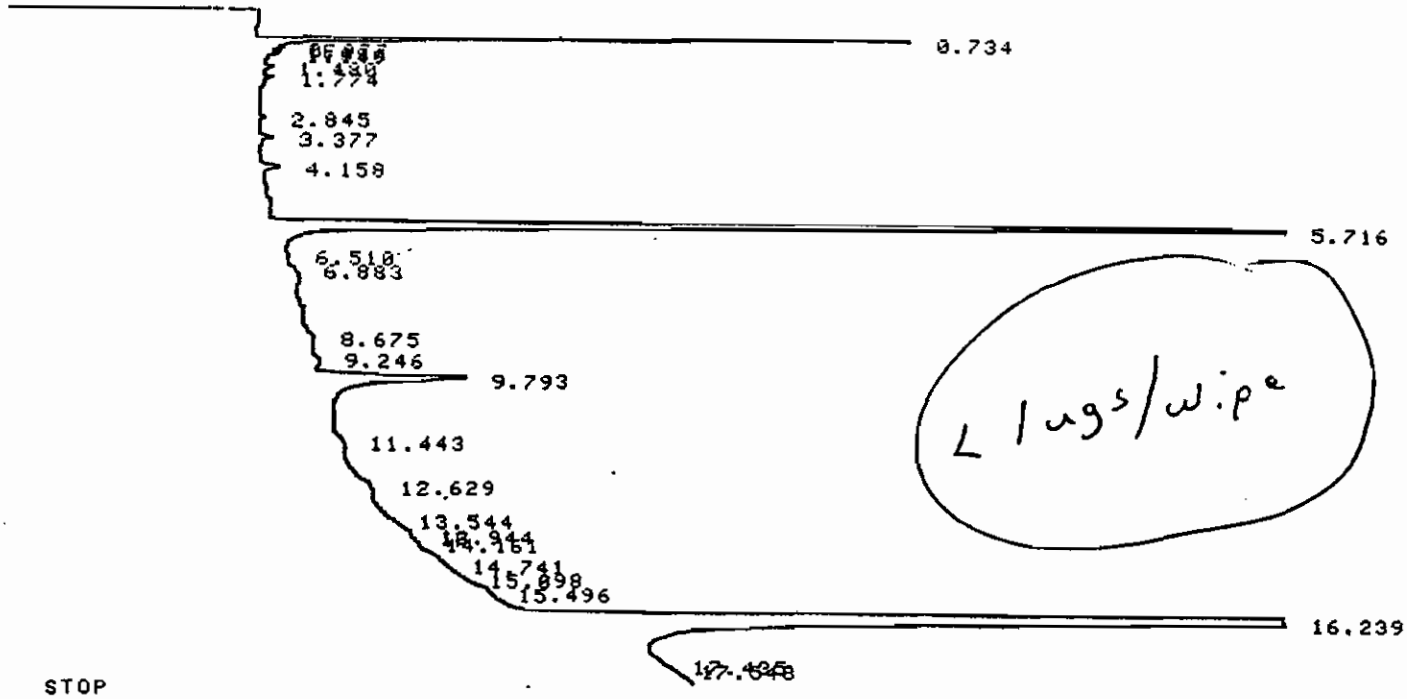
AREA%

RT	AREA	TYPE	WIDTH	AREA%
.732	34562	PB	.855	7.42710
.981	1063	BB	.045	.22843
1.145	965	BB	.040	.20737
1.475	697	PB	.059	.14978
2.030	1362	PB	.078	.29268
4.170	1906 I	PB	.135	.40958
4.648	29	PB	.044	.00623
5.724	40553	VB	.133	8.71452
6.509	2715	PB	.185	.58343
7.150	5015	PB	.268	1.07768
7.738	881	PB	.136	.18932
7.987	2273	BB	.140	.48845
8.688	4242	VB	.195	.91157
9.338	4194	PB	.213	.90126
9.791	20754	BB	.292	4.45987
11.739	6795	PB	.222	1.46019
12.608	615 I	PB	.123	.13216
13.395	22	PB	.122	.00473
13.633	15	BB	.083	.00322
13.880	43	BB	.072	.00924
14.160	563	PB	.168	.12098
14.804	2040	PB	.141	.43838
15.428	385	PB	.128	.08273
16.222	333588	PB	.208	71.68541
17.309	73	PB	.243	.01569

TOTAL AREA= 465350

$\frac{19320}{45537} \times 0.025 \times 20$
$= 0.2$ ugs/wipe

RUN #13340 AUG 19, 1997 13:27:50
START



STOP

RUN# 13340 AUG 19, 1997 13:27:50

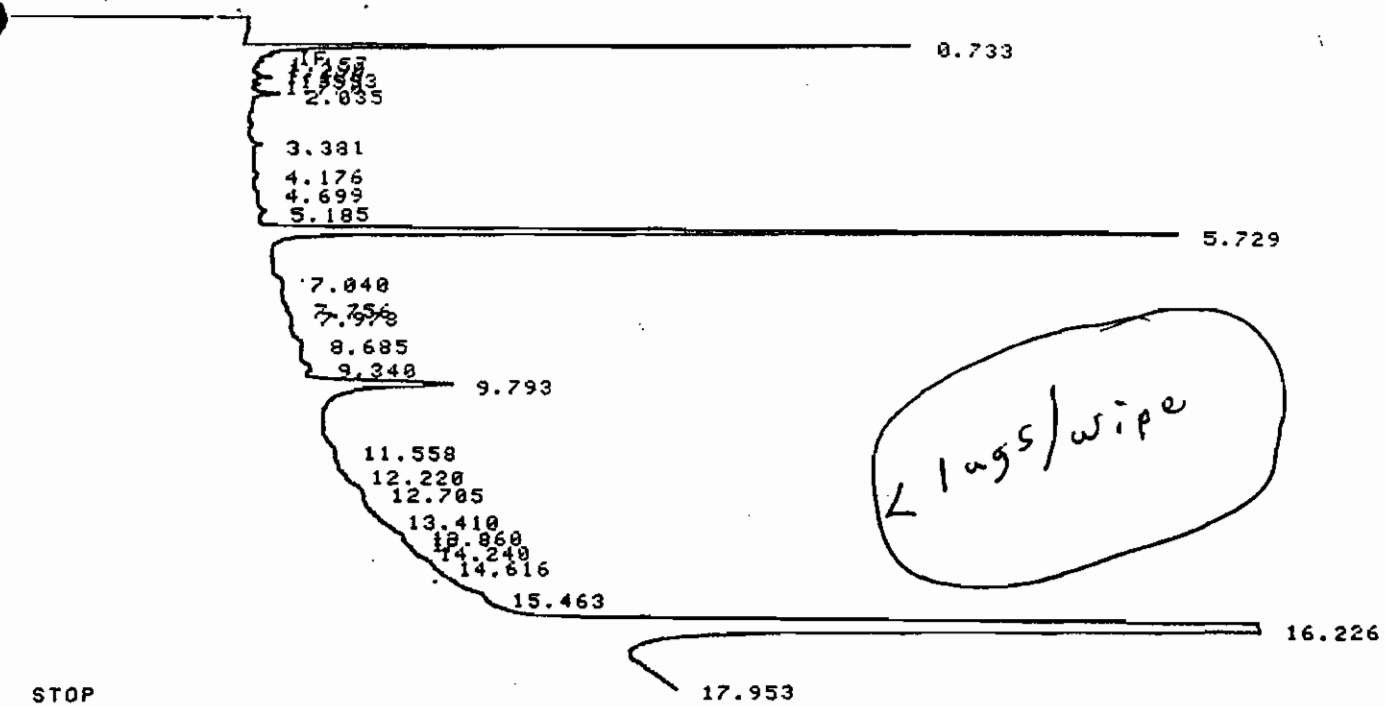
SAMPLE NAME: 818ED7 SAMPLE# 7

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.734	31710	PB	.058	6.19086
.977	319	BB	.053	.06228
1.090	209	BB	.026	.04080
1.149	318	BB	.039	.06208
1.480	497	PB	.067	.09703
1.774	564	PB	.063	.11011
2.845	536	PB	.080	.10465
3.377	1012	PB	.092	.19758
4.158	1948	PB	.127	.38031
5.716	158968	PB	.128	31.03589
6.510	464	PB	.143	.09059
9.793	28452	PB	.236	5.55478
11.443	68	PB	.227	.01328
13.944	573	BB	.503	.11187
15.496	404	BB	.962	.07887
16.239	286130	BB	.212	55.86218
17.548	35	BB	.194	.00683

TOTAL AREA= 512207
MUL FACTOR=1.0000E+00

RUN #13341 AUG 19, 1997 13:49:36
START



STOP

RUN# 13341 AUG 19, 1997 13:49:36

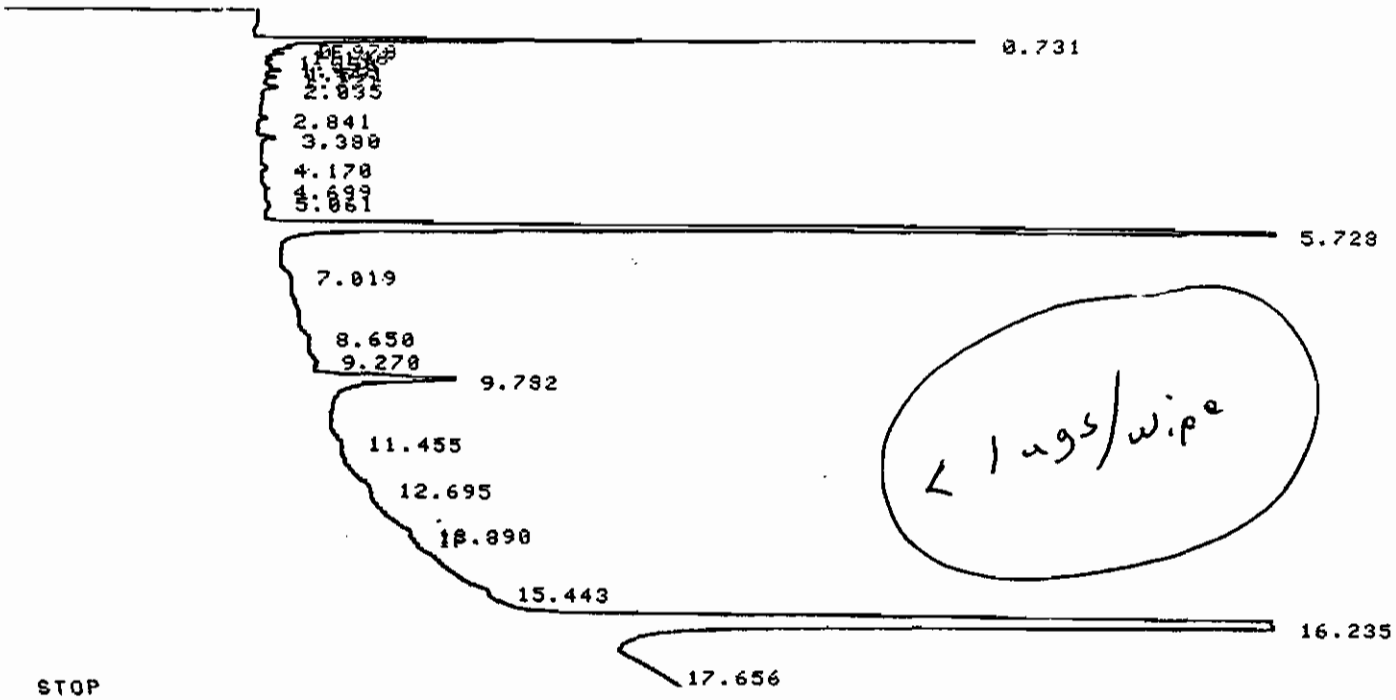
SAMPLE NAME: 818ED8 SAMPLE# 8

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.733	34301	PV	.060	6.27239
1.157	608	VB	.067	.11118
1.290	240	BB	.046	.04389
1.480	234	BB	.053	.04279
1.593	825	BB	.053	.15086
1.770	371	BB	.061	.06784
2.035	2405	BB	.101	.43979
3.381	639	PB	.089	.11685
4.176	642	PB	.135	.11740
5.185	879	VB	.163	.16074
5.729	100892	PB	.129	18.44943
7.040	106	PB	.088	.01938
7.756	78	PB	.130	.01426
8.685	936	VB	.175	.17116
9.793	28677	PB	.241	5.24397
11.558	99	PB	.413	.01810
12.705	147	PB	1.225	.02688
13.410	25	PB	.139	.00457
14.240	65	PB	.271	.01189
16.226	374502	PB	.214	68.48262
17.953	186	PB	.620	.03401

TOTAL AREA= 546857
MUL FACTOR=1.0000E+00

RUN #13342 AUG 19, 1997 14:11:24
START



STOP

RUN# 13342 AUG 19, 1997 14:11:24

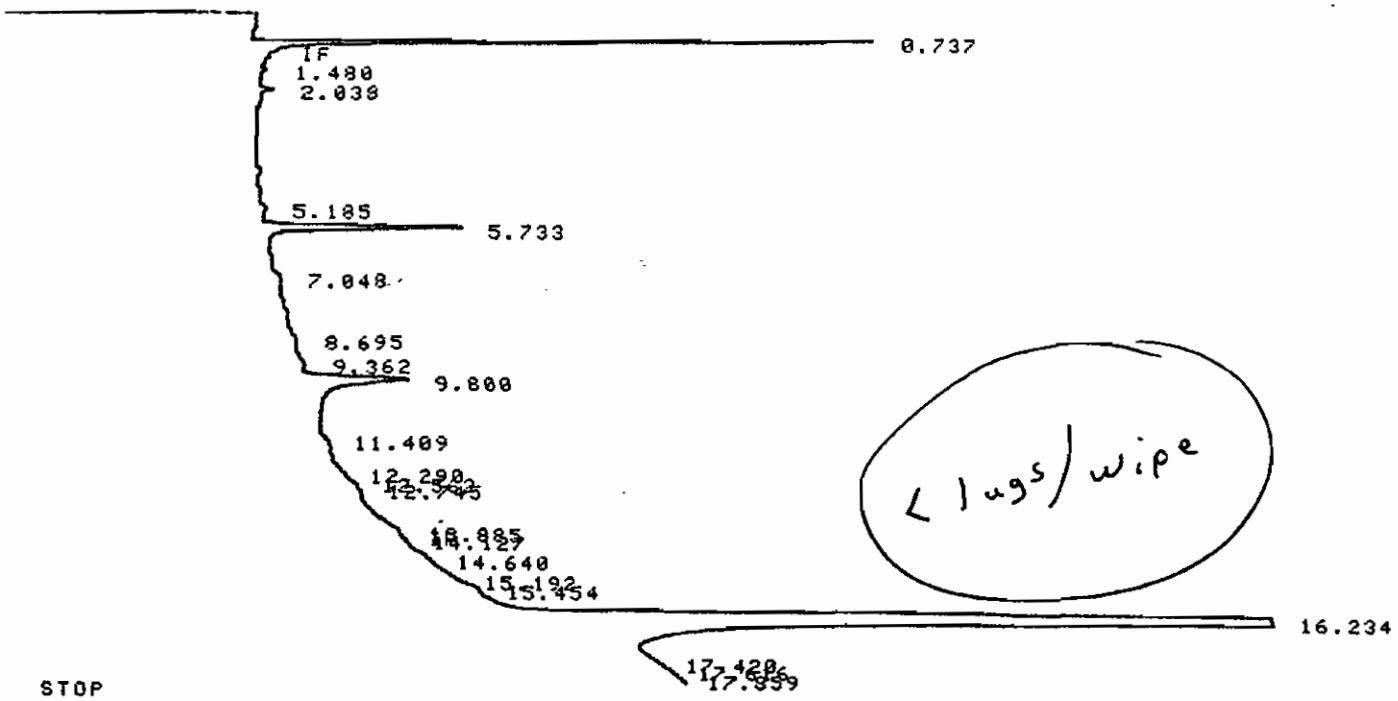
SAMPLE NAME: 818ED9 SAMPLE# 9

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.731	35975	PB	.059	5.83305
.978	312	BB	.042	.05059
1.146	454	BB	.042	.07361
1.290	280	BB	.055	.04540
1.478	205	BB	.044	.03324
1.591	505	BB	.046	.08188
1.771	621	BB	.061	.10069
2.035	1417	BB	.105	.22976
2.841	634	PB	.079	.10280
3.380	1229	BB	.089	.19927
4.170	796	VB	.126	.12906
4.699	568	PB	.141	.09210
5.728	174861	PB	.127	28.35228
7.019	71	PB	.091	.01151
9.782	23802	BB	.212	3.85930
13.890	250	PB	.076	.04054
16.235	374764	PB	.205	60.76493

TOTAL AREA= 616744
MUL FACTOR=1.0000E+00

RUN #13343 AUG 19, 1997 14:33:11
START



STOP

RUN# 13343 AUG 19, 1997 14:33:11

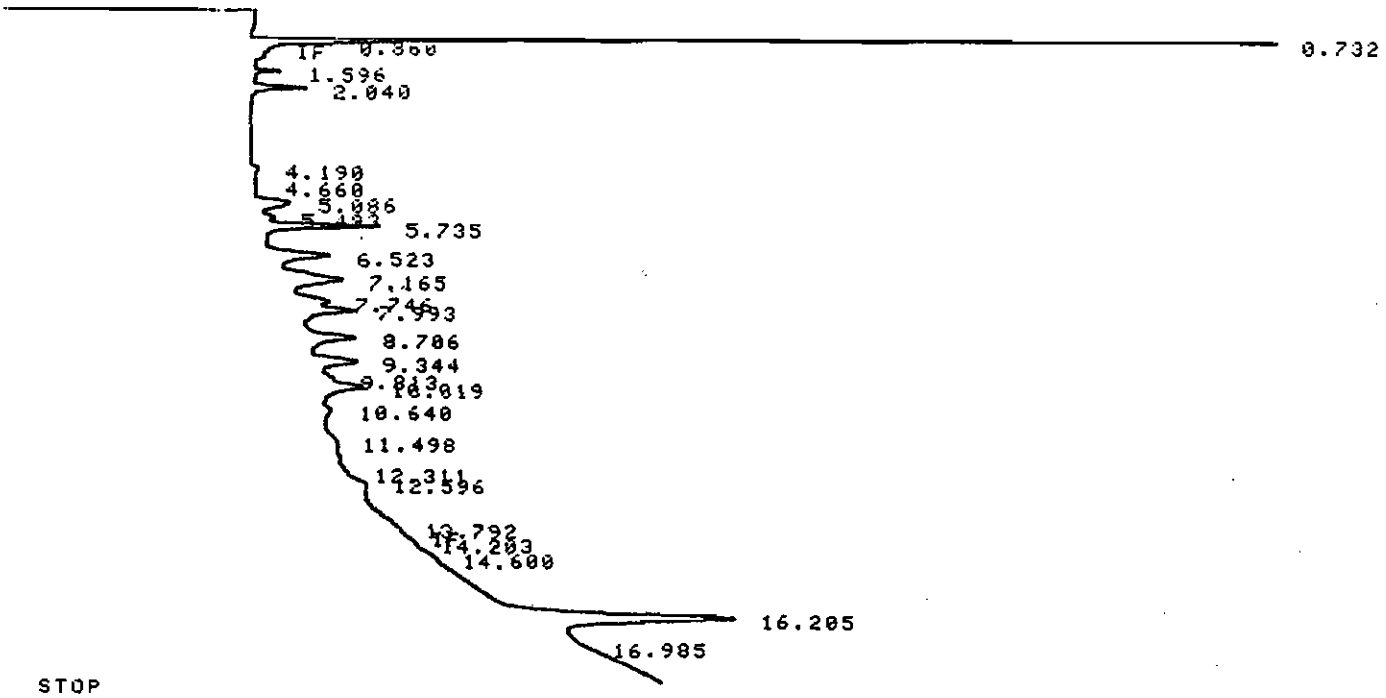
SAMPLE NAME: 818ED10 SAMPLE# 10
6

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.737	31498	PB	.060	7.79499
1.480	587	PB	.088	.14527
2.038	1257	PB	.098	.31108
5.185	761	VB	.159	.18833
5.733	22472	I PB	.136	5.56128
7.048	87	PB	.104	.02153
8.695	37	PB	.206	.00916
9.800	18264	PB	.222	4.51990
12.745	106	BB	.353	.02623
15.454	240	BB	.667	.05939
16.234	328656	PB	.223	81.33437
17.616	66	BB	.220	.01633
17.859	49	BB	.163	.01213

TOTAL AREA= 404080
MUL FACTOR=1.0000E+00

RUN #13344 AUG 19, 1997 14:55:01
START



STOP

RUN# 13344 AUG 19, 1997 14:55:01

SAMPLE NAME: 1254 0.025 SAMPLE# 11

AREA%

RT	AREA	TYPE	WIDTH	AREA%
.732	61599	PV	.033	36.42842
.860	4512	VV	.066	2.66831
1.596	1498	VB	.064	.88589
2.040	5091	PB	.111	3.01072
4.190	396	PB	.096	.23419
4.660	74	PB	.077	.04376
5.086	5719	BB	.210	3.38210
5.493	887	BB	.106	.52455
5.735	12678	BB	.135	7.49752
6.523	9805	PB	.208	5.79848
7.165	12149	PB	.267	7.18468
7.746	2156	PB	.150	1.27502
7.993	4837	BB	.149	2.86051
8.706	9434	PB	.227	5.57908
9.344	7146	PB	.207	4.22600
9.813	40	BB	.048	.02366
10.019	5472	BB	.192	3.23603
10.640	1126	PB	.174	.66589
11.498	144	PB	.800	.08516
16.205	24333	BB	.153	14.39005

TOTAL AREA= 169096
MUL FACTOR=1.0000E+00

45537 A11 51049