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Research and Development



# Regional Air Pollution Study

Gas Chromatography Laboratory Operation



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# REGIONAL AIR POLLUTION STUDY Gas Chromatography Laboratory Operation

by

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#### **ABSTRACT**

A gas chromatography laboratory was set up to analyze air samples collected in 100-liter Teflon bags and 8-liter stainless steel tanks. Samples were analyzed for total hydrocarbons (THC), methane, and CO on a Beckman Model 6800 gas chromatograph, and for  $\rm C_2\text{-}C_{10}$  hydrocarbons on a Perkin-Elmer Model 900 gas chromatograph equipped with dual columns, connected to a PEP-1 data system. A phenyl isocyanate column resolved the  $\rm C_2\text{-}C_5$  compounds and a support-coated-open-tubular squalane column was used for  $\rm C_6\text{-}C_{10}$  compounds. Experiments were performed to establish optimun temperature programming, flow rates, and column lengths to yield good compound resolution within reasonable elution times.

A total of 455 samples including replicates, were analyzed during the summer and fall, 1976. Of these samples, 292 were collected at 12 of the Regional Air Monitoring System (RAMS) sites to yield data on spatial and temporal distributions of hydrocarbons. All were analyzed for THC, methane, CO, and  $\rm C_2\text{-}C_5$  hydrocarbons. In addition, early morning (0600-0800) samples were analyzed for  $\rm C_6\text{-}C_{10}$  compounds. Portions of these data are included in this report. Sampling results indicate excessive concentration of ethylene within some of the RAMS stations, probably from the ozone monitoring system used at all RAMS sites. Some samples collected during roadway studies were reanalyzed on subsequent days; results showed good reproducibility using the Teflon bags. System reproducibility from quality control checks was good; with analyses of standards indicating deviations generally less than 5 percent.

All data, including the sums of paraffins, olefins, aromatics and total non-methane hydrocarbons are stored in the RAPS Data Bank at Research Triangle Park, North Carolina.

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#### 1.0 INTRODUCTION

The Regional Air Pollution Study (RAPS) is directed toward a quantitative understanding of urban pollution, including the monitoring of ambient levels of pollution, the gathering of micro-meteorological data and of a comprehensive emission inventory.

The gas chromatography laboratory was established to support these investigations. In particular it was charged with the collection and analysis of samples of hydrocarbons  $\mathrm{C}_1$  -  $\mathrm{C}_{10}$ , carbon monoxide and tracer gases in ambient air. In order to find a more accurate and selective method for identifying hydrocarbons  $\mathrm{C}_2$  -  $\mathrm{C}_{10}$ , modifications were made on the Perkin Elmer 900 Gas Chromatograph such that detailed analyses of these hydrocarbons in the parts-per-billion range could be performed. Samples were run and the results were tabulated and classified.

#### 2.0 SUMMARY

The gas chromatography laboratory modified the analysis of  $\rm C_2$  -  $\rm C_{10}$  hydrocarbons from the methods described in the final reports of Task Orders 53 and 103. A Perkin Elmer (P.E.) Model 900 gas chromatograph equipped with dual columns and a PEP-1 data system was used. A phenyl isocyanate (Dura Pak) column gave good resolution for  $\rm C_2$  -  $\rm C_5$  compounds and a Support-Coated-Open-Tubular (SCOT) squalane column was used to analyze hydrocarbons  $\rm C_6$  -  $\rm C_{10}$ . The modifications included shortening the SCOT squalane column from 30.48 meters to 15.24 meters and raising the initial temperature from -3°C to 25°C. These changes permitted the accurate and methodical identification of hydrocarbons  $\rm C_6$  -  $\rm C_{10}$  on the squalane column with an analysis time of 55 minutes. As result of these modifications the precision of the analyses done on the squalene column was improved as indicated by the reproducibility of the analysis of a toulene standard over a three month interval.

A total of 455 samples of ambient air (including triplicate analyses) was analyzed during the summer and fall intensives and for special studies during this task order period. These samples were collected from RAMS stations and roadside locations and were analyzed for total hydrocarbons (THC), carbon monoxide (CO) and methane (CH $_4$ ) on the Beckman Model 6800 gas chromatograph and hydrocarbons C $_2$  - C $_{10}$  on the P.E. 900 gas chromatograph.

Two hundred and ninety-two ambient air samples from twelve RAMS sites were analyzed during the summer and fall intensives. All of the samples were analyzed for THC, CH $_4$ , and CO (Beckman 6800) and C $_2$  - C $_5$  hydrocarbons (P.E. 900). The 0600-0800 (CST) samples were analyzed for THC, CH $_4$  and CO and C $_2$  - C $_{10}$  hydrocarbons (P.E. 900). Tables containing a portion of these data are in Appendix C.

High concentrations of ethylene at RAMS sites 101 and 124 led to a special study of ethylene concentration. The samples were collected in 100 litre Teflon bags (5 mil thickness) and 8 litre stainless steel tanks, in

and around the stations as requested by the EPA Task Coordinator. The results of this special study indicate that in certain cases the values reported for ethylene may be biased by fugitive emission of ethylene from the ozone monitor and associated hardware found in each of the RAMS stations.

Roadway samples were analyzed by the gas chromatography laboratory for a special study conducted by the EPA Task Coordinator. Some of the samples were reanalyzed on subsequent days. Relatively good reproducibility was obtained using the 5 mil Teflon bags.

To insure quality control, standards were run daily and records maintained of the areas and peak heights. The triplicate analyses of the first bag sample each day showed a standard deviation of less than 2.0 ppbC for the task order period.

All of these data, including the sums of the paraffins, olefins, aromatics and total non-methane hydrocarbons, were processed and prepared for entry into the RAPS Computer Data Bank, Research Triangle Park (RTP), North Carolina.

#### 3.0 DEVELOPMENT OF THE SQUALANE COLUMN

In the past, the gas chromatography laboratory experienced some difficulty in the analysis of  ${\rm C}_5$  -  ${\rm C}_{10}$  compounds on the squalane column. During Task Orders 53 and 103 periods the initial 60.96 meters x 0.05 cm squalane SCOT column was reduced to 30.48 meters because of excessive column bleed (1, 2). A Carbowax 20M-TPA on Chromosorb W-AW post column was added during Task Order 53 but was replaced with glass beads during Task Order 103 because of the problems incurred with the Carbowax column. A potassium carbonate ( ${\rm K}_2{\rm CO}_3$ ) pretrap was added during Task Order 53 in order to minimize the effect of water and polar compounds in the squalane system. Figure 1 is an example of hydrocarbons  ${\rm C}_2$  -  ${\rm C}_{10}$  analyzed during both task order periods. Because of the length of time required for analysis and the broadening of the peaks,  ${\rm C}_9$  -  ${\rm C}_{10}$  were not really identified. Alternate methods of analysis were explored.

The literature was searched and the problem examined in a systematic manner with the aim of developing a method of analysis that: (1) provided a reliable basis for identification of hydrocarbons on the squalane SCOT column; (2) could be easily incorporated in the equipment of the laboratory; and (3) preferably employed only one column for the analyses of compounds  $C_2 - C_{10}$ .

#### 3.1 LITERATURE SEARCH

Extensive work on the squalane SCOT column has been reported in the literature. W. O. McReynolds (3) and others described the analyses of hydrocarbons  $\rm C_2$  -  $\rm C_{10}$  on 60.96 and 30.48 meters x 0.05 cm columns. The problem encountered was the length of time required to complete the analyses since in some cases five or more hours were required. L. S. Ettre (4) described the analysis of  $\rm C_1$  -  $\rm C_9$  hydrocarbons on a 30.48 meter x 0.05 cm column requiring only 35 minutes. The method, although attractive, had some shortcomings: it did not identify all the compounds in which the gas

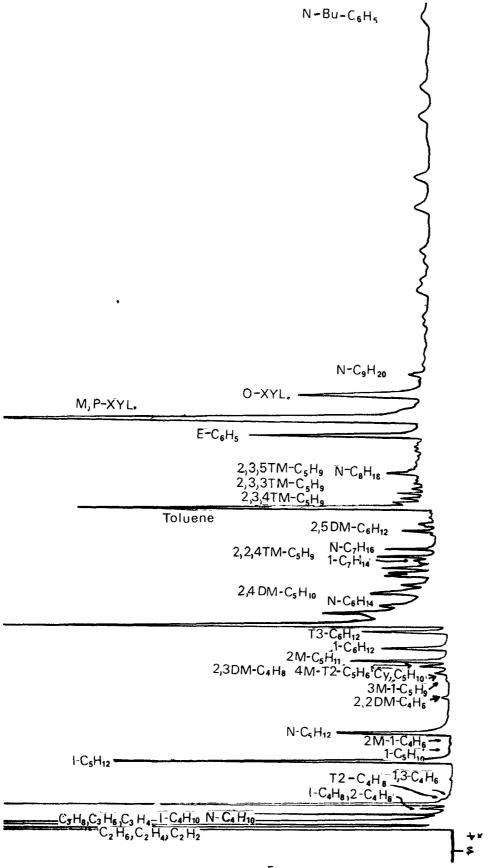


FIGURE 1. SQUALANE COLUMN ANALYSIS OF ATMOSPHERIC SAMPLE FOR  ${
m C_2}$  -  ${
m C_{10}}$  HYDROCARBONS

chromatography laboratory was interested, e.g. ethyl benzene, the xylenes and decanes; therefore, modifications had to be made.

#### 3.2 DEVELOPMENT OF THE 60.96 AND 30.48 METER COLUMNS

#### 3.2.1 Preliminary Experiments

Two 100 litre Teflon bags were filled with 50 litres of Scott-Marrin ultrapure air by using a mass flow meter which was corrected for standard temperature and pressure. Bag 1 was injected with 10 ml of each of the following gases using the appropriate size precision sampling syringe: methane (CH<sub>4</sub>), ethane (C<sub>2</sub>H<sub>6</sub>), ethylene (C<sub>2</sub>H<sub>4</sub>), propane (C<sub>3</sub>H<sub>8</sub>), acetylene (C<sub>2</sub>H<sub>2</sub>), isobutane (I-C<sub>4</sub>H<sub>10</sub>), normal butane (N-C<sub>4</sub>H<sub>10</sub>); and 0.05 ml each of the following liquids: isopentane (I-C<sub>5</sub>H<sub>12</sub>), normal pentane (N-C<sub>5</sub>H<sub>12</sub>), hexane (C<sub>6</sub>H<sub>14</sub>), heptane (C<sub>7</sub>H<sub>16</sub>), toluene (C<sub>7</sub>H<sub>8</sub>), octane (C<sub>8</sub>H<sub>10</sub>), ethyl benzene (C<sub>8</sub>H<sub>10</sub>), meta xylene (C<sub>8</sub>H<sub>10</sub>), ortho xylene (C<sub>8</sub>H<sub>10</sub>), nonane (C<sub>9</sub>H<sub>20</sub>), mesitylene (C<sub>9</sub>H<sub>12</sub>) and decane (C<sub>10</sub>H<sub>22</sub>). After equilibration, one milliliter of this mixture was injected into Bag 2 (dilution bag). The diluted contents of Bag 2 were analyzed on the 60.96 and 30.48 meters x 0.05 cm squalane columns.

The 60.96 and 30.48 meters squalane columns provided acceptable separation for hydrocarbons  $\mathrm{C_1}$  -  $\mathrm{C_7}$ ; but, the heavier hydrocarbons ( $\mathrm{C_8}$  -  $\mathrm{C_{10}}$ ) took an exceptionally long time to elute. The physical conditions (flow rate and temperature) were the same as described in the final report for Task Orders 53 and 103. In an attempt to overcome the elution problem with the heavier hydrocarbons, the initial oven temperature was varied from -3°C to 0°, 3°, 15° and 25°C and the final temperature from 75°C to 90°C and 115°C. The carrier gas flow rate was increased from 12 cc/min. to 15 cc/min. The results of these changes were not satisfactory; an increase in temperature and carrier gas flow rate yielded no data on  $\mathrm{C_1}$  -  $\mathrm{C_6}$  hydrocarbons (they eluted in the initial 15 seconds) and the higher temperatures yielded significant shifts in the baseline of the chromatograph at 90°C and above. These changes had little or no effect on the heavier hydrocarbons.

The 15.24 meter squalane SCOT column was then employed and was found suitable for the analyses of  $\rm C_6$  -  $\rm C_{10}$  hydrocarbons. It was found that hydrocarbons  $\rm C_1$  -  $\rm C_5$  elute too fast (within the first two minutes) to be

properly identified. The hydrocarbons were analyzed under the same physical conditions (temperatures and flow rates) as described for the 60.96 and 30.48 meters columns. The initial temperature was increased to  $25^{\circ}$ C and the final temperature increased to  $90^{\circ}$ C; the carrier gas flow rate remained at 12 cc/min. These conditions proved ideal for the analysis of hexane, which eluted at 3.6 minutes, through normal butyl benzene, which eluted at 55 minutes. It was then decided to continue to employ two columns in the P.E. 900 in the analyses of hydrocarbons  $C_2 - C_{10}$ . Figures 2 and 3 show typical examples of ambient air analyzed with the two columns employed in the P.E. 900 gas chromatograph.

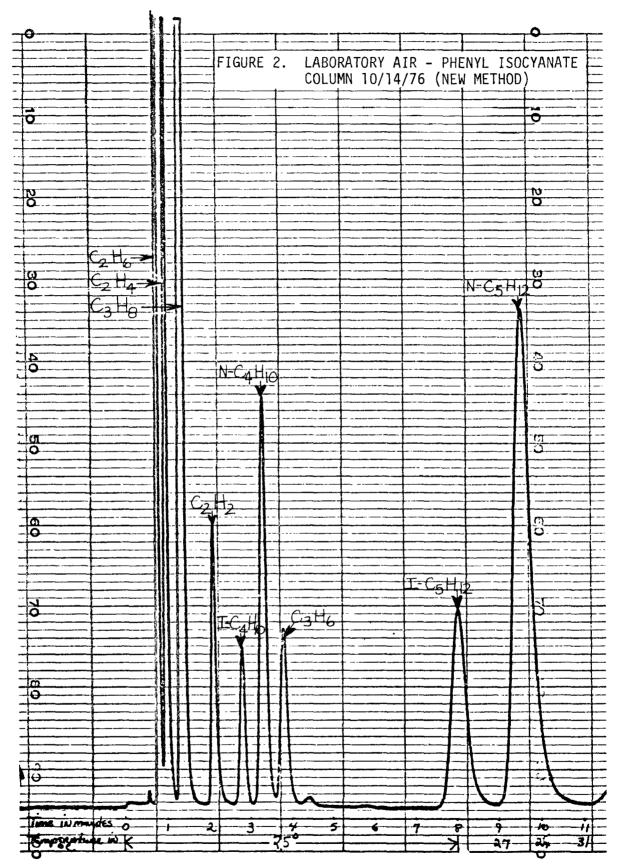
### 3.2.2 $C_2$ - $C_5$ Analysis on the Phenyl Isocyanate (Dura Pak) Column

During the preceding task order period (103)  $\rm C_2$  -  $\rm C_5$  hydrocarbons were analyzed on the 1.83 meter x 0.63 cm ID phenyl isocyanate (Dura Pak) column where the initial oven temperature was held at 0° for 3 minutes then programmed at a rate of 6°/minute until normal pentane (N-C $_5$ H $_{12}$ ), the last compound analyzed on the column, eluted. The whole analysis took 13 minutes to complete.

With the modified procedure, initial oven temperature at 25°C for 8 minutes, the programmed at a rate of 2°/minute and employing the Dura Pak column, normal pentane elutes in 10 minutes. Figures 2 and 4 show that the change in environment had no adverse effect on the  $\rm C_2$  -  $\rm C_5$  analysis.

#### 3.2.3 Use of Dual Columns

The analysis of hydrocarbons  $C_2$  -  $C_{10}$  was accomplished by using two columns connected to two flame ionization detectors in the Perkin Elmer Model 900 gas chromatograph (Figure 5 shows the schematic diagram of the analysis system). Hydrocarbons  $C_2$  -  $C_5$  were analyzed on a 1.83 meter x 0.63 cm ID phenyl isocyanate (Dura Pak) column that was conditioned by heating it at 115°C with helium flowing through over a 24 hour period. Hydrocarbons  $C_6$  -  $C_{10}$  were analyzed on a 15.24 meter x 0.05 cm ID squalane SCOT column that was conditioned at 115°C with helium flowing through over a 60 hour period. Both columns shared the same oven which was held at 25°C for eight minutes, then programmed at a rate of 2°C per minute until the



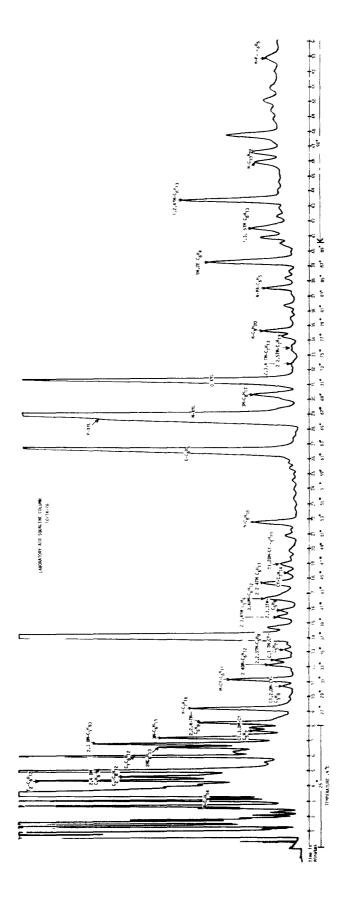


FIGURE 3. SAMPLE OF AMBIENT AIR ON THE SQUALANE COLUMN (NEW METHOD)

. 9

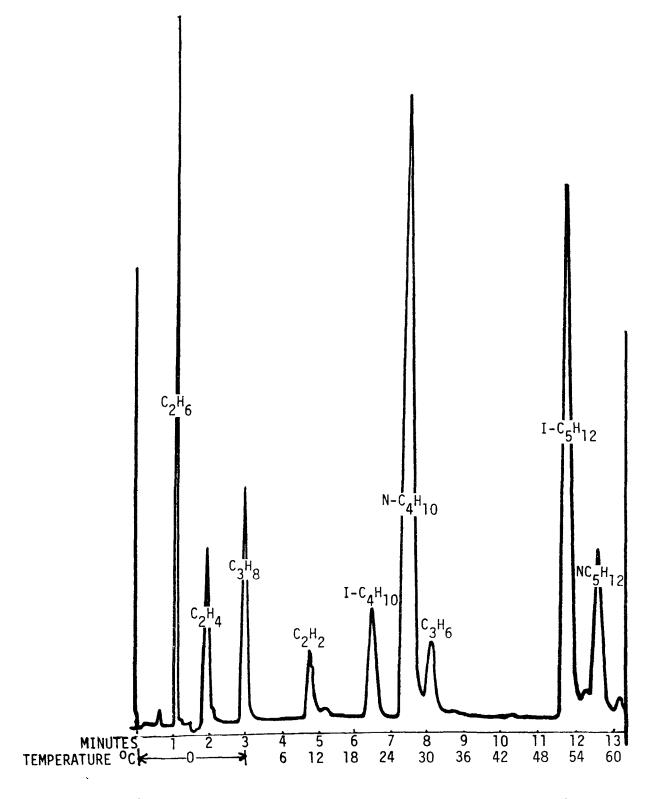


FIGURE 4. HYDROCARBONS  $\rm C_2$  -  $\rm C_5$  ON DURA PAK , 3 MINUTES AT 0°, PROGRAMMED AT 6°/MINUTE (Old Method)

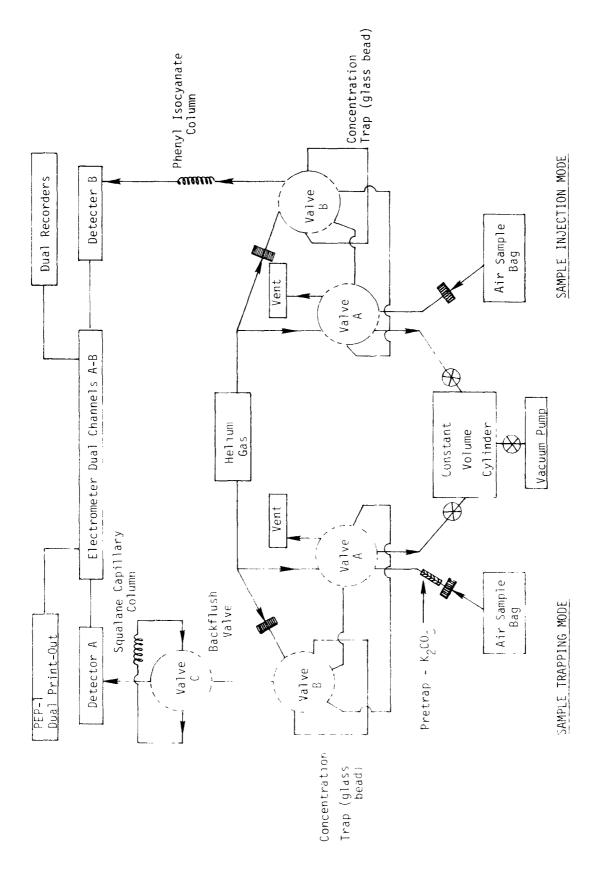


FIGURE 5. ATMOSPHERIC ORGANICS ANALYSIS SYSTEM - P.E. 900

system reached 90°C, where it was held for the remainder of the analysis. The total analysis times were 10 minutes for hydrocarbons  $\rm C_2$  -  $\rm C_5$  on the Dura Pak column and 55 minutes for hydrocarbons  $\rm C_6$  -  $\rm C_{10}$  on the squalane column.

# 3.2.4 Response Factors For Hydrocarbons $C_6 - C_{10}$

In order to obtain accurate response factors (RF) for  $\rm C_6$  -  $\rm C_{10}$  hydrocarbons for the squalane column, two synthetic mixtures of the various compounds were carefully prepared, using the procedure described in Section 3.2.1, then analyzed on the P.E. 900 gas chromatograph. The concentrations of the liquid compounds were computed in the following manner.

Example: Toluene

$$22.4 \times \frac{295^{\circ}}{273^{\circ}} \text{ 1/mole } \times \frac{1 \text{ mole}}{92.15 \text{ gm}} \times 0.8669 \frac{\text{gm}}{\text{m1}} \times 0.05 \text{ m1}$$
Bag-1 = 228 ppm = 50 litres of ultrapure air

Bag-2 
$$\frac{1 \text{ ml of contents of Bag 1 x 228 ppm}}{50 \text{ litres of ultrapure air}}$$
 x 7 carbon number = 31.9 ppbC

The PEP-1 computer was programmed in such a manner (area normalization) that only the areas of the selected compounds were computed. The response factors for the hydrocarbons were calculated in the following manner.

Example: Toluene

$$RF = \frac{\text{concentration}}{\text{area } \times 5}$$

$$RF = \frac{31.9}{2.044 \times 5}$$

$$RF = 3.12$$

Figure 6 shows an example of a chromatograph that was used in calculating the response factors for hydrocarbons  ${\rm C}_6$  -  ${\rm C}_{10}$ . It was observed that although equivalent amounts of each hydrocarbon were in the final mixture, the detector of the P.E. 900 indicated varying integrated peak areas. Table 1 shows experimentally determined response factors for  ${\rm C}_2$  -  ${\rm C}_{10}$  hydrocarbons. In instances where the pure compound could not be obtained for analysis, the response factors were estimated (Table 2).

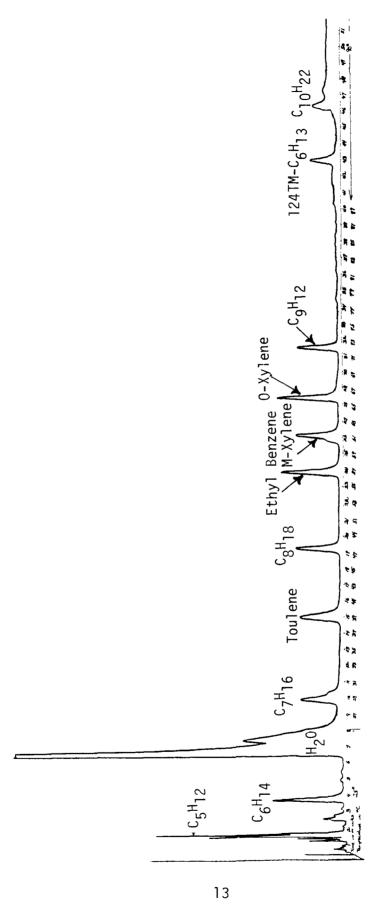


FIGURE 6. QUANTITATIVE STUDY OF HYDROCARBONS  $C_{\rm S}$  -  $C_{\rm 10}$  6/17/76

TABLE 1. EXPERIMENTALLY DETERMINED RESPONSE FACTORS FOR  $C_2$  -  $C_{10}$  HYDROCARBONS

2 10	
COMPOUNDS	RESPONSE FACTOR (RF)
Ethane	3.91
Ethylene	3.85
Propane	4.12
Acetylene	4.84
Isobutane	4.50
N-Butane	4.16
Propylene	4.36
Iso-Pentane	4.11
N-Pentane	4.10
N-Hexane	4.24
Trans 2-Hexene	3.72
Cis 2-Hexene	3.72
Cyclobexane	3.74
N-Heptane	2.98
Toluene	3.12
Octane	3.26
Ethyl Benzene	3.39
Methyl Xylene	3.61
Ortho Xylene	3.67
Nonane	2.99
N-Propyl Benzene	3.94
Mesitylene	3.88
N-Decane	4.34
N-Butyl Benzene	4.16

TABLE 2. ESTIMATED RESPONSE FACTORS FOR C<sub>6</sub> - C<sub>10</sub> HYDROCARBONS

COMPOUNDS	RESPONSE FACTOR (RF)	
Olefins $C_6$ to $C_{10}$	3.70	
Paraffins C <sub>6</sub> to C <sub>10</sub>	3.50	
Alkyl Aromatics	3.80	

# 4.0 ANALYSIS OF RAMS STATIONS SAMPLES DURING THE SUMMER AND FALL INTENSIVES OF 1976

During the period of performance of Task Order 113, the gas chromatography laboratory analyzed atmospheric samples from various RAMS stations. Sampling commenced on 23 June 1976 with a collection of two or four samples per site, five sites per day, two days per week. The choice of sites and times of sampling were determined by the EPA Task Coordinator. The sampling period ended on 18 November 1976.

#### 4.1 SELECTION OF SAMPLING BAGS AND TANKS

Ambient air samples were collected in Teflon bags and stainless steel tanks. The 100 litre Teflon bags (5 mil thickness) were checked for leaks by filling the bags with approximately 60 litres of helium, then going over them thoroughly (especially the seams and Teflon fittings) with a Gow Mac helium leak detector. The detector was zeroed, then the detector's nozzle was passed over the bag. If the gauge remained at zero, the bag was leak free; but, if the gauge moved from zero, the bag contained a leak. Leaks that occurred around the fittings were repaired by tightening them. with leaks in their seams were set aside to be repaired at RTP in Durham, N.C. The "leak free" bags, filled to capacity with helium, were left on the shelves overnight, then again checked for leaks. These bags were purged three times with helium then filled with approximately 60 litres of ultrapure air and again left on the shelves overnight. The bags were analyzed on the Beckman 6800 for THC,  $\mathrm{CH_4}$  and  $\mathrm{CO}$  and the P.E. 900 for  $C_2$  -  $C_{10}$  hydrocarbons. Bags possessing more than 2 ppbC of hydrocarbons were put aside; those having less than 2 ppbC hydrocarbon contamination were placed into circulation to be used in the RAMS sampling system. A supply of at least 25 "good bags" were kept in the system at all times. Bags were replaced in the system as needed. Figures 7 and 8 show the background chromatographs of a typical good bag.

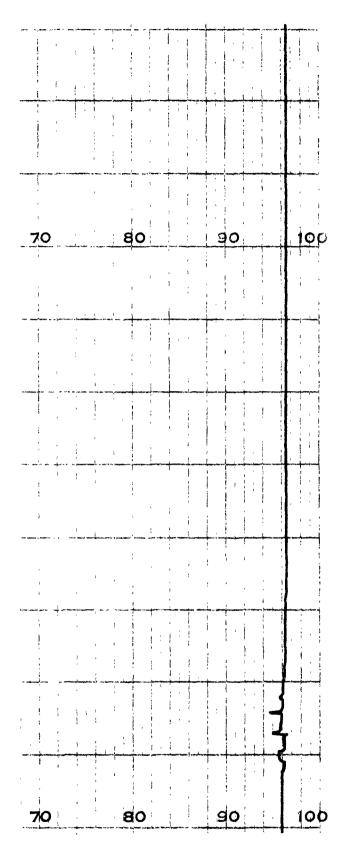
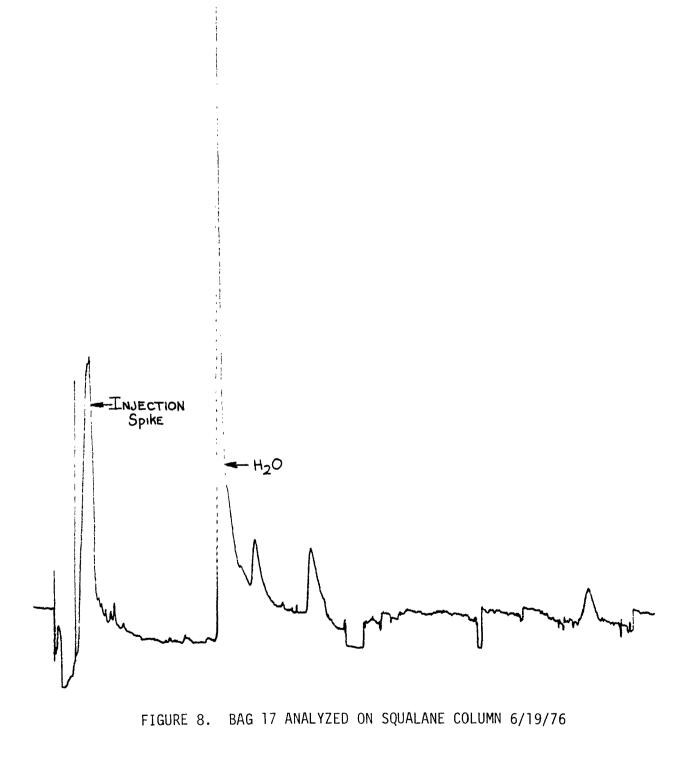


FIGURE 7. BAG 17 ANALYZED ON THE PHENYL ISOCYANATE COLUMN 6/9/76



During the 1976 fall intensive and special studies periods, four 8 litre stainless steel tanks were introduced into the sampling system. The tanks were fitted with a vacuum gauge on the rear and a two way needle valve on the front. The tanks were purged with helium, pressurized to approximately 30 psi with ultrapure air, and left to stand overnight in order to ascertain the amount of leakage and the degree of hydrocarbon contamination. The contents of the tank were analyzed on the P.E. 900 gas chromatograph the following day. The air in the tanks was removed by attaching a regulator with a quick connect fitting to the two-way valve. The valve was opened all the way so that fractionating of the gases would not occur and the regulator was partially opened. The quick connect fitting was joined to its counterpart on the P.E. 900 in injection mode (see Figure 5). Three of the four tanks were found suitable for sampling. Figure 9 shows an example of a chromatogram of a hydrocarbon-free stainless steel tank.

#### 4.2 AMBIENT AIR SAMPLING

The gas chromatography laboratory was responsible for the bag preparation, placing bags at the RAMS stations, sample transportation, sample analyses, data validation and computer tape and printout distribution of the data. These were accomplished as follows:

#### 4.2.1 Bag Preparation and Sample Collection

In order for the Teflon bags to be in place at the RAMS stations at 0600 hours CST, they were purged three times with helium and stored in a box at 0400 on the sampling day. The bags were transported to the various sites in a van and placed in the bag boxes at the RAMS stations. Figure 10 shows the sampling system and the flow diagram of the system at each site. A glass wool filter was used for the removal of ozone and particulants.

One bag was placed at each site in each bag box and upon command from the central computer at 0600 CST, the port between 1 and 2 would open on box 1 (see Figure 10) and air would flow through for two hours at an approximate rate of 833 cc/minute. (This rate varied somew twhat from station to station owing to the idiosyncrasy of the equipment at that station.) The port between 1 and 2 would open and ambient air would flow through 2

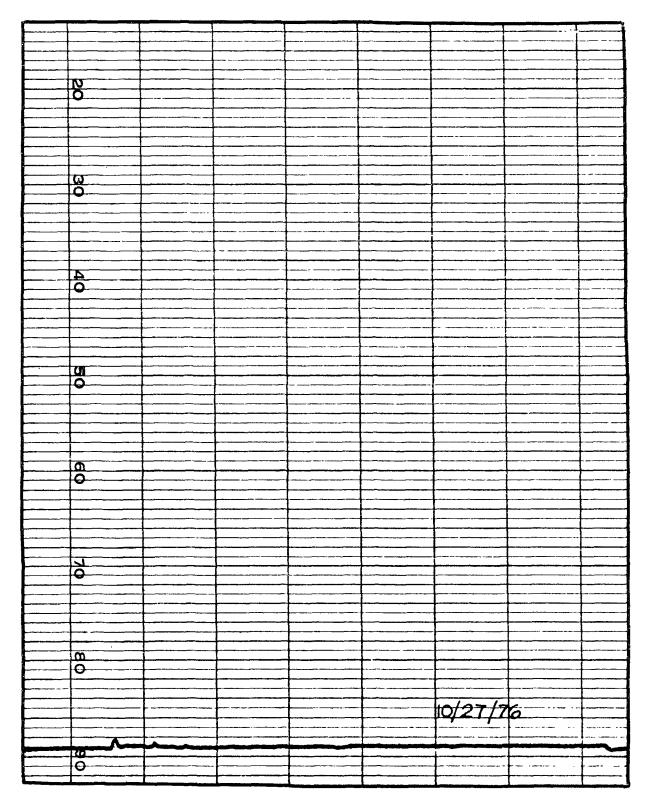
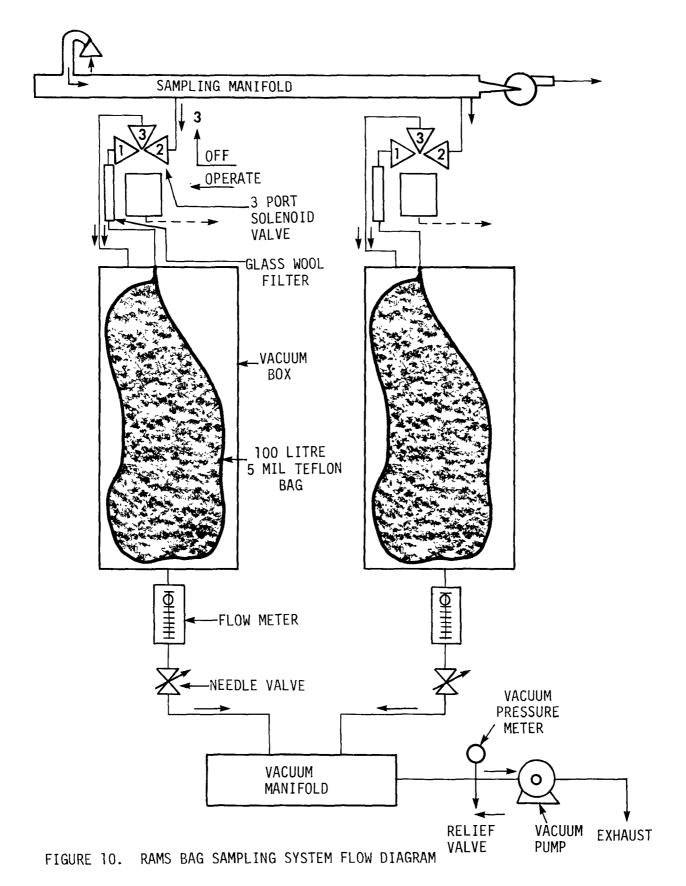


FIGURE 9. BACKGROUND OF STAINLESS STEEL TANK



to 1, then into the bag. Although port 3 was closed, the vacuum pump continued to evacuate the air in the air box; this caused the ambient air to fill the bag.

After two hours, a command from central would shut off the air flow on box 1 and turn on the air flow of box 2 and the procedure would be repeated.

The 0600 ambient air samples were removed and an empty bag put in its place in order to collect a sample at 1000-1200 hours. This procedure continued until the final samples were collected at all stations at 1200-1400 hours. The ambient air samples, protected from direct sunlight, were transported back to the laboratory for analyses.

#### 4.3 AMBIENT AIR SAMPLING

Ambient air samples, which were collected in the 100 litre Teflon bags at the RAMS stations over a two hour period of time, were analyzed for THC,  ${
m CH_4}$  and CO on the Beckman 6800 and for hydrocarbons  ${
m C_2}$  -  ${
m C_{10}}$  on the Perkin Elmer 900 (procedures are described in Appendices A and B). Initially all bag samples were analyzed for all compounds but this proved too difficult due to the late arrivals of the 0600-0800 and 0800-1000 samples at the laboratory (13:00-14:00 hours), the number of samples (maximum of 16) and the time required to complete the squalane analysis (1.5 hours including printout and turnaround times). The following changes were made by the EPA Task Coordinator: ambient air samples that were collected at 0600-0800 hours were analyzed for THC, CO and  $\mathrm{C}_1$  -  $\mathrm{C}_{10}$  and subsequent bags were analyzed for THC, CO and  $\mathrm{C_1}$  -  $\mathrm{C_5}$ . Table 3 shows the number of ambient air samples analyzed by the gas chromatography laboratory during this Task Order period. The analyses commenced immediately upon arrival of the bags at the laboratory and continued until all analyses were complete for all samples, which required the operation of two shifts at the laboratory.

#### 4.4 DATA VALIDATION, INTERPRETATION AND DISTRIBUTION

Each chromatogram of ambient air samples of the summer and fall intensive was visually analyzed by the EPA Task Coordinator and laboratory personnel working together. At the conclusion of the Intensive and data validation, the arithmetic mean and standard deviation of the concentrations of all time segments were calculated for total hydrocarbons (THC), methane ( $CH_q$ ),

TABLE 3. AMBIENT AIR SAMPLES ANALYZED IN THE GAS CHROMATOGRAPHY LABORATORY 1976

<u>MONTH</u>	NUMBER	
June	20	
July	98	
August	96	
September	9	
October	0	
November	69	
TOTAL	292	

carbon monoxide (CO) and acetylene ( ${\rm C_2H_2}$ ). Table 4 shows data on a station by station basis. The sums of the parafins, olefins, aromatics and total non-methane hydrocarbons per sample were calculated and recorded. The validated data, on one copy of a 600 foot, 9 track, 800 BPI, odd parity magnetic data tape, along with two copies of the printout were given to the EPA Task Coordinator. Appendix C contains some of the results of the data collected during the summer and fall intensives.

The carbon monoxide to acetylene ratios were computed from the arithmetic mean for each station where acetylene was consistently analyzed. These data are shown in Table 5. It should be noted that acetylene was not present in sufficient quantities to be analyzed at all times for many of the RAMS sites.

TABLE 4. SUMMARY OF SUMMER AND FALL INTENSIVES 1976\*

	THC		CH <sub>4</sub>	4	00		C <sub>2</sub> H <sub>2</sub>	
RAMS STATIONS	×	ν×	×	ر ×	×	۰×	×	××
Summer 101	3283	1137	1880	193	1101	530	12	8
Summer 102**	2660	306	1972	97	935	214		
Summer 103	2328	424	1860	273	763	412	4	2
Summer 108	2223	270	1872	98	603	388		
Summer 113	1874	78	1912	82	957	167		
Summer 114	2613	912	1851	308	599	302	7	6
Summer 115	2427	806	1788	147	498	213	က	2
Summer 118	2791	1310	1770	133	479	301	က	2
Summer 121	1960	311	1872	44	364	183		
Summer 122	2939	1771	1864	212	209	240	2	2
Summer 124	2264	382	1931	402	579	214	က	<b>,</b>
Fall 101	2906	788	1734	240	1815	1322	33	26
Fall 103	2676	817	1739	367	1073	844	15	14
Fall 114	2451	1299	1541	581	1187	1392	19	22
Fall 115	2321	417	1637	225	999	388	6	7

 $\overline{X}$  Mean S<sub>x</sub> Standard Deviation

\* Concentrations in ppbC

\*\* Values artificially high due to small amount of data

TABLE 5. AVERAGE CARBON MONOXIDE TO ACETYLENE RATIOS

RAMS STATION	RATIO OF CO TO C <sub>2</sub> H <sub>2</sub>
Summer 101	96
Summer 103	177
Summer 114	84
Summer 115	143
Summer 118	142
Summer 122	127
Summer 124	196
	$\overline{X} = 138  S_{X} = 40$
Fall 101	56
Fall 103	72
Fall 114	62
Fall 115	71
	$\overline{X} = 65$ $S_X = 8$

#### 5.0 SPECIAL STUDIES

Special studies and audits were carried out by the gas chromatography laboratory, as requested by the EPA Task Coordinator. These included road-way sampling, ethylene concentration verifications and audits on various gas samples.

#### 5.1 COLLECTION AND ANALYSES OF ROADWAY SAMPLES

The EPA Task Coordinator requested and collected roadway samples for the gas chromatography laboratory to analyze THC, CO and  $\rm C_1$  -  $\rm C_{10}$  hydrocarbons. A mobile air monitoring laboratory was used as the vehicle from which the samples were collected from heavily traveled thoroughfares during the morning rush hours. The bags used as containers for ambient air samples were purged three times with helium, filled with approximately 60 litres of helium, then left with the EPA Task Coordinator the day prior to sampling. Prior to sampling, a metal bellows pump was used to purge the bag of the helium, fill the bag with approximately 60 litres of ambient air, then purge the bag again. The 50-60 litre ambient air sample for analysis was collected by using the metal bellows pump with a Teflon sample line ( $\sim$  1.2 m long) extended out of the driver's side of the mobile laboratory for periods of 5-10 minutes. The samples were transported back to the laboratory for analysis. These roadway samples were collected mainly from I-270, I-40, Olive Road, Manchester Road and Babler Park. Detailed analytical results are contained in Appendix D.

The carbon monoxide to acetylene ratios were computed for each sample collected. The arithmetic mean of the ratio, 68.5, (Table 6) is similar to the carbon monoxide to acetylene ratio  $(63.4 \pm 6.1)$  computed for automobile emissions in the Lincoln Tunnel by EPA researchers (5).

TABLE 6. CARBON MONOXIDE TO ACETYLENE RATIOS OF ROADWAY SAMPLES

Location	Date	Ratio of CO to CoH2
Manchester Road	9/22/76	70
I-40	9/22/76	78
I-270	9/22/76	60
N on I-270 (I-40 to Olive)*	9/27/76	(41)
W on Manchester	9/27/76	55
W on Olive	9/27/76	76
N on I-270 Olive to Page	9/27/76	83
W on Manchester (I-270 to Weidman Road)	10/29/76	60
W on Manchester Weidman Road to Ries Rd.	10/29/76	61
S on I-270 (Manchester to I-44)	10/29/76	67
N on I-270 (I-44 to Manchester)	10/29/76	75
		$\overline{X} = 68.5$ $S_{X} = 9.35$
Babler Park**	9/22/76	96
Olive to Eatherton (west of airport	:)**9/22/76	61

<sup>\*</sup> Extremely high acetylene - not used in determining  $\overline{\mathbf{X}}$  and  $\mathbf{S}_{\mathbf{X}}$ 

<sup>\*\*</sup>Rural Samples

In order to ascertain the quality of data obtained from ambient air samples stored in Teflon bags over a period of time, the EPA Task Coordinator requested that some of the roadway samples be analyzed on the day collected and again 48 and 168 hours later. Table 7 shows samples collected from Manchester Road, I-40 and I-270 at the requested times.

#### 5.2 ETHYLENE STUDIES

Ethylene studies were carried out at RAMS stations 101 and 124 on 20, 22, 28 and 31 October 1976 and 1 and 4 November 1976 because of periodic abnormally high concentrations of that compound found in these stations during the summer intensive. Ethylene was of particular interest because of its use with the ozone analyzers at the RAMS stations. For these studies, the ambient air samples were collected in 100 litre Teflon bags and in 8 litre stainless steel tanks. The samples were taken from the following locations on various days:

- A. From RAMS station bag box with a sample taken outside simultaneously, 10-17 meters upwind and 2-3 meters above the ground using a metal bellows pump.
- B. Inside the station between the pump box and bag box.
- C. From the outlet of the center most blower of the pump box.
- D. From the outlet of the vacuum pump in pump box.

The bags and tanks were immediately transported back to the laboratory and analyzed on the phenyl isocyanate column of the Perkin Elmer Model 900 gas chromatograph for  ${\rm C_2-C_5}$  hydrocarbons. Detailed analytical results are contained in Appendix D.

#### 5.3 SPECIAL AUDITS

At the request of the EPA Task Coordinator, the cylinders MM11435, MM11437, MM11436 (from Scott-Marrin) and cylinder number FF3753 (from AIRCO Industrial Co.) were analyzed for CO, THC and CH<sub>4</sub> and RSG-80-4802 and RSG-80-7384 (from AIRCO Industrial Co.) for isobutane and normal hexane respectively. The audits were done in the following manner.

TABLE 7. STORAGE OF ROADSIDE SAMPLES IN TEFLON BAGS\*

	MANCHESTER ROAD		INTERSTATE 40		INTERSTATE I-270			
	†C9/22/76	10/21/76	MO/20/76	A9/22/76	10/21/76	A0 /22 /76	10/21/76	10.100.176
Dura Pak		A3/24/70	R9/20/70	A9/22/10	H9/24/70	A9/22/70	A9/24/76	A9/28/76
<sup>С</sup> 2 <sup>Н</sup> 6	22.3	22.2	24.0	18.9	19.1	42.0	47.1	44.9
C <sub>2</sub> H <sub>4</sub>	157.1	148.6	158.5	127.3	121.0	253.0	280.1	254.4
с <sub>3</sub> н <sub>8</sub>	27.6	26.7	29.0	14.4	14.2	54.8	61.5	56.8
C2H2	111.5	99.5	105.4	83.1	78.0	190.1	208.2	185.0
IsoC <sub>4</sub> H <sub>10</sub>	15.2	14.5	15.8	11.3	11.1	19.8	24.2	22.0
N-C4H10	91.0	88.0	94.0	68.0	66.0	131.0	149.0	137.0
С <sub>3</sub> Н <sub>6</sub>	73.7	69.8	74.2	60.3	57.3	112.7	125.0	113.5
IsoC <sub>5</sub> H <sub>12</sub>	125.0	119.0	130.0	84.0	81.0	172.0	191.0	179.0
N-C <sub>5</sub> H <sub>12</sub>	68.0	64.0	72.0	41.0	38.0	86.0	85.0	94.0
Squalane								
N-C <sub>6</sub> H <sub>14</sub>	46.0	41.0		64.0	27.0	38.0	87.0	
N-C <sub>7</sub> H <sub>16</sub>	15.0	15.0		32.0	14.0	51.0	28.0	
Toluene	72.0	71.0		165.0	67.0	239.0	140.0	
N-C <sub>8</sub> H <sub>18</sub>	12.0	14.0		16.0	12.0	45.0	25.0	
E-C <sub>6</sub> H <sub>5</sub>	30.0	36.0		68.0	37.0	107.0	66.0	
M-XYL	77.0	93.0		136.0	90.0	277.0	131.0	
0-XYL	30.0	36.0		65.0	32.0	91.0	56.0	
N-C <sub>9</sub> H <sub>20</sub>	4.0	4.0		4.0	1.0	21.0	10.0	
N-P-C <sub>6</sub> H <sub>5</sub>	5.0	8.0		6.0	2.0	18.0	10.0	
N-C <sub>10</sub> H <sub>22</sub>		9.0			5.0	41.0	25.0	

<sup>\*</sup> Concentration in ppbC

<sup>+</sup> C - Collected

<sup>§</sup> A - Analyzed

### 5.3.1 Total Hydrocarbon, Methane and Carbon Monoxide Audits

A five point calibration (Figure 11) was performed on the Beckman 6800 as described in Appendix A. The tanks to be audited were analyzed on the gas chromatograph and their concentrations determined from the 5-point calibration curve.

The results of the audit on 10/21/76 are shown in Tables 8 and 9.

TABLE 8. CARBON MONOXIDE AUDITS

Tank #	Manufacturer's Conc. in ppm	Rockwell Conc. in ppm	% Difference Rockwell vs. Manufacturer	EPA Conc. in ppm	% Difference Rockwell vs. EPA
MM11435	15.35	16.24	5.5	15.56	4.2
MM11436	8.00	8.31	3.7	8.17	1.7
MM11437	5.02	5.23	4.0	5.16	1.3
*FF3753		21.62		20.32	6.0
L-2359	5.06	5.33	5.1	5.29	0.7
L-1749	5.11			5.34	

<sup>\*</sup>Tank prepared by EPA at RTP

TABLE 9. METHANE AUDITS

Tank #	Manufacturer's Conc. in ppm	Rockwell Conc. in ppm	% Difference Rockwell vs. Manufacturer	EPA Conc. in ppm	% Difference Rockwell vs. EPA
MM11435	8.04	7.81	2.9	8.35	6.5
MM11436	4.98	4.83	3.0	5.23	7.6
MM11437	2.02	1.95	3.5	2.08	6.3
*FF3753		5.85		6.33	7.6
L-1749	1.99	1.81	9.0	2.09	13.4
L-2359	1.95	1.82	6.7	2.20	17.3

<sup>\*</sup>Tank prepared by EPA at RTP

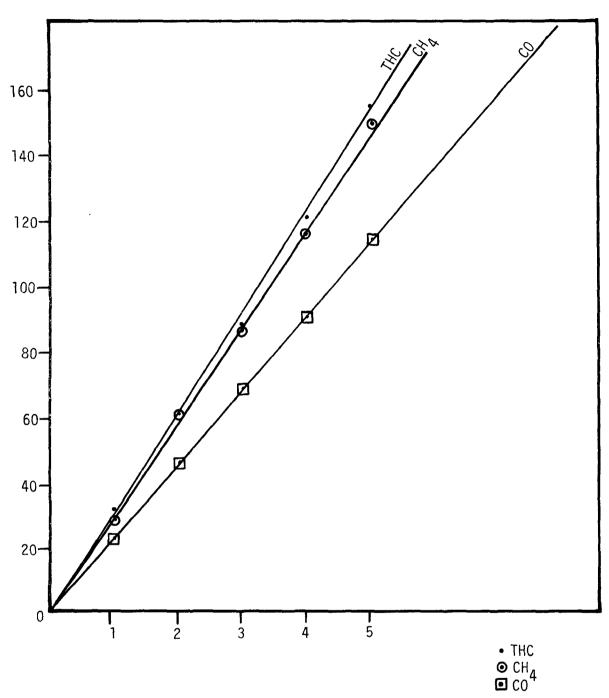


FIGURE 11. FIVE POINT CALIBRATION OF BECKMAN MODEL 6800 GAS CHROMATOGRAPH

#### 5.3.2 Isobutane, Normal Hexane and Synthetic Bag Mixture Audits

Audits were performed on AIRCO cylinders RGS-80-7384 and RGS-80-4802 containing isobutane and normal hexane respectively in nitrogen and on a synthetic mixture of carbon monocide and methane in ultrapure air. In order to perform these audits, primary standards were prepared of I-C $_4$ H $_{10}$  and N-C $_6$ H $_{14}$  as described in Appendix B and a 5-point calibration was performed to facilitate the analysis of the synthetic bag mixture of CO and CH $_4$  which was prepared by the EPA Task Coordinator. The results are given in Table 10.

TABLE 10. ISOBUTANE, N-HEXANE AND SPECIAL BAG AUDITS

	AIRCO RGS-80-7384	AIRCO RGS-80-4802
	Isobutane	Hexane
	Conc. in ppm	Conc. in ppm
Manufacturer	57.0	50.0
Rockwell	62.1	36.5
EPA	58.4	47.0

The results of the special bag samples are:

		Value of
	Measured	EPA Std.
CH <sub>4</sub>	4.5	4.59
CO T	8.9	9.32

#### 5.4 SPECIAL STUDIES AT RAMS 103 AND 107

Special studies were conducted at RAMS stations 103 and 107 in order to compare concentrations of compounds collected simultaneously in Teflon bags and stainless steel tanks; determine contamination at stations by collecting samples upwind; and to ascertain the accuracy of the data obtained from the gas chromatography laboratory. The samples were analyzed using the Dura Pak Column.

The bag and tank samples were collected by the same methods described in sections 4.2.1 and 4.1 respectively. The actual collection time varied from sample to sample, but bag and tank samples that were collected simultaneously

were collected over the same time interval. Tank samples were collected upwind of the stations.

Table 11 shows the results of replicate analyses of the samples, a comparisons of duplicate bag samples, and a comparison of station and upwind samples.

#### 5.5 DATA VALIDATION, INTERPRETATION AND DISTRIBUTION

Each chromatogram of ambient air samples collected for the roadway, ethylene and special studies at RAMS 103 and 107 was visually analyzed by the EPA Task Coordinator and laboratory personnel jointly. The validated data on one copy of a 6800 foot, 9 track, 800 BPI, odd parity magnetic data tape, along with 2 copies of the printout were given to the EPA Task Coordinator. These data contain the sums of the paraffins, olefins, aromatics and total non-methane hydrocarbons per sample and are furnished in Appendix D of this text.

## A. Reproducibility of Analyses

RAMS 103 Sample 103-1

Collection Time and Date: 7:15 - 9:05, November 24, 1976

Dura Pak Column. Concentrations in ppbC

Compound	First Analysis	Repeat Analysis	Duplicate Analysis	Average	Standard Deviation
С <sub>2</sub> Н <sub>6</sub>	20.0	19.7	20.2	20.0	0.25
C <sub>2</sub> H <sub>4</sub>	48.7	48.3	49.6	48.9	0.67
C <sub>3</sub> H <sub>8</sub>	56.4	55.4	57.3	56.4	0.95
C <sub>2</sub> H <sub>2</sub>	29.5	26.6	28.4	28.2	1.46
I-C <sub>4</sub> H <sub>10</sub>	16.1	16.6	17.0	16.6	0.45
N-C <sub>4</sub> H <sub>10</sub>	53.1	54.5	55.2	54.3	1.07
C <sub>3</sub> H <sub>6</sub>	9.6	11.3	12.8	11.2	1.60
I-C <sub>5</sub> H <sub>12</sub>	43.7	44.8	45.5	44.7	0.90
N-C <sub>5</sub> H <sub>12</sub>	15.1	17.2	15.7	16.0	1.08

RAMS 107 Sample 107-1

Collection Time and Date: 7:22 - 9:12, November 30, 1976

Dura Pak Column. Concentrations in ppbC

Compound	First <u>Analysis</u>	Repeat Analysis	Duplicate Analysis	Average	Standard Deviation
<sup>C</sup> 2 <sup>H</sup> 6	14.7	14.5	14.8	14.7	0.17
C <sub>2</sub> H <sub>4</sub>	47.3	46.9	47.6	47.3	0.30
C <sub>3</sub> H <sub>8</sub>	12.2	11.7	12.1	12.0	0.29
$C_2H_2$	37.3	38.8	43.1	39.7	3.01
I-C <sub>4</sub> H <sub>10</sub>	9.2	8.9	9.2	9.1	0.17
N-C4H10	41.4	40.8	42.3	41.5	0.75
C <sub>3</sub> H <sub>6</sub>	13.8	12.9	14.2	13.7	0.71
I-C <sub>5</sub> H <sub>12</sub>	40.5	40.6	40.6	40.6	0.06
N-C <sub>5</sub> H <sub>12</sub>	16.2	15.7	18.3	16.8	1.40

(continued)

#### B. Duplication of Samples

RAMS 107 Samples 107-1, 107-2

Collection Time and Date: 7:22 - 9:12, November 24, 1976

Dura Pak Column. Concentrations in ppbC

Compound	Bag #1	Bag #2	Average	Standard Deviation
С <sub>2</sub> Н <sub>6</sub>	29.3	29.8	29.6	0.35
C <sub>2</sub> H <sub>4</sub>	76.2	74.8	75.5	0.99
C <sub>3</sub> H <sub>8</sub>	25.3	25.7	25.5	0.28
C <sub>2</sub> H <sub>2</sub>	72.3	72.6	72.5	0.21
I-C <sub>4</sub> H <sub>10</sub>	23.5	20.6	22.1	2.05
N-C <sub>4</sub> H <sub>10</sub>	94.0	90.4	94.2	2.54
C3H6	27.5	28.4	28.0	0.64
I-C <sub>5</sub> H <sub>12</sub>	84.9	82.3	83.6	1.83
N-C <sub>5</sub> H <sub>12</sub>	35.2	34.9	35.1	0.21

RAMS 107

Samples 107-1, 107-2

Collection Time and Date: 7:22 - 9:12, November 30, 1976

Dura Pak Column. Concentrations in ppbC

Compound	Bag #1	Bag #2	Average	Standard Deviation
С <sub>2</sub> Н <sub>6</sub>	14.8	14.7	14.8	0.07
C <sub>2</sub> H <sub>4</sub>	47.3	56.1	51.7	6.22
С <sub>3</sub> Н <sub>8</sub>	12.2	12.9	12.6	0.49
C <sub>2</sub> H <sub>2</sub>	37.3	38.7	37.7	0.56
I-C <sub>4</sub> H <sub>10</sub>	9.2	8.8	9.0	0.28
N-C <sub>4</sub> H <sub>10</sub>	41.4	41.1	41.3	0.21
C <sub>3</sub> H <sub>6</sub>	13.8	13.4	13.6	0.28
I-C <sub>5</sub> H <sub>12</sub>	40.6	40.1	40.4	0.35
N-C <sub>5</sub> H <sub>12</sub>	16.2	15.9	16.1	0.21

(continued)

#### C. Comparison of Upwind and Station Data

**RAMS 103** 

Samples 103-1 (average), 103 Tank

Collection Time and Date: 7:15 - 9:05, November 23, 1976

Upwind Sample Collected in Stainless Steel Tank

Dura Pak Column. Concentrations in ppbC

Compound	Station Sample (average)	Upwind Sample (S.S. Tank)	<u>%                                    </u>
<sup>C</sup> 2 <sup>H</sup> 6	20.0	19.5	2.5
C <sub>2</sub> H <sub>4</sub>	48.9	27.5	43.7
C <sub>3</sub> H <sub>8</sub>	56.4	52.3	7.3
C <sub>2</sub> H <sub>2</sub>	28.2	26.7	5.3
$I-C_4H_{10}$	16.6	14.1	15.1
N-C <sub>4</sub> H <sub>10</sub>	54.3	47.0	13.4
<sup>C</sup> 3 <sup>H</sup> 6	11.2	11.1	0.9
I-C <sub>5</sub> H <sub>12</sub>	44.7	41.7	6.7
N-C <sub>5</sub> H <sub>12</sub>	16.0	15.0	6.2

RAMS 107

Samples 107-1 (average), 107 Tank Collection Time and Date: 7:22 - 9:12, November 24 and 30, 1976

Upwind Sample Collected in Stainless Steel Tank

Dura Pak Column. Concentrations in ppbC

		11/24/76		11/30/76		
Compound	Station Sample	Upwind Sample	<u>%                                    </u>	Station Sample	Upwind Sample	<u>% Δ</u>
С <sub>2</sub> Н <sub>6</sub>	29.6	30.2	-2.0	14.7	15.1	-2.7
C <sub>2</sub> H <sub>4</sub>	75.5	63.8	15.5	50.2	33.7	32.8
C <sub>3</sub> H <sub>8</sub>	25.5	25.1	1.6	12.3	12.4	-0.8
C2H2	72.5	76.2	-5.1	39.3	38.8	1.3
I-C <sub>4</sub> H <sub>10</sub>	22.1	19.4	12.2	9.0	10.7	-1.9
N-C <sub>4</sub> H <sub>10</sub>	94.2	85.2	9.6	41.4	42.3	-2.2
<sup>C</sup> 3 <sup>H</sup> 6	28.0	27.3	2.5	13.6	12.3	9.6
I-C <sub>5</sub> H <sub>12</sub>	83.6	76.7	8.3	40.5	41.7	-3.0
N-C <sub>5</sub> H <sub>12</sub>	35.1	31.8	9.4	16.6	15.0	9.6

#### 6.0 QUALITY CONTROL

To insure accurate results for all analyses done in the laboratory, quality control procedures were employed. The instruments were standardized using standard gases from Scott-Marrin for the Beckman 6800 and accurately prepared laboratory standards for the P.E. 900. (See Appendix B, Sec. 1.3) Records of area sizes and peak heights were maintained and checked daily, duplicate and repeat analyses were performed on the samples that were collected.

The first daily sample was analyzed three times in order to insure that the Perkin Elmer 900 was functioning properly. The first repeat analysis came directly after the original analysis, to provide a check on reproducibility of the machine; the second duplicate analyses were performed at the end of the day in order to ascertain whether the gas chromatograph maintained its stability. These duplications were usually within  $\pm$  5% of each other; if not, the samples were reanalyzed.

#### 6.1 STANDARDS FOR BECKMAN 6800

Scott-Marrin gases were chosen as standards because of the good reputation of the company and the availability of the THC, CH<sub>4</sub>, and CO gases in stable all-aluminum cylinders. EPA Quality Assurance Branch at the Research Triangle Park provided the calibrations used to determine the concentrations of the gases used as primary and secondary standards.

#### 6.2 STANDARDS FOR PERKIN ELMER 900

Appendix B describes the methods employed to prepare standards. Propane was used on the phenyl isocyanate column and toluene on the squalane column in the P.E. 900 gas chromatograph. The gases were stored in an stainless steel cylinders in order to insure their stability.

#### 6.3 DAILY STANDARDS

A tank containing laboratory air was used as a secondary standard.

Its composition was checked against the primary standards. The tank was refilled periodically because of the size of the cylinder and the volume of gas used per analysis, reanalyzed, then standardized against the primary standards.

Tables 12 and 13 show that from 20 July 1976 through 30 November 1976, the standard deviation of the analysis of propane, in two tank standards, were 0.08 ppbC and 1.07 ppbC with a standard error of 0.02 ppbC and 0.32 ppbC, respectively. For the same period, toluene exhibits a standard deviation of 0.58 ppbC and a standard error of 0.15 ppbC.

The comparison of areas of the standards was used as a quality control criterion as well as to detect the presence of leaks in the system. This approach provided a high level of confidence for the work done over this task order period.

TABLE 12. PROPANE STANDARD\*

DATE	AREA	DATE	AREA
7/20/76	14.7193	8/24/76	2.2904
7/22/76	13.7971	8/26/76	2.3351
7/27/76	16.7500	8/31/76	2.3192
7/29/76	16.7449	9/2/76	2.3775
8/3/76	16.7033	9/22/76	2.3649
8/4/76	17.0707	9/24/76	2.4480
8/5/76	16.9472	9/27/76	2.3218
8/10/76	17.1686	9/28/76	2.2637
8/12/76	16.7462	10/20/76	2.1580
8/17/76	16.5964	10/22/76	2.1924
8/19/76	16.7571	10/28/76	2.2105
	$\overline{X} = 16.36$	10/29/76	2.2111
	S <sub>x</sub> = 1.07	11/1/76	2.1606
	$S_{\bar{x}} = 0.32$	11/2/76	2.2017
	X	11/3/76	2.1917
		11/4/76	2.2104
		11/8/76	2.2781
		11/10/76	2.2583
		11/12/76	2.2524
		11/16/76	2.1953
		11/18/76	2.2167
		11/24/76	2.1489
		11/30/76	2.2672
			$\overline{X} = 2.26$
			$S_{X} = 0.08$
			$S_{\bar{x}} = 0.02$
			X

<sup>\*</sup> These areas are directly proportional to the response factors and the concentrations. Areas in arbitrary units.

TABLE 13. TOLUENE STANDARD\*

DATE	AREA	DATE	AREA	
7/20/76	0.9344	8/26/76	12.9900	
7/22/76	1.0363	8/31/76	13.2486	
7/27/76	1.8308	9/2/76	13.0400	
8/3/76	1.4842	9/22/76	13.0912	
8/5/76	1.6171	9/24/76	13.1302	
8/12/76	0.9089	9/27/76	13.2902	
8/17/76	1.4664	10/29/76	13.1078	
8/19/76	0.8082	11/8/76	12.6534	
<del>-</del>	$\overline{X} = 1.26$	11/10/76	12.6560	
	$S_{x} = 0.38$	11/12/76	12.5260	
	$S_{\overline{x}} = 0.14$	11/16/76	12.1836	
	X	11/18/76	13.0540	
		11/29/76	14.1280	
		11/30/76	14.3846	
		$\overline{X}$	= 13.1100	
		S	_ = 0.58	
			$\frac{x}{x} = 0.15$	
		•	^	

<sup>\*</sup> These areas are directly proportional to the response factors and the concentrations. Areas in arbitrary units.

#### 7.0 DATA PROCESSING

Data generated by the RAPS Gas Chromatography Laboratory were processed and submitted to the RAPS Computer Data Bank, Research Triangle Park (RTP), North Carolina. Data processing from analysis to submission to the data bank was performed as described in the following sections.

#### 7.1 DATA TABULATION

It was planned to perform approximately thirty-two analyses per week for up to sixty components. The data were initially recorded in the form of strip chart chromatograms, punched tape and/or teletype printouts. Next, the data were given a first quality review by visitally inspecting the data for general chromatographic form. Quantitative values for each component were then established. Following review and approval, the data were tabulated on a special preprinted form for keypunching.

#### 7.2 KEYPUNCHING AND PROCESSING

At the end of the analysis period, the data forms were presented to the EPA Task Coordinator for review and approval and subsequently keypunched and keypunch validated. Keypunching errors were corrected by computer operators at the RAMS Computer Facility.

Data processing entailed checking the cards for index number consistency, as provided for by the form, and then producing a triple copy printout of labelling information: the name, code number, concentration (PPB), ratio relative to CO, and flags if the concentration or ratio was outside an upper and lower set of limits provided by EPA. Four quantities aggregated by the software were treated as components in all respects: sum of nonmethane paraffins, olefins, aromatics, and nonmethane hydrocarbons. Validation of the data was carried out by visual inspection and comparison of the data with the chromatogram and original tabulated data. Also, special attention was directed to flagged data for validity and proper annotation.

Upon completion of data validation, three copies of a 600 foot, 9 track, 800 BPI, odd-parity magnetic data tape were prepared. One copy was sent to RTP, along with a copy of the printout. The other copy of the tape and a printout was delivered to the EPA Task Order Coordinator (St. Louis) and a third copy of the printout and tape retained by the RAMS Central Computer Facility.

#### LITERATURE CITED

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- 5. Lonneman, et al, Environmental Science and Technology, Vl. 8, pg. 231.

# APPENDIX A OPERATION OF BECKMAN 6800 GAS CHROMATOGRAPH

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#### A.O TOTAL HYDROCARCONS, CARBON MONOXIDE AND METHANE ANALYSES

Analyses of CO, THC and  $\mathrm{CH}_4$  were done on a Beckman Model 6800 Gas Chromatograph in the following manner:

The recorder was turned on, set at 10 mv range and zeroed with the zero volt button. The auto-zero switch was actuated in order to zero the electrometer of the chromatograph. The pump on the back of the chromatograph was turned on and a sample bag was connected to commence analyses.

To analyze total hydrocarbons, the attenuation was normally set at 4 and the range at 10 and the valve B toggle switch was actuated for approximately 15-20 seconds. The recorder was allowed to return to zero, aided with the auto-zero toggle switch. The attenuation was changed from 4 to 1 for the methane and carbon monoxide analyses and the valve A toggle switch actuated for 45 seconds.

#### A.1 REPRODUCIBILITY OF THE BECKMAN 6800

Reproducibility of the results of the analyses of THC,  $CH_4$  and CO was checked by running duplicate analyses; that is, the first sample and the standard were rerun after the analysis of all other samples. During the course of this task order, these duplicates agreed closely with the initial analysis (+5%).

#### A.2 LINEARITY OF DETECTOR RESPONSE

Five point calibrations were carried out in duplicate to insure that the detector response was linear with respect to sample concentration. The tests were performed for THC, CH<sub>4</sub> and CO. These checks were determined from synthetic bag mixtures prepared in the laboratory. The procedure was as follows:

Five bags were prepared. The bags were labeled and filled with 100 liters of zero air using a mass flow meter. The appropriate amounts of  $\mathrm{CH}_4$  and  $\mathrm{CO}$  were injected into each bag using a gas-tight calibrated syringe and allowed to diffuse throughout the bag. The bags were then analyzed. The make-up of the mixtures is shown below:

BAG #	ZERO AIR	CH <sub>4</sub>	<u>co</u>	CONC.
1	100 litres	0.1 ml	0.1 ml	1 ppm
2	100 litres	0.2 ml	0.2 ml	2 ppm
3	100 litres	0.3 ml	0.3 ml	3 ppm
4	100 litres	0.4 ml	0.4 ml	4 ppm
5	100 litres	0.5 ml	0.5 ml	5 ppm

#### A.3 PRIMARY STANDARD

A cylinder containing small amounts of methane and carbon monoxide in air (Scott-Marrin L 1749) was analyzed by the EPA at Research Triangle Park, North Carolina. Its concentrations was computed to be  $5.33~\rm ppm$  CO and  $1.99~\rm ppm$  CH $_4$ . This cylinder served as the primary standard for the Beckman  $6800~\rm and$  was used to calibrate other tanks.

#### A.4 SECONDARY STANDARDS

Two other cylinders (Scott-Marrin L 2327 and L 2359) served as secondary standards for the Beckman 6800 during this task order period. When analyzed against the primary standard Scott-Marrin L 1749, their concentrations were computed to be 5.28 ppm CO, 4.38 ppm  $\rm CH_4$  and  $\rm THC$  for Scott L 2327 and 5.29 ppm CO and ppm  $\rm CH_4$  and 2.20 ppm  $\rm CH_4$  and  $\rm THC$  for Scott L 2349. The secondary standards were checked against the primary ones once a month.

#### A.5 CALCULATIONS

Calculations of the concentrations of THC, CH<sub>4</sub> and CO in various samples are straight forward and were done by hand. When analyses were to be performed, the secondary standard was analyzed; that is, its peak height was measured. Since its concentration was known, it was possible to compute the concentration of unknown once its peak height was measured:

 $\frac{\text{Standard concentration in ppm}}{\text{Peak height of standard in mm}} \ x \ \text{peak height of unknown in mm} = \frac{\text{concentration}}{\text{of unknown}}$ 

Initially, a sample of standard gas was injected into a Teflon bag, then analyzed on the Beckman 6800. It was observed that the peak height varied as much as 42.6% for THC, 9.7% for  $\mathrm{CH_4}$  and 58% for  $\mathrm{CO}$ . These variations were unacceptable; therefore, another method was devised by which the standard gas was passed directly into the Beckman 6800. With this procedure, maximum peak height variation for THC was 8.4%,  $\mathrm{CH_4}$  was 14.9%, and  $\mathrm{CO}$  was 17.9%. The standard deviations were computed to be 4.88 mm (3.8%) for THC, 7.25 mm (6.5%) for  $\mathrm{CH_4}$  and 9.89 mm (9.2%) for  $\mathrm{CO}$  as shown in Table A-1.

	ABLE A-1. TANK L-2	<u> 327 - TANK STANDARD</u>	FOR 6800
DATE	THC	CH <sub>4</sub>	CO
6/23/76	129.5*	105.0	102.0
6/28/76	127.0	103.5	100.5
7/06/76	127.0	106.5	92.0
7/08/76	142.5	104.0	98.5
7/13/76	132.0	102.0	85.5
7/15/76	115.5	97.0	86.8
7/20/76	130.0	106.0	103.8
7/22/76	129.5	112.0	110.7
7/27/76	122.6	117.3	117.0
7/29/76 am	124.8	106.6	96.7
7/29/76 pm	124.7	121.8	122.4
8/03/76 am	126.4	115.1	112.8
8/03/76 pm	129.1	115.0	109.0
8/03/76	125.8	116.5	114.6
8/05/76	126.4	116.9	116.5
8/10/76 <b>a</b> m	132.9	102.3	106.5
8/10/76 pm	130.1	112.5	106.9
8/12/76 <b>a</b> m	128.4	120.9	121.0
8/12/76 pm	127.1	118.3	119.1
8/24/76	127.0	117.2	113.3
8/26/76 am	125.1	117.5	114.7
8/26/76 pm	125.1	118.7	115.8
	$\overline{X} = 127.7$	X = 111.48	$\overline{X}$ = 108.00
	s <sub>x</sub> = 4.88	$S_{x} = 7.25$	S <sub>X</sub> = 9.89
	$S_{\bar{x}} = 1.04$	S <sub>₹</sub> = 1.54	$S_{\bar{x}} = 2.11$

<sup>\*</sup> Response in mm

## APPENDIX B

## OPERATION OF THE PERKIN ELMER 900 GAS CHROMATOGRAPH

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# B.O C<sub>2</sub> - C<sub>10</sub> HYDROCARBON ANALYSIS

The RAMS bag samples were analyzed for  $\mathrm{C_2}\text{-}\mathrm{C_{10}}$  hydrocarbons on the PE 900 gas chromatograph. The ambient air samples were analyzed and the concentrations of the compounds determined by comparing the samples to the standard. One sample was analyzed in duplicate and in some cases triplicate to insure the reproducibility of the entire set of samples and the precision and accuracy of the gas chromatograph.

#### B.1 GENERAL PROCEDURES

The hydrogen and air sources were turned on and the flame ionization detectors lighted. The oven was cooled from 90°C (overnight and weekend temperature) to 25°C by turning on the liquid nitrogen. The recorders were turned on, checked for positive response, then zeroed; the vacuum line was turned on. Directives of identification and sampling time were entered on the teletype to the PEP-1 integrator.

The stainless steel concentration trap was placed in a Dewar flask which contained liquid oxygen. The appropriate vacuum line was connected to the vacuum source and the bag or tank connected to the inlet line. The vacuum source was opened and a small sample was drawn through the inlet line to purge the line of any air from a previous sample. The inlet valve was switched to the trapping mode and a measured volume of the sample was cold-trapped into the liquid oxygen cooled concentration trap.

After the sample was trapped, the ready light on the PEP-1 interface was actuated, the valve switched to the injection mode and a Dewar flask of hot water applied to the concentration trap. The "ready light" on the gas chromatograph was immediately pressed to initiate the temperature programming. The chromatograph was programmed to hold  $25^{\circ}$ C for 8 minutes. The temperature was then raised at a rate of  $2^{\circ}$ C per minute to  $90^{\circ}$ C, where it was held for the remainder of the run. The valve was returned from the injection mode to the backflush (neutral) position after 2 minutes. The analyses times were  $10^{\circ}$ 

minutes for the phenyl isocyanate column ( $\rm C_2-\rm C_5$ ) and 55 minutes for the squalane column ( $\rm C_5-\rm C_{10}$ ).

Upon completion of the analyses, the PEP-1 integrator tabulated the area measured, the concentration and the identification of peaks. The reset button on the gas chromatograph was pressed and the above procedure was repeated. The analyses and turnaround time for trapping sample, analysis and computer printout times for  $C_2$ - $C_5$  were approximately 0.5 hours on the phenyl isocyanate column and 1.35 hours for  $C_5$ - $C_{10}$  on the squalane column. The files of the memory bank of the PEP-1 computer were normally erased at the end of the working day.

#### B.2 REPRODUCIBILITY OF THE PHENYL ISOCYANATE COLUMN

To ascertain and maintain reproducibility of this column, the propane standard and triplicate sample analyses were used as indicators. Any major change in the area size of the propane standard or major deviation from the concentrations of nine compounds analyzed on the column signaled problems in the system. The magnetic tapes and the computer printouts demonstrate the reproducibility of the triplicate analyses done on the first daily bag samples. The reproducibility of the daily duplicate and triplicate analyses were generally within  $\pm$  5%.

#### B.3 LINEARITY OF DETECTOR RESPONSE OF PHENYL ISOCYANATE COLUMN

Five point calibrations were carried out on the P.E. 900 is insure that the detector response was linear with respect to sample concentration. Five mil (0.005 inch) Teflon bags were filled with ultrapure air and injected with accurately measured quantities of propane and toluene. After allowing half an hour for diffusion, samples were withdrawn from this bag ("A") and injected into five other bags, each of which had been filled with 50 l of ultrapure air according to the schedule shown on the following page:

Bag #	Zero Air	<u>C<sub>3</sub>H<sub>8</sub>(99% purity)</u>	Conc.	Toluene (99% purity)	Conc.
Α	50 liters	5 ml of propane	100 ppm	0.05 ml toluene	226.9 ppm
7	50 litres	1 ml from Bag A	6 ppbC	l ml from Bag A	31.9 ppbC
2	50 litres	2 ml from Bag A	12 ppbC	2 ml from Bag A	63.8 ppbC
3	50 litres	3 ml from Bag A	18 ppbC	3 ml from Bag A	95.7 ppbC
4	50 litres	4 ml from Bag A	24 ppbC	4 ml from Bag A	127.6 ppbC
5	50 litres	5 ml from Bag A	30 ppbC	5 ml from Bag A	159.5 ppbC

The bags were analyzed in descending order, then reanalyzed in the same order for duplicate analyses. Figures B-l and B-2 and Tables B-l and B-2 are examples of the 5 point calibrations of propane and toluene respectively.

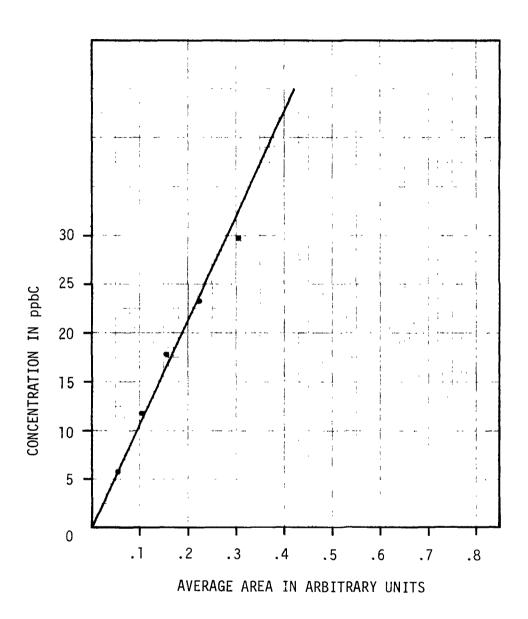
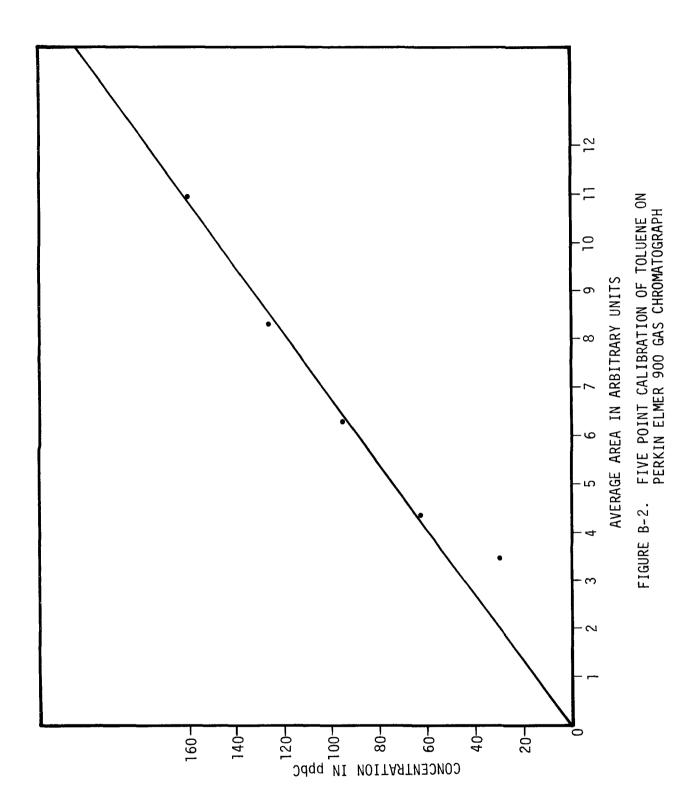


FIGURE B-1. FIVE POINT CALIBRATION OF PROPANE ON PERKIN ELMER 900 GAS CHROMATOGRAPH



## TABLE B-1. FIVE POINT CALIBRATION OF PROPANE

Bag A: 5.0 ml gas 50L of ultrapure air Bag l to 5: 50 litres of ultrapure air each

	Amount Injected		AREA					
Number	From Bag A	Run - 1	Run - 2	Average	Conc.			
1	1.0 ml	0.0642	0.0641	0.0641	6 ppbC			
2	2.0 ml	0.1068	0.1061	0.1064	i2 ppbC			
3	3.0 ml	0.1631	0.1652	0.1641	18 ppbC			
4	4.0 ml	0.2288	0.2297	0.2292	24 ppbC			
5	5.0 ml	0.3080	0.3082	0.3081	30 ppbC			

## TABLE B-2. FIVE POINT CALIBRATION OF TOLUENE

Bag A: 0.0500 ml liquid 50L of ultrapure air Bag l to 5: 50 litres of ultrapure air each

	Amount Injected		AREA				
Number	From Bag A	Run - 1	Run - 2	Average	Conc.		
1	1.0 ml	3.5380		3.5380	31.9 ppbC		
2	2.0 ml	4.4350	4.4036	4.4193	63.8 ppbC		
3	3.0 ml	6.6937	6.4601	6.5769	95.7 ppbC		
4	4.0 ml	8.5107	8.1254	8.3181	127.6 ppbC		
5	5.0 ml	10.9798	10.7468	10.8633	159.5 ppbC		

#### B.4 PEP-1 COMPUTER CALCULATIONS FOR P.E. 900

The PEP-1 computer was programmed in such a manner that it could detect, file, store, compute the concentration of selected compounds and print out pertinent data of samples that were analyzed on the P.E. 900.

In order to accurately compute the concentrations of the label compounds, the following information was input to the computer:

- reference peak(s)
- 2) relative retention times for all compounds
- 3) response factor for all compounds
- 4) response factor update

#### B.1.4.1 Reference Peak(s)

N-Butane and toluene were chosen as the reference peaks for the phenyl isocyanate and squalane columns respectively because both compounds occur abundantly in ambient air samples and no large peaks appear nearby, making identification easy. When writing the programs for the phenyl isocyanate and squalane columns, n-butane was assigned a reference number of one (1) and toluene was assigned a reference number of five (5). Along with the reference numbers, time spans for the elutions of the two compounds were provided.

#### B.4.2 Relative Retention Times

Relative retention times may be obtained two ways: manual computation or the PEP-1 system. The G. C. Laboratory employs both methods; the procedure and results of both are identical. As an example, the calculation of RRT of  $\rm C_2H_6$  is as follows:

X = Relative retention time of n-butane = 1.0.

T = The elution time in minutes of n-butane = 2.5 minutes.

 $X^{1}$  = The unknown relative retention time of  $C_{2}H_{6}$ .

 $T^1$  = The known elution time of the  $C_2H_6 = 0.87$  minutes.

$$\frac{X}{T} = \frac{X}{T}$$

$$\frac{1}{2.5} \times .87 = X^{1}$$

$$X^{1} = .348$$

It should be noted here that once the reference peak has been designated and its elution time standardized, the PEP-1 automatically assigned the relative retention times to all other compounds occurring in each sample. The same procedure was used for toluene.

#### B.4.3 Response Factor

To obtain response factors (RF) for hydrocarbons  $\rm C_2$  -  $\rm C_{10}$  a primary standard was made up and analyzed in the laboratory. The PEP-1 was programmed in such a manner that only the areas of the selected compounds were computed. After manually calculating the concentrations of the hydrocarbons and obtaining the various areas from the computer, it was possible to calculate the response factors for each compound.

The PEP-1 used the following formula:

Area x response factor x = 5 = concentration

To calculate the response factors of the hydrocarbons:

$$RF = \frac{Concentration}{Area \times 5}$$

Example: Toluene

$$RF = \frac{31.9}{2.0440 \times 5}$$

$$RF = 3.12$$

Response factors for all selected  ${\rm C_2}$  -  ${\rm C_{10}}$  hydrocarbons were determined in the above fashion.

After all of the calculations are done and the computer has been properly programmed, the PEP-1 uses; area x response factor x 5 to compute the concentrations of unknowns against the concentration of a standard.

#### B.4.4 Response Factor Update

Once the concentration of the standard was calculated and an acceptable range  $(\pm\ 3\%)$  for propane and  $\pm\ 3\%$  for toluene) for the area size of each compound had been established, the response factor update feature of the PEP-1 data system was used to correct the small changes that occur in the detector from day to day. The daily computed concentrations for the propane and toluene standards were compared to the calculated value for these standards; the relative change was fed to the computer and the response factor for each compound that is analyzed is multiplied by that factor. During this task order period, the detectors showed little or no change. The changes that occurred were random with a standard deviation of less than  $+\ 3\%$ .

#### B.5 CALCULATIONS USED TO DETERMINE STANDARD CONCENTRATIONS

The preparation of primary standards and the secondary standard was done at the same time. Only one secondary standard was needed because ambient air contains the propane and toluene that was used by the Gas Chromatography Laboratory on the phenyl isocyanate and squalane columns, respectively. The PEP-1 computer system uses the area x response factor x 5 method for computing the concentrations of compounds.

#### B.6 PRIMARY STANDARDS (Propane and Toluene) P.E. 900

The primary standards for the P.E. 900 were synthetically prepared in the laboratory by using the same reagents and equipment described in Section B.3 of this appendix. It should be noted here that only one dilution bag was used.

#### B.7 SECONDARY STANDARDS (Propane and Toluene) P.E. 900

Since the hydrocarbons to be analyzed occurred in laboratory air, secondary standards were prepared by simply storing a pressurized sample of ambient air in an 8 litre stainless steel tank, equipped with the appropriate valves. A sample of the secondary standard was then analyzed and the concentration of propane and toluene computed.

# APPENDIX C DATA FROM SUMMER AND FALL INTENSIVES

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#### C.O INTRODUCTION

Appendix C contains data obtained from samplings at RAMS stations. The results are in table form in order to provide an overview of the data from RAMS stations. The arithmetic mean  $(\overline{X})$  and standard deviation  $(S_X)$  were calculated for total hydrocarbon (THC), methane  $(CH_4)$ , carbon monoxide (CO) and acetylene  $(C_2H_2)$  for each table so that comparisons could be made between and among the RAMS stations. Not all compounds are listed in these tables. The magnetic tape of data delivered to the EPA contains the concentration for all compounds as well as sums for olefins, paraffins, etc. The following marks and abbreviations were used: "Trace" indicates less than 1 ppbC in the squalane analyses, "BDL" below detection limits, "NM" not measurable (because of interference, etc.). All concentrations in the following tables are in ppbC and the time of samples is Central Daylight Time (CDT) from April 25 to October 30 and Central Standard Time (CST) after October 30.

Figure C-1 shows the header for that tape and lists the various compounds which are reported.

```
001TOTAL SULFUR
002HYDROGEN SULFIDE
003SULFUR DIOXIDE
004METHYL MERCAPTAN
005NITROGEN OXIDES (NOX)
006SULFUR HEXAFLUORIDE
007TRICHLORO-FLUORD METHANE
008DICHLORO-DIFLUORO METHANE
009TOTAL ORGANICS
010METHANE
011CARBON MONCKIDE
012ETHYLENE
013ETHANE
014ACETYLENE
015PROPANE
016PROPYLENE
017 ISOBUTANE
018ISOBUTYLENE + BUTENE-1
019N-BUTANE
020T-2-BUTENE
021C-2-BUTENE
0223-M-1 BUTENE
023ISOPENTANE
024PENTENE-1
0252-M-1 BUTENE
026N-PENTANE
027T-2-PENTENE
028C-2-PENTENE
0292-M-2 BUTENE
0302,2-DM-BUTANE
031CYCLOPENTENE
0323-M-1-PENTENE + 4-M-1-PENTENE
0334-M-C-2-PENTENE
034CYCLOPENTANE
0352,3-DM-BUTANE + 4-M-T-2 PENTENE
0362-M-PENTANE
0372-M-1-PENTENE
0383-M-PENTANE + HEXENE-1 + 2-E-1-BUTENE
039T-3-HEXENS
0402-M-2-PENTENE + 3-M-CYCLOPENTANE
0413-M-C-2 PENTENE
042N-HEXANE
043T-2-HEXENE
044C-2-HEXENE
0453-M-T-2-PENTENE
046METHYLCYCLOPENTANE + 3,3-DM-1-PENTENE
047BENZENE
0482,4-DM-PEHTENE
0492,2,3-TM-BUTANE
0502,4-DM-1-PENTENE
0511-METHYLCYCLOPENTENE + 2-M-C-3-HEXENE
0522,4-DM-2-PENTENE + 3-E-1-PENTENE + 3-M-1-HEXENE
0532-M-T-3-HEXENE + 5-M-1-HEXENE
                                                       (continued)
054CYCLOHEXANE + 4-M-C-2-HEXENE
```

FIGURE C-1. LISTING OF ALL COMPOUNDS RECORDED ON MAGNETIC TAPES

```
0554-M-1-HEXENE + 4-M-T-2-HEXENE
0563-M-2-E-1-BUTENE + 5-M-T-2-HEXENE
057CYCLOHERSNE
0582-M-MEXANE + 5-M-C-2-HEXENE
0592.3-DM-PENTANS + 1,1-DM-CYCLOPENTANE
0603-M-HEX-HE
8611-C-3-DM-CYCLOPENTANE + 2-M-1-MEXANE
0621-T-3-001-CMCLOPENTANE + 1-MEPTENE + 2-E-1-PENTAME
0633-E-PENTONE + 3-M-T-2-HEXENS
0642,2,4-TH-PERTANE + T-3-HERTENE
8650-3-HEPTENE
0663-M-C-3-HEXEME + 2-M-2-PEXEME + 3-M-T-3-PEXEME
0673-E-2-FENTENE
068N-HEPTANE
0692,3-DIT-2-PENTEDE + C-2-HEPTENE
9701-C-2-DM-CYCLOPENTAME
071M-CYCLOHEXANE + 2,2-DM-HEXAME + 1,1,3-TM-CYCLOPENTANE
0724-M-CYCLOHEXENE
0732,5-DM-HEXANE
074ETHYLCYCLOPENTANE
0752.4-DM-HEXAME
0762,2,3-TM-PENTAME
0771-T-2-0-4-TM-CYCLOPENTANE
078TOLUENE
0792,3,4-TM-PEHTONE
0802,3,3-TM-PENTONE
0812,3-DM-HEXAME + 2-M-3-5-PENTAME
0822-M-HEPTANE
0934-11-HEPTANE
9843,4-DM-REXAME + 1-C-2-T-4-TM CYCLOPENTAME
0853-M-HEPTONE 4 3-M-3-E-PENTARE
0862,2,5-TM-HEMANE + 1-C-2-C-4-TM GYCLOPENTANE
0871-T-4-DM-CYCLOHEX9ME
0891-M-T-3 -E-CYC_OPENTAME
0892, 2, 4-TM HEXAME
090CYCLOHEPTANE + 1-M-1-E-CYCLOPENTANE
0911-T-2-DM-CYCLOHEXANE + 1-C-2-C-3-TM-CYCLOPENTANE
092N-OCTANE
0931-T-3-DM-CYCLOHEXANE
0942,4,4-TM-HENAME
0952,3,5-TM-HEXAME
0962.2-DM-HEPTANE
0972,4-PM-MEPTRNE + 2,2,3-TM-HEXANE
0982,2-DM-3-PENTANE + 2-M-4-E-HEWANE
0992,6-DM-FEPTAKE + 1-8-2-DM-CYCLOHEXANE
100N-PROPYLCYCLOPENTANE
101ETHYLCYCLOHEXANE
1022,5-DM-HEPTANS + 3,5-DM-HEPTANE
103ETHYLBENZENE
1043, 3-DM-HEPTANE
1052,3,3-TM-HEPTANE
106P-XYLENE
107M-XYLENG
                                                       (continued)
1084-M-OCTANE
```

FIGURE C-1 (continued)

```
1092-M-OCTANE
1103-E-HEPTANE
1113-M-00TANE
1120-XYLENE
1132.2, 4-TM-HEPTANE
1142,2,5-TM-HEPTANE + 2,2,5-TM-HEPTANE
1152,5,5ITM-HEPTANE + 2,4,4ITM-HEPTANE
116N-MONANC
117N-PROPYLIGENZENE
1182,2,3,3-TM-HEMANE
1191-M-2-E-BEHZENE
1201,3,5-TH- BENZENE
121TERT-PUTYLBENZENE
1221,2,4-TH-BENZENS
123SEC-BUTYLBENGENE + ISOBUTYLBENZENE
124N-DECANE
1251,2,3-TM-SENZENE + 1-M-4-ISOPROPYLBENZENE
126N-BUTYLBENZENE
127PARAFFINS
1280LEFINS
129AROMATICS
130TOTAL NON-METHANE HYDROCARSONS
```

FIGURE C-1 (continued)

# SUMMER INTENSIVE

	Mean	Standard Deviation
THC	3720 ppbC	494 ppbC
CH <sub>4</sub>	2016 ppbC	262 ppbC
co	1621 ppbC	634 ppbC
<sup>C</sup> 2 <sup>H</sup> 2	19.8 ppbC	11 ppbC

Date	6/23/76 7 <b>-</b> 9	6/23/76 9-11	6/23/76 11-13	6/23/76 13-15	7/13/76 9-11	7/15/76 7-9	7/15/76 9-11
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6					8.7	8.0	5.9
C <sub>2</sub> H <sub>4</sub>					996.5	967.5	18.8
C <sub>3</sub> H <sub>8</sub>					22.8	8.2	5.3
C2H2					8.7	8.6	5.0
IsoC <sub>4</sub> H <sub>10</sub>					12.6	4.1	4.5
N-C4H10					62.6	17.7	17.3
C <sub>3</sub> H <sub>6</sub>					2.4	3.0	BDL*
IsoC <sub>5</sub> H <sub>12</sub>					55.0	22.0	14.0
N-C <sub>5</sub> H <sub>12</sub>					14.0	9.0	7.0
Beckman 6800							
THC	2690	2600	3550	2540	3850	3640	2730
CH <sub>4</sub>	1960	1940	1900	1900	2060	2030	2090
CO	1520	1010	1050	1220	2210	1280	1460

<sup>\*</sup> Below detection limits

TABLE C-1 (continued)

Date			7/20/76 7-9		7/20/76 11-13		7/22/76 7-9
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	5.3	4.1	8.4	8.0	4.9	4.5	12.6
C <sub>2</sub> H <sub>4</sub>	1055.6	12.7	1345.9	30.0	1253.4	11.7	1553.2
<sup>C</sup> 3 <sup>H</sup> 8	6.0	3.4	10.4	9.7	5.2	4.1	16.7
C <sub>2</sub> H <sub>2</sub>	4.5	6.3	10.9	5.5	4.8	2.1	19.3
IsoC <sub>4</sub> H <sub>10</sub>	3.5	4.1	7.2	6.4	5.4	4.1	15.1
N-C <sub>4</sub> H <sub>10</sub> '	12.6	15.6	32.7	24.5	32.3	21.3	72.9
<sup>C</sup> 3 <sup>H</sup> 6	1.3	1.0	6.3	3.0	BDL*	BDL*	11.5
IsoC <sub>5</sub> H <sub>12</sub>	14.0	13.0	36.0	25.0	26.0	18.0	73.0
N-C <sub>5</sub> H <sub>12</sub>	7.0	4.6	18.0	8.0	8.0	6.0	28.5
Squalane							
<sup>N-C</sup> 6 <sup>H</sup> 14			12				27
<sup>N-C</sup> 7 <sup>H</sup> 16			5				9
Toluene			50				62
N-C <sub>8</sub> H <sub>18</sub>			15				5
E-C <sub>6</sub> H <sub>5</sub>			18				25
M-XYL			88				67
0-XYL			19				22
N-C <sub>9</sub> H <sub>20</sub>			40				2
N-P-C <sub>6</sub> H <sub>5</sub>			1				3
N-C <sub>10</sub> H <sub>22</sub>			189				5
Beckman 6800							
THC	3680	2470	3690	7950	3560	2270	3960
CH <sub>4</sub>	1960	1870	1920	1800	1780	1720	1920
CO	730	700	1320	926	712	610	2120

<sup>\*</sup> Below detection limits

TABLE C-1 (continued)

Date	7/22/76 9-11	7/22/76 13-15	7/27/76 7-9	7/27/76 9-11	7/27/76 11-13	7/27/76 13-15	7/29/76 7-9
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	9.4	5.8	14.5	10.4	12.3	8.7	11.3
C <sub>2</sub> H <sub>4</sub>	33.5	21.9	1240.8	30.1	1137.4	14.3	1044.0
C <sub>3</sub> H <sub>8</sub>	9.1	5.5	23.4	18.6	12.0	9.0	12.0
C <sub>2</sub> H <sub>2</sub>	10.8	4.2	17.3	11.9	11.2	8.5	12.4
IsoC <sub>4</sub> H <sub>10</sub>	6.6	6.2	13.8	12.4	5.0	4.4	4.4
N-C4H10	27.3	28.9	45.6	36.9	22.1	23.3	24.2
С <sub>3</sub> Н <sub>6</sub>	3.6	BDL*	34.0	43.0	66.0	25.0	15.0
IsoC <sub>5</sub> H <sub>12</sub>	28.0	24.0	43.0	30.0	22.0	21.0	26.0
N-C <sub>5</sub> H <sub>12</sub>	19.0	10.0	17.0	11.0	10.0	6.0	9.9
Squalane							
N-C6 <sup>H</sup> 14			10				13
N-C7H16			4				2
Toluene			48				117
N-C8H18			3				4
E-C <sub>6</sub> H <sub>5</sub>			13				30
M-XYL			40				73
0-XYL			11				24
N-C9 <sup>H</sup> 20			2				3
N-P-C <sub>6</sub> H <sub>5</sub>			1				2
N-C <sub>10</sub> H <sub>22</sub>							3
Beckman 6800							
THC	2370	2520	4010	2600	4860	2260	3630
CH <sub>4</sub>	1870	1800	1630	1480	2110	1710	2080
co	1060	525	1130	796	758	781	1190

<sup>\*</sup> Below detection limits

TABLE C-1 (continued)

Date	7/29/76 9-11	7/29/76 11-13	7/29/76 13-15	8/3/76 7-9	8/3/76 9-11	8/3/76 11-13	8/3/76 13-15
<u>Dura Pak</u>							
с <sub>2</sub> н <sub>6</sub>	11.7	12.2	14.1	9.8	6.6	5.0	6.0
с <sub>2</sub> н <sub>4</sub>	26.9	1402.6	41.3	1345.6	22.9	1151.5	20.7
С <sub>3</sub> Н <sub>8</sub>	16.5	19.2	20.8	17.7	11.4	7.2	8.6
C <sub>2</sub> H <sub>2</sub>	12.7	13.4	12.9	10.7	5.6	3.7	4.7
IsoC <sub>4</sub> H <sub>10</sub>	8.5	10.4	10.3	4.2	4.2	2.7	6.1
N-C <sub>4</sub> H <sub>10</sub>	43.3	63.6	53.0	22.7	15.1	10.6	22.2
С <sub>3</sub> Н <sub>6</sub>	11.5	14.0	8.0	6.8	3.1	1.5	1.3
IsoC <sub>5</sub> H <sub>12</sub>	38.0	50.0	42.0	23.0	10.0	9.2	19.0
N-C <sub>5</sub> H <sub>12</sub>	15.0	18.0	18.0	12.0	5.0	4.3	9.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>				8			
N-C <sub>7</sub> H <sub>16</sub>				2			
Toluene				81			
N-C <sub>8</sub> H <sub>18</sub>				4			
E-C <sub>6</sub> H <sub>5</sub>				9			
M-XYL				24			
0-XYL				8			
N-C <sub>9</sub> H <sub>20</sub>				1			
N-P-C <sub>6</sub> H <sub>5</sub>				BDL*			
N-C <sub>10</sub> H <sub>22</sub>				2			
Beckman 6800							
THC	3320	2960	6440	4060	2470	3180	2240
CH <sub>4</sub>	2000	1740	1710	2090	1920	1680	1720
<u>co</u>	1250	1070	1120	1040	754	585	645

<sup>\*</sup>Below detection limits

TABLE C-1 (continued)

Date	8/5/76 7-9	8/5/76 9-11	8/5/76 11-13	8/5/76 13-15	8/12/76 7-9	8/12/76 9-11	8/12/76 11-13
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	9.0	7.0	6.1	6.0	8.6	8.9	8.3
C <sub>2</sub> H <sub>4</sub>	1321.0	14.9	15.8	19.9	24.9	1864.0	43.8
С <sub>3</sub> Н <sub>8</sub>	11.2	7.3	5.5	4.9	19.4	18.4	7.7
С <sub>2</sub> Н <sub>2</sub>	14.4	8.3	6.5	6.2	29.7	10.3	7.4
IsoC <sub>4</sub> H <sub>10</sub>	8.2	5.0	6.7	5.1	14.0	14.0	5.5
N-C4H10	48.3	26.7	41.0	28.0	70.6	47.2	24.9
<sup>C</sup> 3 <sup>H</sup> 6	5.1	2.5	1.0	1.1	7.3	3.4	1.8
IsoC <sub>5</sub> H <sub>12</sub>	49.0	29.0	30.0	25.0	54.0	30.0	23.0
N-C <sub>5</sub> H <sub>12</sub>	21.0	11.0	13.0	11.0	28.0	13.0	10.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>	9				23		
N-C <sub>7</sub> H <sub>16</sub>	3				7		
Toluene	32				73		
N-C <sub>8</sub> H <sub>18</sub>	2				8		
E-C <sub>6</sub> H <sub>5</sub>	9				55		
M-XYL	32				199		
0-XYL	12				47		
N-C <sub>9</sub> H <sub>20</sub>	1				8		
N-P-C <sub>6</sub> H <sub>5</sub>	1				3		
N-C <sub>10</sub> H <sub>22</sub>	3				20		
Beckman 6800							
THC	3810	2350	2300	2200	3110	3790	2390
CH <sub>4</sub>	1820	1730	1720	1720	1820	1690	1740
<u>co</u>	1220	811	770	635	1810	850	630

TABLE C-1 (continued)

Date	8/17/76 7-9	8/17/76 9-11	8/17/76 11-13	8/19/76 7-9 Repeat Analysis	8/19/76 7 <b>-</b> 9	8/19/76 11-13
<u>Dura Pak</u>						
<sup>C</sup> 2 <sup>H</sup> 6	14.1	7.9	5.6	19.5	19.2	8.2
C <sub>2</sub> H <sub>4</sub>	158.1	83.1	79.4	108.3	108.1	30.8
C3H8	21.0	11.7	6.6	32.8	32.9	9.0
C <sub>2</sub> H <sub>2</sub>	21.6	13.5	6.6	19.6	18.5	7.5
IsoC <sub>4</sub> H <sub>10</sub>	10.7	7.2	3.0	9.1	9.0	4.8
N-C <sub>4</sub> H <sub>10</sub>	43.8	30.4	14.2	37.7	37.4	21.2
<sup>C</sup> 3 <sup>H</sup> 6	11.2	8.4	2.3	11.0	11.1	2.3
IsoC <sub>5</sub> H <sub>12</sub>	40.0	29.0	13.0	31.0	35.0	19.0
N-C <sub>5</sub> H <sub>12</sub>	19.0	13.0	6.0	25.0	21.0	10.0
Squalane						
N-C <sub>6</sub> H <sub>14</sub>	18				13	
N-C <sub>7</sub> H <sub>16</sub>	8				8	
Toluene	48				57	
N-C <sub>8</sub> H <sub>18</sub>	5				6	
E-C <sub>6</sub> H <sub>5</sub>	14				25	
M-XYL	39				42	
O-XYL	8				13	
N-C <sub>9</sub> H <sub>20</sub>	5				4	
N-P-C <sub>6</sub> H <sub>5</sub>	2				2	
N-C <sub>10</sub> H <sub>22</sub>	25				19	

TABLE C-1 (continued)

Date	8/24/76	8/24/76	8/24/76	8/24/76	8/26/76	8/26/76	8/26/76
	7-9	9-11	11-13	13-15	7-9	9-11	11-13
Dura Pak							
C <sub>2</sub> H <sub>6</sub>	30.2	10.0	9.0	7.0	18.6	12.3	10.3
C <sub>2</sub> H <sub>4</sub>	65.5	35.4	103.5	17.3	89.3	117.4	31.1
C <sub>3</sub> H <sub>8</sub>	70.8	13.4	9.6	9.3	23.9	14.4	12.9
C <sub>2</sub> H <sub>2</sub>	23.3	10.9	9.9	8.0	48.1	20.9	14.0
IsoC <sub>4</sub> H <sub>10</sub>	88.6	8.3	6.6	5.0	27.2	14.7	10.9
N-C <sub>4</sub> H <sub>10</sub>	240.2	35.2	41.3	29.8	144.3	98.9	61.8
C <sub>3</sub> H <sub>6</sub>	34.8	3.5	2.0	0.8	19.8	7.9	3.7
IsoC <sub>5</sub> H <sub>12</sub>	163.0	24.0	25.0	19.0	147.0	80.0	49.0
N-C <sub>5</sub> H <sub>12</sub>	98.0	14.0	14.0	10.0	61.0	33.0	21.0
$\begin{array}{c} \underline{\text{Squalane}} \\ N-C_6H_{14} \\ N-C_7H_{16} \\ \underline{\text{Toluene}} \\ N-C_8H_{18} \\ \underline{\text{E-C_6H_5}} \\ M-XYL \\ \underline{\text{O-XYL}} \\ N-C_9H_{20} \\ N-P-C_6H_5 \\ N-C_{10}H_{22} \end{array}$					26 8 59 6 23 66 23 5 4		
Beckman 6800 THC CH <sub>4</sub> CO	4570 2600 2020	2750 1960 780	2490 1880 860	2060 1740 460	3750 2310 3180	3200 1940 1520	2760 1960 1200

7751 L	1	irm,	Ched's
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are an agency of the second of the second of		1751	( Call Color
Date	8/26/76 13-11		
		• •	
oura Pak			
Calto	9.9	4 8	
C <sub>2</sub> H <sub>4</sub>	112.2	11.21	
C <sub>2</sub> H <sub>4</sub> - H <sub>8</sub> - C <sub>2</sub> H <sub>2</sub>	9 4	1.1	
- Spity	0.01	,' -4	
TEOC4H,	7,0	8.5	
N C4H10	40.0		
: 31 ó	3 1	,	
1500 <sub>5</sub> d <sub>12</sub>	43.6	29.0	
4 (5H <sub>12</sub>	7 ()	1 1	
Brokman 6800			
· HC	3290	****	
CH4	1840	1200	
(1)	1010	211	

TABLE C-2. RAMS STATION 102

SUMMER INTENSIVE									
	Mean Standard Deviation								
THC		2660 ppbC			306	ppbC			
CH <sub>4</sub>			1972 ppt	OC O	97	ppbC			
CO T		9 <b>3</b> 5 ppbC			214	ppbC			
Date	6/23/76 7-9	6/23/76 9 <b>-</b> 11	6/23/76 11 <b>-</b> 13	6/23/76 13-15					
Beckman 6800									
THC	3010	2820	2440	2370					
CH <sub>4</sub>	1970	2110	1900	1910					
co	1100	1140	750	750					

# SUMMER INTENSIVE

THC CH <sub>4</sub> CO C <sub>2</sub> H <sub>2</sub>			Mean 2327 ppb 1860 ppb 763 ppb 4 ppb	C C C	424 273 412	Deviation ppbC ppbC ppbC ppbC	
Date  Beckman 6800 THC CH <sub>4</sub> CO	6/28/76	6/28/76	6/28/76	6/28/76	6/30/7	76 6/30/76	6/30/76
	7-9	9-11	11-13	13-15	7-9	9-11	11-13
	3670	2530	2280	2350	2210	2070	2140
	3200	1920	1900	1850	1900	1880	1820
	1400	870	880	750	740	630	660

TABLE C-3 (continued)

Date	6/30/76 13 <b>-</b> 15	7/13/76 11-13	7/13/76 13-15	7/15/76 7-9	7/15/76 9-11	7/20/76 7-9	7/20/76 9-11	
Dura Pak				· -				
<sup>C</sup> 2 <sup>H</sup> 6		5.8	7.5	8.6	5.7	6.9	7.2	
C <sub>2</sub> H <sub>4</sub>		26.2	14.3	23.8	1128.0	26.8	224.2	
С <sub>3</sub> Н <sub>8</sub>		10.9	13.4	7.6	5.9	9.0	8.6	
$C_2H_2$		3.3	4.7	5.0	2.1	5.6	3.7	
IsoC <sub>4</sub> H <sub>10</sub>		3.4	7.3	4.2	2.0	6.5	3.7	
N-C <sub>4</sub> H <sub>10</sub>		12.7	31.8	18.2	8.5	<b>42.</b> 8	17.5	
C <sub>3</sub> H <sub>6</sub>		BDL*	Trace	BDL*	BDL*	3.7	BDL*	
IsoC <sub>5</sub> H <sub>12</sub>		9.0	22.0	14.0	6.5	27.0	12.0	
N-C <sub>5</sub> H <sub>12</sub>		1.0	9.0	4.5	1.4	8.0	4.8	
<u>Squalane</u>								
N-C <sub>6</sub> H <sub>14</sub>						NM**		
N-C <sub>7</sub> H <sub>16</sub>						2		
Toluene						49		
N-C <sub>8</sub> H <sub>18</sub>						2		
E-C <sub>6</sub> H <sub>5</sub>						31		
M-XYL						138		
0-XYL						27		
N-C <sub>9</sub> H <sub>20</sub>						2		
N-P-C <sub>6</sub> H <sub>5</sub>						BDL*		
N-C <sub>10</sub> H <sub>22</sub>						36		
Beckman 6800								
THC	2090	2090	2240	2470	3570	2730	2650	
CH <sub>4</sub>	1840	1910	1870	2030	1990	1880	1780	
co	370	610	800	1220	840	1170	990	
* Below detection limits (continued)								

<sup>\*\*</sup> Not measurable (because of interference etc.)

TABLE C-3 (continued)

Date	7/20/76 11 <b>-</b> 13	7/20/76 13-15	7/22/76 7-9	7/22/76 9-11	7/22/76 13-15	7/27/76 7-9	7/27/76 9-11
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	4.8	3.9	10.7	8.1	5.2	11.8	6.9
C <sub>2</sub> H <sub>4</sub>	38.9	13.6	69.0	32.8	26.5	20.1	19.9
C <sub>3</sub> H <sub>8</sub>	4.9	3.9	34.8	16.5	18.0	37.9	11.5
C <sub>2</sub> H <sub>2</sub>	1.5	1.5	9.1	5.5	2.2	6.6	4.3
IsoC <sub>4</sub> H <sub>10</sub>	3.7	3.1	23.1	4.5	2.6	15.6	6.7
N-C <sub>4</sub> H <sub>10</sub>	15.8	13.6	62.9	18.4	13.1	37.8	19.1
с <sub>3</sub> н <sub>6</sub>	BDL*	BDL*	5.9	2.8	BDL*	49.8	40.4
IsoC <sub>5</sub> H <sub>12</sub>	10.0	8.0	38.0	15.0	5.0	26.0	15.0
N-C <sub>5</sub> H <sub>12</sub>	4.8	3.0	15.0	6.0	4.0	13.0	6.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>			25			7 7	
N-C7 <sup>H</sup> 16			211			3	
Toluene			586			16	
N-C <sub>8</sub> H <sub>18</sub>			1008			2	
E-C <sub>6</sub> H <sub>5</sub>			1262			14	
M-XYL			1695			39	
O-XYL			3423			10	
N-C9 <sup>H</sup> 20			1426			2	
N-P-C <sub>6</sub> H <sub>5</sub>			6			2	
N-C <sub>10</sub> H <sub>22</sub>			24			BDL*	
Beckman 6800							
THC	2340	2160	1390	2310	1980	2500	2180
CH <sub>4</sub>	1760	1720	2000	1800	1760	1520	1470
сн	380	430	2510	1050	440	550	410

<sup>\*</sup> Below detection limits

TABLE C-3 (continued)

Date Dura Pak	7/27/76 11-13	7/27/76 13-15		11 Bags 7/29/76 1515- 1530	7/29/76 7-9	7/29/76 9-11	7/29/76 11-13
C <sub>2</sub> H <sub>6</sub>	7.1	6.5	55.4	55.3	9.7	10.1	12.0
C <sub>2</sub> H <sub>4</sub>	15.2	13.7	68.6	61.4	24.7	27.6	48.8
C <sub>3</sub> H <sub>8</sub>		8.9	19.2	18.9	19.3	18.0	19.8
C <sub>2</sub> H <sub>2</sub>		3.0		+ 7.0	7.6	7.0	8.5
IsoC <sub>4</sub> H <sub>10</sub>	2.8		11.2	9.5	5.6	5.2	8.0
N-C <sub>4</sub> H <sub>10</sub>	9.8	7.6	117.7	98.9	19.2	18.1	30.9
C <sub>3</sub> H <sub>6</sub>	35.6		8.9	1.8	12.2	12.7	10.0
IsoC <sub>5</sub> H <sub>12</sub>	7.8	4.0	88.0	64.0	19.0	18.0	26.0
N-C <sub>5</sub> H <sub>12</sub>	3.0	1.0	15.0	14.0	7.3	7.7	11.0
Squalane	3.0	1.0	13.0	14.0	7.3	/ • /	11.0
N-C <sub>6</sub> H <sub>14</sub>					5		
N-C <sub>7</sub> H <sub>16</sub>					1		
Toluene					27		
N-C <sub>8</sub> H <sub>18</sub>					2		
E-C <sub>6</sub> H <sub>5</sub>					10		
M-XYL					29		
O-XYL					8		
N-C <sub>9</sub> H <sub>20</sub>					2		
					2		
N-P-C <sub>6</sub> H <sub>5</sub> N-C <sub>10</sub> H <sub>22</sub>					2		
1022					_		
Beckman 6800							
THC	2690	2090	† 2650	2790	2190	2220	2420
CH <sub>4</sub>	1740	1680	1970	1760	1880	1920	1710
CO	410	390	2070	2230	810	840	811

+Values are not included in calculations

TABLE C-3 (continued)

Date	8/3/76 7-9	8/3/76 9-11	8/3/76 11-13	8/3/76 13-15	8/5/76 7-9	8/5/76 9-11	8/5/76 11-13
Dura Pak							
<sup>С</sup> 2 <sup>Н</sup> 6	11.2	4.7	4.0	5.9	6.8	8.4	6.0
C <sub>2</sub> H <sub>4</sub>	37.3	18.3	8.4	19.5	8.4	29.4	17.9
С <sub>3</sub> Н <sub>8</sub>	24.3	7.2	4.9	7.5	31.0	17.6	12.4
C <sub>2</sub> H <sub>2</sub>	4.1	2.9	2.2	2.7	3.3	5.8	3.4
IsoC <sub>4</sub> H <sub>10</sub>	3.5	2.5	2.3	4.8	15.8	8.0	6.6
	11.5	9.5	7.2	4.8	47.6	59.2	22.4
с <sub>3</sub> н <sub>6</sub>	5.1	11.8	2.2	4.0	2.0	2.0	1.0
IsoC <sub>5</sub> H <sub>12</sub>	10.0	8.0	3.0	6.0	40.0	32.0	17.0
N-C <sub>5</sub> H <sub>12</sub>	4.0	3.0	NM**	3.0	9.0	10.0	8.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>	3				<b>3</b> 8		
N-C <sub>7</sub> H <sub>16</sub>	1				1		
Toluene	8				16		
N-C8H18	BDL*				3		
E-C <sub>6</sub> H <sub>5</sub>	4				7		
M-XYL	13				29		
0-XYL	2				5		
N-C <sub>9</sub> H <sub>20</sub>	Trace				1		
N-P-C <sub>6</sub> H <sub>5</sub>	Trace				7		
N-C <sub>10</sub> H <sub>22</sub>	1				3		
Beckman 6800							
THC	2270	1940	1950	2100	2360	2360	2360
CH <sub>4</sub>	1920	1750	1830	1710	1800	1770	1720
CO	570	530	450	560	670	820	660

Below detection limits Not measurable (because of interference etc.)

TABLE C-3 (continued)

Date	8/5/76 13-15		
Dura Pak			
<sup>C</sup> 2 <sup>H</sup> 6	5.3		
C <sub>2</sub> H <sub>4</sub>	19.2		
C <sub>3</sub> H <sub>8</sub>	8.5		
C <sub>2</sub> H <sub>2</sub>	2.5		
IsoC <sub>4</sub> H <sub>10</sub>	4.9		
N-C4H10	26.2		
C <sub>3</sub> H <sub>6</sub>	1.0		
IsoC <sub>5</sub> H <sub>12</sub>	18.0		
N-C <sub>5</sub> H <sub>12</sub>	6.0		

TABLE C-4. RAMS STATION 108

SUMMER INTENSIVE										
Mean Standard Deviation										
THC		2		270 բ	opbC					
CH <sub>4</sub>		1	872 ppbC		98 p	орьС				
CO T		603 ppbC 388 p								
Date	6/23/76 7 <b>-</b> 9	6/23/76 11 <b>-</b> 13	6/23/76 13 <b>-</b> 15	6/28/76 9-11	6/28/76 11 <b>-</b> 13					
Beckman 6800	<u>.</u>									
THC	2870	2260	2210	2260	2210					
CH <sub>4</sub>	2130	1840	1840	1840	1840					
CO T	1480	680	520	680	560					
Date	6/30/76 7 <b>-</b> 9	6/30/76 9-11	6/30/76 11 <b>-</b> 13	6/30/76 13-15						
Beckman 6800	<u>)</u>									
THC	2190	2100	1980	1930						
CH <sub>4</sub>	1880	1820	1840	1820						
CO	410	740	180	180						

# TABLE C-5. RAMS STATION 113

## SUMMER INTENSIVE

		Mean		Standard Deviation	
THC	1874 ppbC			78 ppbC	
CH <sub>4</sub>		1912 ppbC		82 ppbC	
co		957 ppbC		167 ppbC	
Date	7/6/76 7-9	7/6/76 9-11	7/6/76 13-15		
Beckman 6800					
THC	1963	1816	1843		
CH <sub>4</sub>	1995	1912	1830		
co	1150	861	861		

# SUMMER INTENSIVE

THC CH <sub>4</sub> CO C <sub>2</sub> H <sub>2</sub>		Mean 2613 ppbC 1851 ppbC 599 ppbC 7.1 ppbC			Standard Deviation 912 ppbC 310 ppbC 302 ppbC 8.9 ppbC			
Date	6/23/76 9-11	6/23/76 11-13	6/23/76 13-15	7/15/76 9-11	7/15/76 11-13	7/15/76 13-15	7/20/76 9-11	
<u>Dura Pak</u>								
$C_2H_6$				4.8	4.6	2.4	5.7	
<sup>C</sup> 2 <sup>H</sup> 6 <sup>C</sup> 2 <sup>H</sup> 4 <sup>C</sup> 3 <sup>H</sup> 8				16.4	21.8	10.9	20.5	
C3H8				4.8	4.8	2.8	6.3	
C <sub>2</sub> H <sub>2</sub>				1.7	1.4	0.7	3.0+	
IsoC <sub>4</sub> H <sub>10</sub>				1.0	1.0	1.0	3.6	
N-C <sub>4</sub> H <sub>10</sub>				6.1	6.6	4.4	15.4	
C <sub>3</sub> H <sub>6</sub>				BDL*	BDL*	0.5	BDL*	
IsoC <sub>5</sub> H <sub>12</sub>				5.0	4.0	2.0	10.5	
N-C <sub>5</sub> H <sub>12</sub>				1.0	1.0	1.0	2.3	
Beckman 6800								
THC	2200	2750	2160	2450	2330	2350	2490	
CH <sub>4</sub>	1820	1840	1840	1960	1920	1650	1760	
co	520	560	540	550	400	330	460	

<sup>\*</sup> Below detection limits

TABLE C-6 (continued)

Date	7/20/76 11-13	7/20/76 13-15	7/22/76 9-11	7/22/76 11-13	7/22/76 13-15	7/27/76 9-11	7/27/76 11-13
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	5.0	3.9	8.2	7.5	5.5	7.5	6.6
C <sub>2</sub> H <sub>4</sub>	32.6	9.7	15.5	41.9	10.8	19.2	27.8
C <sub>3</sub> H <sub>8</sub>	4.9	3.7	8.9	7.6	4.9	14.7	10.1
C <sub>2</sub> H <sub>2</sub>	2.0	2.0	4.7	2.9	1.0	3.7	2.6
IsoC <sub>4</sub> H <sub>10</sub>	3.0	1.8	4.1	2.6	2.1	8.6	3.3
N-C4H10	12.8	9.8	14.2	12.4	11.2	19.0	9.9
С <sub>3</sub> Н <sub>6</sub>	BDL*	BDL*	0.7	BDL*	BDL*	31.0	51.7
IsoC <sub>5</sub> H <sub>12</sub>	10.0	6.8	14.0	10.0	9.0	11.0	4.0
N-C <sub>5</sub> H <sub>12</sub>	3.0	3.0	5.0	4.0	3.0	6.0	6.0
Beckman 6800							
THC	2650	2240	2080	2040	1950	2090	2470
CH <sub>4</sub>	1740	1610	1820	1800	1760	1390	1660
co ·	410	480	660	410	480	340	340
Date	7/27/76 13-15	7/29/77 11-13	8/3/76 11-13	8/3/76 13 <b>-</b> 15	8/5/76 9-11	8/5/76 11-13	8/5/76 13-15
<u>Dura Pak</u>							
<sup>С</sup> 2 <sup>Н</sup> 6	6.8	14.9	7.7	5.5	13.3	5.7	4.6
C <sub>2</sub> H <sub>4</sub>	39.9	29.6	113.3	17.0	52.9	102.5	10.5
C <sub>3</sub> H <sub>8</sub>	8.8	25.8	18.6	11.2	12.1	5.5	3.5
C <sub>2</sub> H <sub>2</sub>	2.0	5.6	2.8	2.4	11.5	5.3	2.7
IsoC <sub>4</sub> H <sub>10</sub>	2.5	14.3	16.2	13.1	7.4	3.8	1.8
N-C4H10	8.4	35.7	48.1	35.2	34.9	17.3	8.5
С <sub>3</sub> Н <sub>6</sub>	35.0	13.0	24.5	10.3	4.5	6.5	BDL*
IsoC <sub>5</sub> H <sub>12</sub>	2.0	25.0	27.0	35.0	31.0	13.0	6.0
N-C <sub>5</sub> H <sub>12</sub>	3.0	11.0	14.0	11.0	15.0	7.0	3.0
Beckman 6800							
THC		2300	2450	3870	2110	2740	2120
CH <sub>4</sub>		1680	1790	2230	1710	1720	1670
* Polow date		510	340	650	840	840	730

<sup>\*</sup> Below detection limits

TABLE C-6 (continued)

Date	8/10/76 9-11	8/10/76 11-13	8/10/76 13-15	8/12/76 7-9	8/12/76 9-11	8/12/76 13-15	8/17/76 7-9
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	5.5	51.9	4.2	37.6	6.7	5.4	6.2
C <sub>2</sub> H <sub>4</sub>	19.6	26.6	10.2	20.5	20.8	11.5	26.1
с <sub>3</sub> н <sub>8</sub>	6.1	26.9	7.5	19.8	7.4	5.7	15.1
C <sub>2</sub> H <sub>2</sub>	5.7	49.4	3.8	18.1	6.7	21.7	5.3
IsoC <sub>4</sub> H <sub>10</sub>	4.0	21.6	4.6	9.1	3.0	1.0	7.1
N-C <sub>4</sub> H <sub>10</sub>	18.7	56.3	19.8	29.0	12.5	4.7	27.4
с <sub>3</sub> н <sub>6</sub>	0.9	13.0	BDL*	6.0	1.5	BDL*	2.2
IsoC <sub>5</sub> H <sub>12</sub>	15.0	59.0	14.0	26.0	11.0	3.0	13.0
N-C <sub>5</sub> H <sub>12</sub>	6.0	53.0	6.0	39.0	5.0	2.0	9.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>				9			9
N-C <sub>7</sub> H <sub>16</sub>				14			3
Toluene				45			14
N-C <sub>8</sub> H <sub>18</sub>				3			1
E-C <sub>6</sub> H <sub>5</sub>				30			5
M-XYL				89			19
0-XYL				24			5
N-C <sub>9</sub> H <sub>20</sub>				3			7
N-P-C <sub>6</sub> H <sub>5</sub>				1			NM**
N-C <sub>10</sub> H <sub>22</sub>				8.			1
Beckman 6800							
THC				2370	2110	2090	
CH <sub>4</sub>				1880	1690	1670	
co				590	580	290	···········

Below detection limits
Not measurable (because of interference etc.)

TABLE C-6 (continued)

Date	8/17/76 9-11	8/17/76 11-13	8/18/76 13-15	8/19/76 7-9	8/19/76 9-11	8/19/76 11-13	8/19/76 13-15
Dura Pak							
С <sub>2</sub> Н <sub>6</sub>	6.5	5.2	4.8	15.2	13.1	6.1	5.8
C <sub>2</sub> H <sub>4</sub>	30.8	14.9	13.2	19.6	42.7	14.3	15.0
C <sub>3</sub> H <sub>8</sub>	13.0	8.4	6.6	55.5	53.5	6.1	8.8
C <sub>2</sub> H <sub>2</sub>	4.2	5.0	6.3	4.9	5.8	3.7	2.2
IsoC <sub>4</sub> H <sub>10</sub>	4.7	4.3	2.5	44.3	11.1	2.3	2.1
N-C <sub>4</sub> H <sub>10</sub>	15.8	20:1	11.3	131.1	46.5	8.1	8.0
C <sub>3</sub> H <sub>6</sub>	2.1	1.3	1.0	BDL*	BDL*	BDL*	BDL*
IsoC <sub>5</sub> H <sub>12</sub>	11.0	15.0	10.0	65.0	26.0	5.0	5.0
N-C <sub>5</sub> H <sub>12</sub>	5.0	6.0	3.0	42.0	11.0	3.0	3.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>				24			
N-7 <sup>H</sup> 16				17			
Toluene				21			
N-C <sub>8</sub> H <sub>18</sub>				9			
E-C <sub>6</sub> H <sub>5</sub>				12			
M-XYL				30			
0-XYL				10			
N-C <sub>9</sub> H <sub>20</sub>				2			
N-P-C <sub>6</sub> H <sub>5</sub>				1			
N-C <sub>10</sub> H <sub>22</sub>				5			

<sup>\*</sup> Below detection limits

TABLE C-6 (continued)

Date	8/26/76 7-9	8/26/76 9-11	8/31/76 7-9	8/31/76 9-11	9/2/76 7-9	9/2/76 9-11	9/2/76 13-15
Dura Pak							
<sup>С</sup> 2 <sup>Н</sup> 6	15.8	14.2	57.3	13.7	15.4	7.6	4.6
C <sub>2</sub> H <sub>4</sub>	35.4	37.0	24.9	55.0	18.2	25.3	31.6
С <sub>3</sub> Н <sub>8</sub>	19.5	15.9	40.1	35.6	51.1	24.4	5.1
C <sub>2</sub> H <sub>2</sub>	10.1	15.3	21.5	16.3	3.2	2.0	2.3
IsoC <sub>4</sub> H <sub>10</sub>	8.3	7.8	38.0	23.8	73.9	20.1	2.0
N-C4H10	30.0	37.0	173.3	61.1	210.2	77.6	7.5
С <sub>3</sub> Н <sub>6</sub>	3.9	4.7	10.7	6.1	19.7	73.9	0.7
IsoC <sub>5</sub> H <sub>12</sub>	25.0	32.0	193.0	52.0	95.0	45.0	8.0
N-C <sub>5</sub> H <sub>12</sub>	11.0	14.0	98.0	25.0	69.0	19.0	2.0
<u>Squalane</u>							
N-C6H14	11		15		36		
N-C7H16	2		5		15		
Toluene	23		56		21		
N-C <sub>8</sub> H <sub>18</sub>	2		4		2		
E-C <sub>6</sub> H <sub>5</sub>	11		52		17		
M-XYL	32		131		55		
0-XYL	9		37		14		
N-C <sub>9</sub> H <sub>20</sub>	1		2		2		
N-P-C <sub>6</sub> H <sub>5</sub>	1		11		2		
N-C <sub>10</sub> H <sub>22</sub>	8		7		1		
Beckman 6800							
THC	2850	2910	6140	3000	2800	2020	1780
CH <sub>4</sub>	2290	2200	3100	2220	1690	1720	1690
co	830	1200	1330	1540	450	350	410

# SUMMER INTENSIVE

			Mean		Standar	d Deviati	on
THC			2427 ppb	С	908	ppbC	
CH₄			1788 ppb	С	147	ppbC	
co			498 ppb	С	213	ppbC	
<sup>C</sup> 2 <sup>H</sup> 2			3 ppb	С	2	ppbC	
Date	6/28/76 9-11	6/28/76 11-13	6/28/76 13-15	6/30/76 9-11	6/30/76 11 <b>-</b> 13	6/30/76 13-15	
Beckman 6800							
THC	2660	2490	2550	2310	2330	2120	
CH <sub>4</sub>	1880	1880	1900	1840	1800	1840	
co	550	650	430	630	260	260	
Date	7/13/76 13-15	7/15/76 9-11	7/15/76 11 <b>-1</b> 3	7/15/76 13 <b>-</b> 15	7/20/76 9-11	7/20/76 11 <b>-</b> 13	7/20/76 13-15
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	7.6	4.7	3.4	2.5	5.7	4.8	4.2
C <sub>2</sub> H <sub>4</sub>	20.6	33.9	28.5	36.2	34.0	52.8	31.2
C <sub>3</sub> H <sub>8</sub>	14.8	4.6	3.1	2.1	6.6	5.0	4.3
C <sub>2</sub> H <sub>2</sub>	3.0	1.7	2.2	0.9	2.5	2.3	2.4
IsoC <sub>4</sub> H <sub>10</sub>	4.9	1.5	1.4	0.9	3.1	2.0	2.3
N-C <sub>4</sub> H <sub>10</sub>	10.9	5.6	5.3	4.2	12.9	8.4	9.6
С <sub>3</sub> Н <sub>6</sub>	2.0	BDL*	BDL*	BDL*	BDL*	BDL*	BDL*
IsoC <sub>5</sub> H <sub>12</sub>	8.0	3.0	2.0	3.0	5.0	7.0	8.0
N-C <sub>5</sub> H <sub>12</sub>	2.9	1.0	BDL*	BDL*	3.0	2.0	2.0
Beckman 6800							
THC	2160	2520	2260	2140	2220	2170	2090
CH <sub>4</sub>	1870	1930	1870	1830	1760	1720	1720
CO	860	490	330	610	690	410	410

<sup>\*</sup> Below detection limits

TABLE C-7 (continued)

Date	7/22/76	7/22/76	7/27/76	7/27/76		7/29/76	7/29/76
Duna Dak	9-11	13-15	9-11	11-13	13-15	9-11	11-13
Dura Pak	9.9	6.0	6.6	6.8	5.7	12.0	11 0
<sup>C</sup> 2 <sup>H</sup> 6						13.9	11.0
<sup>C</sup> 2 <sup>H</sup> 4	38.8	24.2	33.2	39.9	17.6	19.8	25.4
C3H8	14.6	5.2	9.0	8.8	5.1	22.8	16.2
C <sub>2</sub> H <sub>2</sub>	6.3	1.3	3.1	2.0	1.5	6.9	3.9
IsoC <sub>4</sub> H <sub>10</sub>	7.0	2.1	4.8	2.5	1.2	9.4	7.8
N-C <sub>4</sub> H <sub>10</sub>	24.1	10.6	12.1	8.4	4.8	27.9	22.5
<sup>C</sup> 3 <sup>H</sup> 6	BDL*	BDL*	44.9	35.0	19.3	14.2	11.0
IsoC <sub>5</sub> H <sub>12</sub>	19.0	9.0	6.0	2.0	2.0	26.0	15.0
N-C <sub>5</sub> H <sub>12</sub>	9.0	2.0	2.0	4.0	BDL*	10.0	7.0
Beckman 6800							
THC	2210	2010	3830	2460	2080	2030	2860
CH <sub>4</sub>	1920	1770	1350	1610	1580	2060	1650
co	160	380	280	330	330	950	450
Date	7/29/76 13-15	8/3/76 9-11	8/3/76 11-13	8/3/76 13-15	8/5/76 9-11		
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	11.3	3.2	8.0	2.9	7.4		
C <sub>2</sub> H <sub>4</sub>	12.9	21.6	36.0	17.0	27.8		
C <sub>3</sub> H <sub>8</sub>	16.9	3.1	5.2	1.9	10.1		
C <sub>2</sub> H <sub>2</sub>	4.5	2.6	10.4	1.9	6.8		
IsoC <sub>4</sub> H <sub>10</sub>	8.0	2.1	6.0	2.0	4.0		
N-C <sub>4</sub> H <sub>10</sub>	23.7	10.9	37.1	7.3	22.6		
с <sub>3</sub> н <sub>6</sub>	8.1	14.4	2890.9	6.7	19.2		
IsoC <sub>5</sub> H <sub>12</sub>	14.0	10.0	28.0	3.0	18.0		
N-C <sub>5</sub> H <sub>12</sub>	7.0	3.0	6.0	2.0	9.0		
Beckman 6800							
THC	1060	1980	6200	2000	2230		
$CH_4$	1670	1690	1980	1800	1790		
co	440	700	350	560	940		

<sup>\*</sup> Below detection limits

SUMMER INTENSIVE								
		Mean			Standard Deviation			
THC			2791 p	pbC	1311 ppbC			
CH₄			1770 p	pbC		133 ppb0		
co			479 p	pbC		301 ppb0	•	
С <sub>2</sub> Н <sub>2</sub>			3.4 p	pbC		2.3 ppb0	,	
Date	7/13/76 13-15	7/15/76 7-9	7/15/76 13-15	7/20/76 7-9	7/22/76 7-9	7/22/76 13-15	7/27/76 7-9	
Dura Pak								
с <sub>2</sub> н <sub>6</sub>	6.2	5.0	2.7	4.8	6.6	4.6	6.3	
C <sub>2</sub> H <sub>4</sub>	41.4	15.0	7.9	24.8	9.3	13.5	18.2	
С <sub>3</sub> Н <sub>8</sub>	27.6	6.8	2.0	4.6	8.3	3.9	7.7	
C <sub>2</sub> H <sub>2</sub>	8.5	2.7	0.6	1.5	1.1	0.2	5.6	
IsoC <sub>4</sub> H <sub>10</sub>	2.4	1.2	0.5	1.1	2.4	1.2	3.3	
N-C <sub>4</sub> H <sub>10</sub>	6.6	7.9	2.4	6.4	9.7	3.9	14.6	
С <sub>3</sub> Н <sub>6</sub>	BDL	BDL	BDL	BDL	BDL	0.7	12.7	
IsoC <sub>5</sub> H <sub>12</sub>	3.0	5.0	1.0	2.0	4.0	2.0	8.0	
N-C <sub>5</sub> H <sub>12</sub>	1.0	2.0			14.0	1.0	2.0	
<u>Squalane</u>								
N-C <sub>6</sub> H <sub>14</sub>				NM			2	
N-C7 <sup>H</sup> 16				8			1	
Toluene				26			4	
N-C <sub>8</sub> H <sub>18</sub>				23			1	
E-C <sub>6</sub> H <sub>5</sub>				19			5	
M-XYL				84			18	
0-XYL				21			6	
N-C <sub>9</sub> H <sub>20</sub>				37			Trace	
N-P-C <sub>6</sub> H <sub>5</sub>				7			Trace	
N-C <sub>10</sub> H <sub>22</sub>				126			1	
Beckman 6800								
THC	3850	2330	2280	2340		2670	2080	
CH <sub>4</sub>	1870	1900	1870	1820		1740	1480	
co	1380	580	210	356		191	380	

TABLE C-8 (continued)

Date	7/27/76 13-15	7/29/76 7-9	7/29/76 13-15	8/03/76 7-9	8/03/76 13 <b>-</b> 15		8/05/76 13-15
Dura Pak							
<sup>С</sup> 2 <sup>Н</sup> 6	7.8	6.4	8.4	6.2	3.8	5.3	4.2
<sup>C</sup> 2 <sup>H</sup> 4	4.3	25.2	22.7	32.5	11.2	27.2	12.6
с <sub>3</sub> н <sub>8</sub>	5.5	7.6	10.2	22.7	7.0	4.7	3.0
С <sub>2</sub> Н <sub>2</sub>	4.1	4.7	5.0	5.2	2.8	1.9	1.3
Iso€ <sub>4</sub> H <sub>10</sub>	1.8	2.8	3.1	6.5	3.5	1.8	0.8
N-C <sub>4</sub> H <sub>10</sub>	7.8	15.3	15.8	20.0	13.5	5.6	3.2
<sup>C</sup> 3 <sup>H</sup> 6	10.0	24.4	19.4	92.0	BDL*	BDL*	BDL*
IsoC <sub>5</sub> H <sub>12</sub>	6.0	11.0	11.0	12.0	9.0	3.0	2.0
N-C5 <sup>H</sup> 12	1.0	2.0	3.0	9.0	4.0	5.2	1.2
Squalane							
N-C <sub>6</sub> H <sub>14</sub>		2		5	1		
N-C7H16		1		2		Trace	
Toluene		13		6		3	
N-C <sub>8</sub> H <sub>18</sub>		2		2		7	
E-C <sub>6</sub> H <sub>5</sub>		10		3		3	
M-XYL		43		11		17	
0-XYL		9		4		2	
N-C <sub>9</sub> H <sub>20</sub>		1		ī		Trace	
N-P-C <sub>6</sub> H <sub>5</sub>		1		Trace		BDL*	
N-C <sub>10</sub> H <sub>22</sub>		21		2		1	
Beckman 6800							
THC	6070	2030	2610	2280	2130	2060	2100
CH <sub>4</sub>	1990	1780	1610	1810	1750	1720	1670
CO	460	490	430	530	590	410	230

<sup>\*</sup> Below detection limits

TABLE C-9. RAMS STATION 121

SUMMER INTENSIVE								
			Mean		Standar	d Deviation		
THC			1960 ppb0	2	311	ppbC		
CH <sub>4</sub>			1872 ppb0	C	43 ppbC			
co		364 ppbC			183 ppbC			
Date	7/6/76 9-11	7/6/76 11-13	7/6/76 13-15	7/8/76 9-11	7/8/76 13-15			
Beckman 6800								
THC	1790	1880	2500	1910	1720			
CH <sub>4</sub>	1870	1930	1830	1900	1830			
co	660	370	717	350	270			

	<u>SU</u>	MMER INTE	NSIVE			
		Mean		Standa	rd Deviat	ion
THC		2939 ppb	С	1771	ppbC	
CH <sub>4</sub>		1864 ppb	С	212	ppbC	
CO		607 ppb	C	240	ppbC	
С <sub>2</sub> Н <sub>2</sub>		5 ppb	C	3	ppbC	
Date	8/10/76 13-15	8/12/76 7-9	8/12/76 13-15	8/17/76 7 <b>-</b> 9	8/17/76 13-15	8/19/76 13-15
<u>Dura Pak</u>						
<sup>C</sup> 2 <sup>H</sup> 6	4.5	12.3	7.4	4.2	2.9	4.9
<sup>C</sup> 2 <sup>H</sup> 4	6.4	15.4	16.1	6.6	5.3	6.5
C3H8	6.0	26.1	6.8	8.5	3.4	4.3
С <sub>2</sub> Н <sub>2</sub>	5.0	8.4	4.4	4.1	2.0	2.4
IsoC <sub>4</sub> H <sub>10</sub>	3.3	20.4	2.3	2.3	0.9	1.3
N-C <sub>4</sub> H <sub>10</sub>	15.8	47.2	7.8	9.1	3.4	4.2
<sup>C</sup> 3 <sup>H</sup> 6	BDL*	4.9	1.0	1.1	BDL*	BDL*
IsoC <sub>5</sub> H <sub>12</sub>	12.0	36.0	7.0	6.0	1.9	2.0
N-C5 <sup>H</sup> 12	4.0	18.0	3.0	2.0	8.0	2.0
<u>Squalane</u>						
N-C <sub>6</sub> H <sub>14</sub>				2		
N-C7 <sup>H</sup> 16				3		
Toluene				12		
N-C <sub>8</sub> H <sub>18</sub>				Trace		
E-C <sub>6</sub> H <sub>5</sub>				8		
M-XYL				28		
0-XYL				5		
N-C <sub>9</sub> H <sub>20</sub>				Trace		
N-P-C <sub>6</sub> H <sub>5</sub>				1		
N-C <sub>10</sub> H <sub>22</sub>				2		
Beckman 6800						
THC	2070	2580	1950			
CH <sub>4</sub>	1810	1840	1670			
CO	450	650	300			

\* Below detection limits

TABLE C-10 (continued)

Date	8/24/76 7-9	8/24/76 13-15	8/26/76 7-9	8/26/76 13-15	8/31/76 7-9	8/31/76 13-15	9/2/76 7-9
<u>Dura Pak</u>							
<sup>C</sup> 2 <sup>H</sup> 6	7.5	5.8	9.4	7.7	18.1	8.4	0.7
C <sub>2</sub> H <sub>4</sub>	18.9	8.1	43.3	25.6	16.8	5.7	3.1
C <sub>3</sub> H <sub>8</sub>	10.1	4.6	10.0	7.4	29.1	28.2	4.4
C <sub>2</sub> H <sub>2</sub>	3.9	3.0	10.0	3.9	10.6	5.3	1.3
IsoC <sub>4</sub> H <sub>10</sub>	2.0	1.7	7.1	2.1	12.4	23.4	1.0
N-C4H10	7.8	4.1	31.2	6.7	43.1	76.3	4.2
<sup>C</sup> 3 <sup>H</sup> 6	0.7	BDL*	1.3	0.4	5.3	1.1	0.1
IsoC <sub>5</sub> H <sub>12</sub>	5.0	3.0	22.0	5.0	37.0	46.0	2.0
N-C <sub>5</sub> H <sub>12</sub>	3.0	2.0	13.0	2.0	17.0	27.0	7.0
Squalane							
N-C6 <sup>H</sup> 14			6		10		5
N-C7H16			2		2		5
Toluene			18		24		4
N-C <sub>8</sub> H <sub>18</sub>			1		2		1
E-C <sub>6</sub> H <sub>5</sub>			16		12		5
M-XYL			48		31		16
0-XYL			14		10		4
N-C <sub>9</sub> H <sub>20</sub>			1		1		1
N-P-C <sub>6</sub> H <sub>5</sub>			2		1		1
N-C <sub>10</sub> H <sub>22</sub>			8		5		2
Beckman 6800							
THC	2270	2080	2520	2210	2750	2160	1600
CH <sub>4</sub>	1890	1690	1900	1800	2390	1800	1650
CO	460	400	760	490	970	620	310

<sup>\*</sup> Below detection limits

TABLE C-10 (continued)

Date	9/2/76 13-15
Dura Pak	
<sup>C</sup> 2 <sup>H</sup> 6	4.7
С <sub>2</sub> Н <sub>4</sub>	5.1
C3H8	5.2
C <sub>2</sub> H <sub>2</sub>	1.9
IsoC <sub>4</sub> H <sub>10</sub>	1.9
N-C <sub>4</sub> H <sub>10</sub>	5.8
C <sub>3</sub> H <sub>6</sub>	
IsoC <sub>5</sub> H <sub>12</sub>	3.0
N-C <sub>5</sub> H <sub>12</sub>	2.0

		<u>SU</u>	MMER INTE	NSIVE			
			Mean		Standa	rd Deviat	ion
THC			2264 ppb	С	382	ppbC	
$CH_{\mathbf{\Delta}}$			1931 ppb	С	402	ppbC	
coʻ			579 ppb	С	214	ppbC	
с <sub>2</sub> н <sub>2</sub>			3 ppb	С	2	ppbC	
Date	8/12/76 7-9	8/12/76 13-15	8/17/76 11-13	8/24/76 7-9	8/26/76 7-9	8/26/76 13-15	8/31/76 7-9
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	4.8	7.6	8.4	10.4	6.0	7.2	1.8
C <sub>2</sub> H <sub>4</sub>	255.8	40.0	112.9	54.1	151.7	113.5	38.5
C <sub>3</sub> H <sub>8</sub>	5.5	6.9	15.6	14.8	5.9	5.9	5.7
C <sub>2</sub> H <sub>2</sub>	2.5	2.6	2.6	7.1	2.6	1.7	3.5
IsoC <sub>4</sub> H <sub>10</sub>		1.8	3.1	5.2	1.1	1.7	1.5
N-C <sub>4</sub> H <sub>10</sub>	4.4	4.3	11.7	30.8	4.5	6.2	8.4
С <sub>3</sub> Н <sub>6</sub>				1.9	0.9	0.5	1.6
IsoC <sub>5</sub> H <sub>12</sub>	3.0	3.0	4.0	14.0	3.0	5.0	5.0
N-C <sub>5</sub> H <sub>12</sub>	2.0	2.0	3.0	6.0	3.0	4.0	3.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>							
N-C <sub>7</sub> H <sub>16</sub>							
Toluene							
N-C <sub>8</sub> H <sub>18</sub>							
E-C <sub>6</sub> H <sub>5</sub>							
M-XYL							
0-XYL							
N-C9 <sup>H</sup> 20							
N-P-C <sub>6</sub> H <sub>5</sub>							
N-C <sub>10</sub> H <sub>22</sub>							
Beckman 6800							
ТНС	2380	1950		2270	2630	2210	
сн <sup>4</sup>	1720	1670		1890	1900	1930	
CO	290	300		460	460	300	

TABLE C-11 (continued)

Date	8/31/76 13-15	9/2/76 7-9	9/2/76 13-15
Dura Pak			
<sup>C</sup> 2 <sup>H</sup> 6	6.3	6.9	5.4
C <sub>2</sub> H <sub>4</sub>	278.6	111.0	180.8
C <sub>3</sub> H <sub>8</sub>	4.3	13.1	7.7
C <sub>2</sub> H <sub>2</sub>	2.9	2.4	1.6
IsoC <sub>4</sub> H <sub>10</sub>	2.0	3.2	2.4
N-C4H10	6.9	10.5	7.1
C <sub>3</sub> H <sub>6</sub>	0.7	0.9	0.5
IsoC <sub>5</sub> H <sub>12</sub>	5.0	5.0	5.0
N-C <sub>5</sub> H <sub>12</sub>	3.0	3.0	3.0
Squalane			
N-C <sub>6</sub> H <sub>14</sub>			
N-C <sub>7</sub> H <sub>16</sub>			
Toluene			
N-C <sub>8</sub> H <sub>18</sub>			
E-C <sub>6</sub> H <sub>5</sub>			
M-XYL			
0-XYL			
N-C <sub>9</sub> H <sub>20</sub>			
N-P-C <sub>6</sub> H <sub>5</sub>			
N-C <sub>10</sub> H <sub>22</sub>			
10 22 Beckman 6800			
THC	2900	1910	1850
	2140	1750	1690
CH <sub>4</sub>	620	280	220
<u>CO</u>	UZU	200	

	TABLE C-12. RAMS STATION	N 101
	FALL INTENSIVE	
	Mean	Standard Deviation
THC	2906 ppbC	789 ppbC
CH	1734 ppbC	240 ppbC

		<del></del>	Mea	an	Standard	Deviatio	n
THC			2906 p	opbC	789 p	pbC	
CH <sub>4</sub>			1734 p	opbC	240 p	pbC	
co ·			1815 p	opbC	1322 p	pbC	
с <sub>2</sub> н <sub>2</sub>			33 p	pbC	26 p	pbC	
Date	11/8/76 13-15	11/10/76 7-9	11/10/ 9-11	76 11/10/76 11-13	11/10/76 13-15	11/12/7 9-11	6 11/12/76 13:30-15
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	18.0	11.9	10.4	7.9	6.4	9.2	7.6
$C_2H_4$	46.4	24.3	20.4	10.7	7.7	21.1	16.5
С <sub>3</sub> H <sub>8</sub>	11.4	6.8	8.8	5.2	4.9	9.0	6.6
C <sub>2</sub> H <sub>2</sub>	40.3	25.8	13.6	5.3	5.7	18.2	10.2
IsoC <sub>4</sub> H <sub>10</sub>	11.8	6.7	5.0	3.1	3.3	4.9	3.1
N-C4H10	47.5	29.6	22.8	15.4	16.0	20.9	12.8
с <sub>3</sub> н <sub>6</sub>	12.4	7.0	6.0	3.4	2.1	4.1	1.5
IsoC <sub>5</sub> H <sub>12</sub>	44.0	28.0	18.0	9.0	10.0	14.0	9.0
N-C <sub>5</sub> H <sub>12</sub>	22.0	13.0	9.0	6.0	5.0	6.0	5.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>		NM				7	
N-C <sub>7</sub> H <sub>16</sub>		3				4	
Toluene		38				42	
N-C8H18		14				3	
E-C <sub>6</sub> H <sub>5</sub>		11				15	
M-XYL		27				37	
0-XYL		9				10	
N-C <sub>9</sub> H <sub>20</sub>		43				2	
N-P-C <sub>6</sub> H <sub>5</sub>		2				1	
N-C <sub>10</sub> H <sub>22</sub>		370				4	
Beckman 6800							
THC		2690					1990
CH <sub>4</sub>		2040					1470
CO		2115				740	250

TABLE C-12 (continued)

		IADL	L U-12 (C	on cinueu)			
Date	11/16/76 7-9	11/16/76 9-11	11/16/76 11-13	11/16/76 13-15	11/18/76 7-9	11/18/76 9-11	11/18/76 11-13
<u>Dura Pak</u>							
<sup>C</sup> 2 <sup>H</sup> 6	40.0	33.5	30.0	10.7	75.6	66.4	23.8
<b>c</b> <sub>2</sub> H <sub>4</sub>	87.9	185.5	120.3	19.7	57.4	40.5	28.7
C3H8	37.0	43.5	31.3	14.8	75.5	73.4	27.0
C <sub>2</sub> H <sub>2</sub>	71.2	62.8	87.2	14.2	60.1	42.0	16.7
IsoC <sub>4</sub> H <sub>10</sub>	23.0	33.4	47.6	8.4	32.0	27.8	21.0
N-C <sub>4</sub> H <sub>10</sub>	101.3	160.8	187.2	30.8	134.2	103.2	95.0
<sup>C</sup> 3 <sup>H</sup> 6	26.0	24.5	57.5	5.8	22.2	15.1	8.8
IsoC <sub>5</sub> H <sub>12</sub>	92.0	108.0	161.0	23.0	99.0	71.0	61.0
N-C <sub>5</sub> H <sub>12</sub>	38.0	46.0	76.0	11.0	54.0	37.0	28.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>	29				31		
N-C7 <sup>H</sup> 16	13				5		
Toluene	87				49		
N-C8H18	10				15		
E-C <sub>6</sub> H <sub>5</sub>	31				16		
M-XYL	<b>6</b> 8				40		
0-XYL	28				17		
N-C <sub>9</sub> H <sub>20</sub>	3				35		
N-P-C <sub>6</sub> H <sub>5</sub>	4				3		
N-C <sub>10</sub> H <sub>22</sub>	15				10		
Beckman 6800							
THC	3900	4020		2310	3860	3300	2540
CH <sub>4</sub>	2120	1980		1490	1970	1830	1470
CO	3660	3810		710	3100	2250	900

TABLE C-12 (continued)

Date	11/18/76 13-15			
Dura Pak				
<sup>C</sup> 2 <sup>H</sup> 6	46.2			
C <sub>2</sub> H <sub>4</sub>	20.2			
C <sub>3</sub> H <sub>8</sub>	53.3			
C <sub>2</sub> H <sub>2</sub>	17.2			,
IsoC <sub>4</sub> H <sub>10</sub>	22.5			
N-C4 <sup>H</sup> 10	99.4			
<sup>C</sup> 3 <sup>H</sup> 6	9.5			
IsoC <sub>5</sub> H <sub>12</sub>	58.0			
N-C <sub>5</sub> H <sub>12</sub>	28.0			
Beckman 6800				
THC	2670			
CH <sub>4</sub>	1610			
CO	900		 	

		F	ALL INTEN	SIVE			
			Mean		Standard [	Deviation	
THC			2676 ppb		817 ppl		
CH <sub>₄</sub>			1739 ppb		367 ppi		
co <sup>4</sup>			1073 ppb		844 ppl		
C <sub>2</sub> H <sub>2</sub>			15 ppb		14 ppl		
Date	11/8/76 7-9	11/8/76 11-13	11/8/76 13-15	11/10/76 7-9	11/10/76 9-11	11/10/76 11-13	11/10/76 13-15
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	19.1	5.0	45.2	9.7	6.4	9.0	7.8
C <sub>2</sub> H <sub>4</sub>	47.2	15.4	9.4	31.0	20.1	15.7	16.0
C <sub>3</sub> H <sub>8</sub>	13.5	4.8	65.7	7.6	7.8	6.0	6.6
C <sub>2</sub> H <sub>2</sub>	26.0	3.9	7.5	17.7	8.2	5.0	6.6
IsoC <sub>4</sub> H <sub>10</sub>	6.8	2.8	17.7	4.5	3.2	4.9	2.8
N-C4 <sup>H</sup> 10	31.5	9.1	63.8	22.2	12.0	43.0	18.4
C <sub>3</sub> H <sub>6</sub>	7.7	2.5	2.4	6.1	3.0	4.3	3.9
IsoC <sub>5</sub> H <sub>12</sub>	30.0	6.0	30.0	17.0	10.0	19.0	7.0
N-C <sub>5</sub> H <sub>12</sub>	14.0	3.0	10.0	7.0	4.0	7.0	5.0
Squalane							
<sup>N-C</sup> 6 <sup>H</sup> 14	11			16			
N-C7 <sup>H</sup> 16	3			2			
Toluene	90			41			
N-C <sub>8</sub> H <sub>18</sub>	14			2			
E-C <sub>6</sub> H <sub>5</sub>	21			9			
M-XYL	37			22			
0-XYL	15			9			
N-C <sub>9</sub> H <sub>20</sub>	38			2			
N-P-C6H5	3			1			
N-C <sub>10</sub> H <sub>22</sub>	258			5			
Beckman 6800							
THC	3066	1785	1850				
CH <sub>4</sub>	2270	1683	1767				
co	1560	281	336				

TABLE C-13 (continued)

Date	11/12/76 9-11		11/12/76 1330 - 1530		11/16/76 9-11	11/16/76 11-13	11/16/76 13-15
Dura Pak							
<sup>C</sup> 2 <sup>H</sup> 6	7.0	6.7	6.3	42.1	30.4	9.0	11.4
C <sub>2</sub> H <sub>4</sub>	11.9	9.2	9.2	121.6	54.7	7.8	13.2
С <sub>3</sub> Н <sub>8</sub>	7.7	6.5	6.4	223.8	105.7	18.7	34.1
С <sub>2</sub> Н <sub>2</sub>	7.9	4.5	4.8	51.0	39.0	6.0	8.4
IsoC <sub>4</sub> H <sub>10</sub>	2.7	2.9	2.1	65.0	30.7	5.7	8.8
N-C4H10	11.8	11.5	7.6	174.9	148.2	24.9	43.8
<sup>C</sup> 3 <sup>H</sup> 6	2.2	1.5	1.2	24.6	14.4	1.5	2.4
IsoC <sub>5</sub> H <sub>12</sub>	11.0	9.1	6.4	118.0	75.0	11.0	17.0
N-C <sub>5</sub> H <sub>12</sub>	1.0	1.2	1.0	43.0	31.0	4.0	6.0
<u>Squalane</u>							
N-C <sub>6</sub> H <sub>14</sub>	2			48			
N-C7 <sup>H</sup> 16	1			6			
Toluene	6			40			
N-C <sub>8</sub> H <sub>18</sub>	1			5			
E-C <sub>6</sub> H <sub>5</sub>	4			21			
M-XYL	10			48			
0-XYL	4			16			
N-C9 <sup>H</sup> 20				3			
N-P-C <sub>6</sub> H <sub>5</sub>	1			3			
N-C <sub>10</sub> H <sub>22</sub>	1			11			
Beckman 6800							
THC	2010	1990	2960	4470	3360	2090	2280
CH <sub>4</sub>	1500	1480	1470	2550	1890	1440	1480
coʻ	490	250	280	2640	2160	410	530

TABLE C-13 (continued)

Date	11/18/7 7-9	6 11/18/76 9-11
Dura Pak	, ,	J 11
C <sub>2</sub> H <sub>6</sub>	58.6	60.6
C <sub>2</sub> H <sub>4</sub>	23.2	
C <sub>3</sub> H <sub>8</sub>	69.1	
C <sub>s</sub> H <sub>2</sub>	15.9	
IsoC <sub>4</sub> H <sub>10</sub>	31.6	28.5
N-C <sub>4</sub> H <sub>10</sub>	197.3	
C <sub>3</sub> H <sub>6</sub>	6.3	7.8
°3''6 IsoC <sub>5</sub> H <sub>12</sub>	57.0	55.0
	30.0	28.0
N-C <sub>5</sub> H <sub>12</sub>	30.0	20.0
Squalane		
<sup>N-C</sup> 6 <sup>H</sup> 14	6	
N-C7 <sup>H</sup> 16	2	
Toluene	24	
N-C <sub>8</sub> H <sub>18</sub>	2	
E-C <sub>6</sub> H <sub>5</sub>	4	
M-XYL	16	
O-XYL	4	
N-C <sub>9</sub> H <sub>20</sub>	2	
N-P-C <sub>6</sub> H <sub>5</sub>	1	
N-C <sub>10</sub> H <sub>22</sub>	5	
Beckman 6800		
THC	2920	2990
CH <sub>4</sub>	1740	1740
co	1350	1690

TABLE C-14. RAMS STATION 114

			FALL INT	ENSIVE			
			Mean	9	Standard D	eviat <b>i</b> on	
THC			2451 ppl	bC	1299 ppb	С	
CH <sub>4</sub>			1541 pp	bC	581 ppb	С	
co			1187 pp	bC	1392 ppb	С	
${\rm C_2H_2}$			19 ppl	ьС	22 ppb	С	
Date	11/8/76 7-9	11/8/76 13-15	11/10/7 7-9	6 11/10/76 7-9 Tank	5 11/10/76 9-11 Tank	11/10/76 11-13	11/10/76 13-15
<u>Dura Pak</u>					(Upwind)		
<sup>C</sup> 2 <sup>H</sup> 6	11.7	6.7	9.4	6.9	5.1	9.9	4.9
<sup>C</sup> 2 <sup>H</sup> 4	9.6	10.7	16.6	7.8	3.6	5.6	7.0
C3H8	8.0	11.2	7.9	6.4	7.2	8.1	4.7
С <sub>2</sub> Н <sub>2</sub>	9.0	8.0	15.1	9.2	2.3	1.7	1.6
IsoC <sub>4</sub> H <sub>10</sub>	5.0	6.1	4.4	3.4	1.1	2.4	4.0
N-C4 <sup>H</sup> 10	12.2	29.6	16.4	12.2	3.8	11.1	16.8
<sup>C</sup> 3 <sup>H</sup> 6	3.9	2.3	4.1	2.9	1.9	0.9	
IsoC <sub>5</sub> H <sub>12</sub>	10.0	17.0	15.0	10.0	2.0	11.0	9.0
N-C <sub>5</sub> H <sub>12</sub>	10.0	8.0	6.0	4.0	2.0	6.0	2.0
<u>Squalane</u>							
N-C <sub>6</sub> H <sub>14</sub>	8		6	3			
N-C <sub>7</sub> H <sub>16</sub>	3		2	7			
Toluene	19		13	9			
N-C8 <sup>H</sup> 18	7		1	1			
E-C <sub>6</sub> H <sub>5</sub>	7		5	9			
M-XYL	15		16	26			
0-XYL	6		8	10			
N-C <sub>9</sub> H <sub>20</sub>	1		BDL*	1			
N-P-C <sub>6</sub> H <sub>5</sub>	1		1	1			
N-C <sub>10</sub> H <sub>22</sub>	2		1	2			
Beckman 6800							
THC	2654	2254					
CH <sub>4</sub>	2140	1863					
co	1330	536					

<sup>\*</sup> Below detection limits

TABLE C-14 (continued)

Date	11/12/76 7-9	11/12/76 0930 - 1130	11/12/76 1130 - 1330	11/12/76 1330 - 1530	Tank 11/12/76 7-9 (Upwind)	Tank 11/12/76 9-11 (Upwind)	7-9
Dura Pak							
C <sub>2</sub> H <sub>6</sub>	7.8	6.2	5.7	5.4	8.2	6.5	37.0
C <sub>2</sub> H <sub>4</sub>	11.0	8.3	19.5	4.9	5.3	3.3	94.2
C3H8	9.6	6.8	3.4	5.3	10.3	7.4	39.4
C <sub>2</sub> H <sub>2</sub>	7.4	3.9	28.2	9.3	8.2	4.8	72.8
IsoC <sub>4</sub> H <sub>10</sub>	1.7	1.2	2.8	1.7	1.8	1.4	78.0
N-C4H10	7.8	6.2	7.0	5.0	5.9	4.3	235.9
<sup>C</sup> 3 <sup>H</sup> 6	1.6	1.1	1.1	1.1	2.6	1.9	36.1
IsoC <sub>5</sub> H <sub>12</sub>	4.0	4.0	5.0	4.0	4.0	3.0	156.0
N-C <sub>5</sub> H <sub>12</sub>	4.0	2.0	3.0	2.0	3.0	2.0	71.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>	NM**	10			2	5	
N-C <sub>7</sub> H <sub>16</sub>	1				1		
Toluene	4	3			5	3	
N-C <sub>8</sub> H <sub>18</sub>	1	1			1	1	
E-C <sub>6</sub> H <sub>5</sub>	3	1			7	5	
M-XYL	11	4			23	18	
0-XYL	3	1			9	5	
N-C <sub>9</sub> H <sub>20</sub>	Trace	Trace			BDL*	BDL*	
N-P-C <sub>6</sub> H <sub>5</sub>	Trace	Trace			1	BDL*	
N-C <sub>10</sub> H <sub>22</sub>	1	1			2	1	
Beckman 6800							
THC	2030	1890	2060	1970			5170
CH <sub>4</sub>	1540	1470	1460	1450			1880
co	350	210	170	220			4010

<sup>\*</sup> Below detection limits
\*\* Not measurable (because of interference etc.)

TABLE C-14 (continued)

Date Dura Pak	11/16/76 9-11	11/16/76 13-15	11/18/76 7-9	Tank 11/18/76 7-9 (Upwind)	
C <sub>2</sub> H <sub>6</sub>	40.3	6.1	66.2	63.0	
C <sub>2</sub> H <sub>2</sub>	72.9	38.4	80.2	23.9	
C <sub>3</sub> H <sub>8</sub>	72.5	5.3	69.6	66.4	
• •	69.1	16.7	37.1	32.1	
C <sub>2</sub> H <sub>2</sub>	25.3	6.8	19.6	17.3	
IsoC <sub>4</sub> H <sub>10</sub>	108.9	24.2	67.2	57.1	
N-C <sub>4</sub> H <sub>10</sub> C <sub>3</sub> H <sub>6</sub>	23.0	7.1	8.7	10.0	
• •	84.0	18.0	43.0	34.0	
1soC <sub>5</sub> H <sub>12</sub>	39.0	10.0	25.0	19.0	
N-C <sub>5</sub> H <sub>12</sub>	39.0	10.0	23.0	19.0	
Squalane					
N-C <sub>6</sub> H <sub>14</sub>			24	13	
<sup>N-C</sup> 7 <sup>H</sup> 16	9		4	4	
Toluene	57		39	37	
N-C8 <sup>H</sup> 18	6		3	2	
E-C <sub>6</sub> H <sub>5</sub>	17		12	8	
M-XYL	43		27	23	
0-XYL	17		12	8	
N-C <sub>9</sub> H <sub>20</sub>	4		2	1	
N-P-C <sub>6</sub> H <sub>5</sub>	4		1	1	
N-C <sub>10</sub> H <sub>22</sub>	10		NM*	2	
Beckman 6800					
THC	4010		2990		
CH <sub>4</sub>	2190		1770		
co	3280		1550		

<sup>\*</sup> Not measurable (because of interference etc.)

TABLE C-15. RAMS STATION 115

			ALL INTEN Mean		Standard	Deviation	n
THC			2321 p	nbC	417		, <u>•</u>
CH <sub>4</sub>			1637 p		225		
CO CO			666 p		388		
C <sub>2</sub> H <sub>2</sub>			9 p			ppbC	
	11.0.776	11 /0 /76					
Date	11/8/76 7-9:45	13-15	11/10/76 7-9	9-11	11/10//6	13-15	9:30 - 11:30
<u>Dura Pak</u>							
<sup>C</sup> 2 <sup>H</sup> 6	11.7	5.4	8.4	6.4	4.8	5.7	13.0
	40.7	27.6	33.4	32.5	42.6	45.2	28.1
С <sub>2</sub> Н <sub>4</sub> С <sub>3</sub> Н <sub>8</sub>	20.5	5.4	10.3	7.0	5.3	5.8	63.7
C <sub>2</sub> H <sub>2</sub>	14.9	3.8	7.3	2.1	2.7	2.5	7.4
IsoC <sub>4</sub> H <sub>10</sub>	8.6	1.7	5.4	3.2	1.8	2.2	28.5
N-C4H10	24.9	8.1	15.1	8.9	7.9	12.9	53.1
C <sub>3</sub> H <sub>6</sub>	5.7	1.2	3.2	0.9	BDL*	BDL*	17.5
IsoC <sub>5</sub> H <sub>12</sub>	19.0	6.0	11.0	6.0	2.0	8.0	33.0
N-C <sub>5</sub> H <sub>12</sub>	9.0	3.0	4.0	3.0	2.0	3.0	14.0
Squalane							
N-C <sub>6</sub> H <sub>14</sub>	8		17				14
N-C <sub>7</sub> H <sub>14</sub>	2		1				2
Toluene	12		9				9
N-C8H18	1		1				2
E-C <sub>6</sub> H <sub>5</sub>	3		5				4
M-XYL	12		17				9
O-XYL	5		6				3
N-C <sub>9</sub> H <sub>20</sub>	1		7				1
N-P-C <sub>6</sub> H <sub>5</sub>	1		1				TRACE
N-C <sub>10</sub> H <sub>22</sub>	4		5				2
Beckman 6800							
THC	2285		1967				2360
CH <sub>4</sub>	2152		1841				1500
<u>CO</u>	768		350				350

<sup>\*</sup> Below detection limits

TABLE C-15 (continued)

Date	11/12/76		11/16/76		11/16/76	11/16/76	11/18/76
	1 <b>1</b> 30 <b>-</b> 1330	1330 - 1530	7-9	9-11	11-13	13-15	7-9
<u>Dura Pak</u>							
<sup>С</sup> 2 <sup>Н</sup> 6	8.3	8.1	23.9	15.0	9.2	10.6	65.8
C <sub>2</sub> H <sub>4</sub>	24.4	60.0	54.2	30.9	30.1	32.1	37.9
C <sub>3</sub> H <sub>8</sub>	16.4	19.9	48.5	26.9	20.6	19.6	91.0
C <sub>2</sub> H <sub>2</sub>	6.2	6.7	24.5	13.8	5.8	8.6	17.2
IsoC <sub>4</sub> H <sub>10</sub>	14.5	19.0	11.6	8.0	6.0	5.5	24.0
N-C4H10	20.9	30.9	41.4	32.5	21.0	19.4	66.9
<sup>C</sup> 3 <sup>H</sup> 6	4.1	5.8	10.6	5.0	1.2	2.1	12.2
IsoC <sub>5</sub> H <sub>12</sub>	13.0	21.0	32.0	16.0	12.0	13.0	33.0
N-C <sub>5</sub> H <sub>12</sub>	5.0	8.0	16.0	9.0	4.0	5.0	19.0
<u>Squalane</u>							
N-C <sub>6</sub> H <sub>14</sub>			8				8
N-C7 <sup>H</sup> 16			3				3
Toluene			15				14
N-C8 <sup>H</sup> 18			3				2
E-C <sub>6</sub> H <sub>5</sub>			6				6
M-XYL			18				15
0-XYL			8				5
N-C <sub>9</sub> H <sub>20</sub>			1				1
N-P-H <sub>5</sub>			1				1
N-C <sub>10</sub> H <sub>22</sub>			4				3
Beckman 6800							
THC	2000		2620	2500	2100	2150	2720
CH <sub>4</sub>	1450		1710	1580	1450	1500	1770
co	300		1070	840	430	470	980

TABLE C-15 (continued)

Date Dura Pak	11/18/76 9-11		Bag Box 1 11/18/76 13-15	Tank	Bag Box 2 11/18/76 13-15
C <sub>2</sub> H <sub>6</sub>	65.1	48.4	28.1	27.9	27.8
<sup>2</sup> <sup>6</sup> C <sub>2</sub> H <sub>4</sub>	74.7	65.6	35.1	6.9	19.6
C <sub>3</sub> H <sub>8</sub>	91.2	65.7	35.2	35.5	35.2
C <sub>2</sub> H <sub>2</sub>	26.1		5.7	5.7	6.7
IsoC <sub>4</sub> H <sub>10</sub>	24.3	16.5		10.6	10.8
N-C4H <sub>10</sub>	85.5	55.8	40.2	36.9	36.7
C <sub>3</sub> H <sub>6</sub>	7.0	4.5	1.3	2.6	1.0
IsoC <sub>5</sub> H <sub>12</sub> .	46.0	27.0	21.0	18.0	21.0
N-C <sub>5</sub> H <sub>12</sub>	27.0	14.0	10.0	8.0	6.0
Squalane					
N-C <sub>6</sub> H <sub>14</sub>				30	21
N-C <sub>7</sub> H <sub>16</sub>				1	1
Toluene				10	12
N-C <sub>8</sub> H <sub>18</sub>				7	1
E-C <sub>6</sub> H <sub>5</sub>				5	6
M-XYL				17	19
0-XYL				5	5
N-C <sub>9</sub> H <sub>22</sub>				1	1
N-P-C6H5				1	1
N-P-C <sub>6</sub> H <sub>5</sub> N-C <sub>10</sub> H <sub>22</sub>				2	2
Beckman 6800					
THC	3020	2580	2350		2270
CH <sub>4</sub>	1830	1710	1580		1560
CO	1610	690	450		410

# APPENDIX D SPECIAL STUDIES

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#### D.O INTRODUCTION

Appendix D contains the results obtained from special studies in table form in order to provide an overview of the data.

TABLE D-1. ROADWAY SAMPLES COLLECTED AND ANALYZED 9/22/76 \*

	Babler Park	Babler Park	Manchester	Ultrapure	Interstate
	#_ 5	Repeat	Road	Air	40
Dura Pak					
<sup>C</sup> 2 <sup>H</sup> 6	4.1	3.7	22.3	0.3	18.9
C <sub>2</sub> H <sub>4</sub>	5.2	5.4	157.1	2.2	127.3
C <sub>3</sub> H <sub>8</sub>	5.7	5.3	27.6	0.3	14.4
$C_2H_2$	3.0	3.2	111.5	0.9+	83.2
IsoC <sub>4</sub> H <sub>10</sub>	0.8	0.8	15.2	BDL**	11.3
N-C4H10	3.4	3.3	84.1	1.2	62.7
C <sub>3</sub> H <sub>6</sub>	1.8	1.4	73.7	0.7	60.3
IsoC <sub>5</sub> H <sub>12</sub>	1.7	1.8	125.0	0.6	84.0
N-C <sub>5</sub> H <sub>12</sub>	1.5	1.5	68.0	BDL**	41.0
Squalane					
N-C <sub>6</sub> H <sub>14</sub>			46.0	NM+	64
N-C <sub>7</sub> H <sub>16</sub>			15.0	NM+	32
Toluene	6		72.0	9	165
N-C8H18	1		12.0	Trace	16
E-C <sub>6</sub> H <sub>5</sub>	9		30	10	68
M-XYL	30		78	33	136
0-XYL	4		30	6	65
N-C <sub>9</sub> H <sub>20</sub>	Trace		4	Trace	4
N-P-C <sub>6</sub> H <sub>5</sub>	BDL**		5	1	6
N-C <sub>10</sub> H <sub>22</sub>	NM +		16	1	NM +
10 22					
Beckman 68	300				
THC	1930		4010	210	3450
CH <sub>4</sub>	1850		2270		1960
Ć0	290		7850	240	6510

<sup>\*</sup> Concentrations in ppbC \*\*Below detection limits + Not measurable (because of interference etc.)

TABLE D-1 (continued)\*

	Interstate	Babler
	270	Duplicate
Duves Dale		
Dura Pak	42.0	4.6
<sup>C</sup> 2 <sup>H</sup> 6 <sup>C</sup> 2 <sup>H</sup> 4	253.0	
C <sub>2</sub> ''4 C <sub>3</sub> H <sub>8</sub>	54.9	
C <sub>2</sub> H <sub>2</sub>	190.1	
<sup>1</sup> soC <sub>4</sub> H <sub>10</sub>	19.8	
N-C <sub>4</sub> H <sub>10</sub>	121.4	
<sup>C</sup> 3 <sup>H</sup> 6		
IsoC <sub>5</sub> H <sub>12</sub>	159.1	2.9
N-C <sub>5</sub> H <sub>12</sub>	79.7	1.5
<u>Squalane</u>		
N-C <sub>6</sub> H <sub>14</sub>	38	
N-C7 <sup>H</sup> 16	51	
Toluene	239	
N-C <sub>8</sub> H <sub>18</sub>	45	
E-C <sub>6</sub> H <sub>5</sub>	107	
M-XYL	277	
0-XYL	91	
N-C <sub>9</sub> H <sub>20</sub>	21	
N-P-C <sub>6</sub> H <sub>5</sub>	18	
N-C <sub>10</sub> H <sub>22</sub>	41	
	• • •	
Beckman 68		
THC	6250	
CH <sub>4</sub>	3280	
<u>CO</u>	11360	

<sup>\*</sup> Concentrations in ppbC

TABLE D-2. ROADWAY SAMPLES COLLECTED 9/22/76 ANALYZED 9/24/76\*

	I-270	I-40	Manchester Road	Ultrapure Air	I-270 Duplicate
Dura Pak					
C2H6	47.1	19.1	22.2	0.4	40.8
C <sub>2</sub> H <sub>4</sub>	280.1	121.0	148.6	2.8	242.6
C <sub>3</sub> H <sub>8</sub>	61.5	14.2	26.7	0.4	53.1
C <sub>2</sub> H <sub>2</sub>	208.2	78.0	99.5	0.6	184.8
IsoC <sub>4</sub> H <sub>10</sub>	24.2	11.1	14.5	0.2	20.3
N-C <sub>4</sub> H <sub>10</sub>	138.3	61.3	80.6	1.4	118.5
C <sub>3</sub> H <sub>6</sub>	125.0	57.3	76.0	0.9+	108.6
IsoC <sub>5</sub> H <sub>12</sub>	191.0	81.0	119.0	0.7	165.0
N-C <sub>5</sub> H <sub>12</sub>	85.0	38.0	64.0	0.7	77.0
Squalane					
N-C <sub>6</sub> H <sub>14</sub>	89	27	41	1	
N-C <sub>7</sub> H <sub>16</sub>	<b>2</b> 8	14	15	1	
Toluene	140	67	73	12	
N-C8H18	25	12	14	2	
E-C <sub>6</sub> H <sub>5</sub>	66	37	36	16	
M-XYL	131	90	92	53	
0-XYL	56	32	36	10	
N-C <sub>9</sub> H <sub>20</sub>	10	1	4	NM +	
N-P-C <sub>6</sub> H <sub>5</sub>	10	2	5	NM +	
N-C <sub>10</sub> H <sub>22</sub>	25	5	9	NM +	

<sup>\*</sup> Concentrations in ppbC + Not Measurable (because of interference etc.)

TABLE D-3. ROADWAY SAMPLES COLLECTED AND ANALYZED 9/27/76 \*

	Bag 3 No. on 270		West on	West on	N. on 270	Bag 1 (Dup1) West on Manchester
Dura Pak						
<sup>C</sup> 2 <sup>H</sup> 6	38.6	38.4	16.1	17.1	21.4	16.0
$C_2H_4$	306.6	305.3	94.0	72.9	115.9	93.9
C3H8	16.6	16.6	12.0	12.4	14.1	12.2
C <sub>2</sub> H <sub>2</sub>	355.4	361.0	105.5	68.7	67.1	109.6
IsoC <sub>4</sub> H <sub>10</sub>	33.5	33.0	12.0	8.4	13.3	12.0
N-C4H10	180.3	179.1	57.6	32.0	60.8	57.5
C <sub>3</sub> H <sub>6</sub>	141.2	141.4	39.1	26.4	49.0	39.2
	136.0	243.0	75.0	38.0	73.5	60.0
N-C <sub>5</sub> H <sub>12</sub>	135.0	134.0	42.0	19.8	43.4	36.4
Squalane						
N-C <sub>6</sub> H <sub>14</sub>	91		45	30	27	
N-C7H16	31		11	6	11	
Toluene	126		44	30	51	
N-C <sub>8</sub> H <sub>18</sub>	21		52	6	10	
E-C <sub>6</sub> H <sub>5</sub>	55		93	14	22	
M-XYL	123		88	40	98	
0-XYL	61		27	14	22	
N-C <sub>9</sub> H <sub>20</sub>	6		135	2	3	
N-P-C <sub>6</sub> H <sub>5</sub>	22		894	6	5	
N-C <sub>10</sub> H <sub>22</sub>	16		4331	3	5	
Beckman 68	300					
THC	6000		9000	2900	3500	10020
CH <sub>4</sub>	1910		1800	1850	1940	1840
<u>co</u>	14600	<del></del>	5850	5190	5540	5890

<sup>\*</sup> Concentrations in ppbC

TABLE D-4. ROADWAY SAMPLES COLLECTED AND ANALYZED 10/29/76\*

	W. on Manchester I-270 to Weidman	W. on Manchester Weidman to Ries	S. on I-270 Manchester to I-44
Dura Pak			
<sup>C</sup> 2 <sup>H</sup> 6	36.1	53.6	33.0
C <sub>2</sub> H <sub>4</sub>	215.1	259.1	207.9
С <sub>3</sub> Н <sub>8</sub>	47.0	40.5	48.5
C <sub>2</sub> H <sub>2</sub>	205.4	257.8	134.7
Iso-C <sub>4</sub> H <sub>10</sub>	32.0	33.0	19.9
N-C <sub>4</sub> H <sub>10</sub>	153.3	194.3	114.8
С <sub>3</sub> Н <sub>6</sub>	95.6	114.1	93.7
IsoC <sub>5</sub> H <sub>12</sub>	145.2	182.6	116.7
N-C <sub>5</sub> H <sub>12</sub>	61.4	75.8	62.9
Squalane			
N-C <sub>6</sub> H <sub>14</sub>	38	74	44
N-C <sub>7</sub> H <sub>16</sub>	19	25	20
Toluene	94	119	83
N-C <sub>8</sub> H <sub>18</sub>	12	20	20
E-C <sub>6</sub> H <sub>5</sub>	38	41	24
M- XYL	90	129	74
0-XYL	47	52	41
N-C <sub>9</sub> H <sub>20</sub>	5	8	31
N-P-C <sub>6</sub> H <sub>5</sub>	7	12	6
N-C <sub>10</sub> H <sub>22</sub>	9	15	NM**
Beckman 6800			
THC	5230	6410	5000
CH <sub>4</sub>	2350	2690	2090
co	12360	15660	8970

<sup>\*</sup> Concentration in ppbC
\*\* Not measurable (because of interference etc.) (continued)

	W. on Olive I-44 to Manchester	W. on Olive I-270 to Eatherton	Dupl. Analysis W. on Manchester I-270 to Weidman
Dura Pak			
с <sub>2</sub> н <sub>6</sub>	29.8	5.3	37.3
C <sub>2</sub> H <sub>4</sub>	206.0	6.6	220.4
C <sub>3</sub> H <sub>8</sub>	42.3	11.8	47.8
C <sub>2</sub> H <sub>2</sub>	127.4	5.8	208.4
Iso-C <sub>4</sub> H <sub>10</sub>	17.1	2.2	33.0
N-C <sub>4</sub> H <sub>10</sub>	95.5	7.7	156.8
C <sub>3</sub> H <sub>6</sub>	95.9	2.4	95.2
IsoC <sub>5</sub> H <sub>12</sub>	99.5	9.8	148.9
N-C <sub>5</sub> H <sub>12</sub>	51.0	2.2	61.0
<u>Squalane</u>			
N-C <sub>6</sub> H <sub>14</sub>	28	BDL	38
N-C <sub>7</sub> H <sub>16</sub>	15	1	21
Toluene	68	5	99
N-C8H <sup>18</sup>	10	2	13
E-C <sub>6</sub> H <sub>5</sub>	. 21	3	32
M-XYL	62	5	88
0-XYL	34	5	39
N-C9 <sup>H</sup> 20	4	1	5
N-P-C <sub>6</sub> H <sub>5</sub>	9	NM**	3
N-C <sub>10</sub> H <sub>22</sub>	14	NW**	8
Beckman 6800			
THC	4290	2030	
CH <sub>4</sub>	1990	1720	
CO	9580	350	

 $<sup>\</sup>star$  Concentration in ppbC

<sup>\*\*</sup> Not measurable (because of interference etc.)

C2H4 BAG SAMPLES AT STATION 101 COLLECTED & ANALYZED 10/22/76\* TABLE D-5.

BAG B0X 7-8	C2H6 C	гак С <sub>2</sub> Н <sub>4</sub>	$c_3 H_8$	$c_2H_2$	ISO-C4H10	$N-C_4H_{10}$	$c_3H_6$	ISO-C <sub>5</sub> H <sub>12</sub>	N-C <sub>5</sub> H <sub>10</sub>
	27.7	7.07	17.6	62.1	23.8	117.5	26.3	97.0	51.0
BAG BOX (REPEAT)	27.7	70.7	17.5	8.09	22.8	116.7	24.5	0.96	53.0
AUTO (UPWIND) 7-8	29.7	61.1	26.0	68.4	79.2	485.8	26.8	231.0	103.0
	24.0	71.6	19.8	57.3	28.6	143.0	23.4	107.0	52.0
8-9	21.9	53.8	19.1	53.7	21.9	98.4	25.3	91.0	48.0
	12.7	1740.5	18.0	21.5	17.5	134.7	8.4	58.0	29.0
CENTERMOST BLOWER OUTLET 10-11	11.0	9199.4	34.4	8.2	12.0	55.9	5.0	38.0	12.0
INOPERATIVE BLOWER OUTLET	8.2	92.9	16.0	10.1	7.6	27.7	3.6	21.0	10.0
OUTLET OF VACUUM PUMP	22.6	9042.8	32.9	11.6	7.7	32.2	17.5	24.0	12.0
PUMP BOX INTERIOR 13-14	9.5	1486.1	20.1	12.0	12.5	105.3	9.9	39.0	14.0
BAG BOX (DUPLICATE)	28.2	72.5	17.2	62.7	22.5	116.0	25.7	98.0	54.0

\* Concentrations in ppbC

_	TABLE D-6.	- 11	ETHYLENE E	SAG SAMP	BAG SAMPLING AT STATION 101 (WITH AIR SAMPLING PUMP)*	) LOL NOI	WITH AIR	SAMPL ING PI	UMP)*
	$c_2H_6$	C2H6 C2H4	$c_3H_8$	$c_2H_2$	ISO-C <sub>4</sub> H <sub>10</sub>	N-C4H10	C3H6	IS0-C <sub>5</sub> H <sub>12</sub>	N-C <sub>5</sub> H <sub>12</sub>
BAG 1 9-9:15	22.1	36.1	31.2	33.5	27.9	430.2	15.0	84.0	25.0
BAG 1 9-9:15	22.1	35.9	31.2	33.5	27.9	427.4	15.0	82.0	25.0
TANK 1 9-9:15	20.3	271.0	30.4	31.3	27.1	418.8	13.0	73.0	26.0
BAG 2 10:10-10:15	13.7	28.9	20.0	18.5	21.6	207.7	8.0	65.0	25.0
TANK 2 10:10-10:15	13.2	21.9	20.4	19.2	20.8	223.4	8.0	59.0	30.0
BAG 3 10:45-11	10.1	108.2	12.6	12.5	11.7	65.7	5.0	40.0	14.0
TANK 3 10:45-11	9.5	13.0	12.0	12.4	11.7	64.5	5.0	39.0	18.0
BAG BOX (NO PUMP) 9-11		16.6 1765.4	24.9	29.3	23.5	259.5	11.0	75.0	24.0
BAG 1 9-9:15 Duplicate	21.6	36.4	31.3	34.3	27.6	431.5	14.0	77.0	30.0

Collected and Analyzed 11/2/76

\* Concentrations in ppbC

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ETHYLENE BAG SAMPLING AT STATION 124 (WITH AIR SAMPLING PUMP)\* TABLE D-7.

	Dura rak	Pak							
C	C2H6	$C_2H_4$	C3H8	$c_2H_2$	ISO-C4H10	N-C4H10	C3H6	ISO-C <sub>5</sub> H <sub>12</sub>	N-C <sub>5</sub> H <sub>10</sub>
BAG 2 10-10:30	4.3	7.5	4.8	1.6	0.8	4.4	0.7	4.0	2.0
BAG 2 (REPEAT) 10-10:30	4.5	7.5	4.8	1.9	[:	4.5	1.0	4.0	2.0
TANK 2 10-10:30	4.5	6.2	5.6	2.6	2.0	4.0	7	5.0	3.0
BAG 3 10:45-11:10	4.5	7.9	4.4	1.5	6.0	3.3	0.7	3.0	2.0
TANK 3 10:45-11:10	4.5	4.5	4.0	1.9	0.8	2.3	6.0	2.0	2.0
BAG 4 (BAG BOX - NO PUMP) 10-11:15	4.7	21.4	4.8	1.2	1.3	7.2	1.0	0.0	2.0
TANK 1 (OUTSIDE AIR) 9:30-9:45	4.4	2.2	5.8	2.0	2.0	15.2	7	4.0	3.0
BAG 5 (STATION AIR)	5.8	2716.8	7.2	1.9	1.6	0.9	1.1	2.0	2.0
BAG 2 (DUPLICATE) 10-10:30	4.3	7.3	4.9	1.6	9.0	4.2	£0	4.0	2.0

Collected and Analyzed on 11/3/76

\* Concentrations in ppbC

TABLE D-8. ETHYLENE	-8.		BAG SAM	SAMPLING AT		STATION 124 (WITH AIR SAMPLING PUMP)*	R SAMPL	ING PUMP)*	
	Dura Pak C <sub>2</sub> H <sub>6</sub>	k C <sub>2</sub> H <sub>4</sub>	c <sub>3</sub> H <sub>8</sub>	C2H2	ISO-C <sub>4</sub> H <sub>10</sub>	N-C4H10	$c_3$ H $_6$	IS0-C <sub>5</sub> H <sub>12</sub>	N-C <sub>5</sub> H <sub>12</sub>
BAG 2 9:55-10:10	5.4	11.4	4.7	4.9	2.2	8.8	2.0	0.6	3.0
BAG 2 (REPEAT) 9:55-10:10	6.1	11.3	4.7	3.8	1.5	8.5	1.0	8.0	3.0
TANK 2 9:55-10:10	5.5	8.1	4.3	4.5	1.4	5.4	1.0	4.0	2.0
BAG 3 10:45-11	4.7	9.4	3.8	3.3	1.7	7.1	3.0	4.0	3.0
TANK 3 10:45-11	4.7	8.6	3.6	2.8	-	4.1	1.0	3.0	3.0
BAG 4 (STATION AIR -1) 10:20-10:45	5.2	2018.2	7.8	4.1	1.9	& &	2.0	0.9	
TANK 1 (AIR UPWIND OF STATION)	5.5	4.2	5.4	3.5	4.1	34.2	2.0	16.0	3.0
BAG 5 (BAG BOX-NO PUMP) 9:30-11	5.3	13.8	4.7	4.5	1.6	6.9	2.0	4.0	2.0
BAG 6 (STATION AIR -2 LEE SIDE OF STATION)	4.6	923.5	5.3	3.5	1.4	5.7	1.0	3.0	2.0
TANK 4 (AIR UPWIND OF STATION)	4.7	3.3	4.2	3.0	2.2	19.1	2.0	11.0	4.0
BAG 2 (DUPLICATE) 9:55-10:10	5.3	11.4	4.6	4.3	1.7	8.1	1.0	5.0	3.0

\* Concentrations in ppbC

Collected and Analyzed 11/4/76

(P	TECHNICAL REPORT DATA lease read Instructions on the reverse vefore com	oleting)
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#### 16. ABSTRACT

A gas chromatography laboratory was set up to analyze air samples collected in Teflon bags and stainless steel tanks. Samples were analyzed for total hydrocarbons, methane, CO, and  $C_2$ - $C_{10}$  hydrocarbons. A total of 455 samples, including replicates, were analyzed during the summer and fall of 1976. Many samples were collected at 12 of the Regional Air Monitoring Systems (RAMS) sites to yield data on spatial and temporal distributions of hydrocarbons. Additional sampling was performed to study ethylene contamination in and around RAMS stations. Roadway samples were collected to determine the composition of freshly emitted vehicular pollution. Quality control audits indicated good system performance during the study. Replicate samples indicated good reproducibility for samples stored for as long as six days in the Teflon bags.

All data, including sums of paraffins, olefins, aromatic, and total non-methane hydrocarbons are stored in the RAPS Data Bank at Research Triangle Park, North Carolina.

17.	KEY WO	ORDS AND DOCUMENT ANALYSIS	
а.	DESCRIPTORS	b.identifiers/open ended terms	c. COSATI Field/Group
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