CONFECTION

AIR QUALITY
SUMMARY
1976

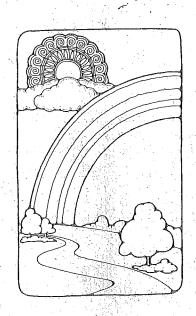




TABLE OF CONTENTS

		TABLE OF CONTENTS	Page
I.	. IN	TRODUCTION	1
	Α.	Total Suspended Particulate and Sulfur Dioxide Trends	1
	В.	Air Monitoring Network	4
	С.	Air Quality Standards	4
	D.	Pollutant Standards Index	6
	Ε.	Quality Assurance	8
II.	ТО	TAL SUSPENDED PARTICULATES	11
III.	SU	LFUR DIOXIDE	24
IV.	OZ	ONE	29
· V.	NI	TROGEN DIOXIDE	35
VI.	CA	RBON MONOXIDE	40
VII.	SP	ECIAL MONITORING STUDIES	44
	Α.	Asbestos Monitoring	44
	В.	Transport of Particulate Matter	45
	С.	Sulfur Dioxide Bubbler Monitoring	46
	D.	Historical Total Suspended Particulate Data	46
	Ε.	Passive Sampling Error	47
	F.	Low Volume TSP Measurements	48
	G.	Chemical Composition of Total Suspended Particulate Matter Samples	49
	Н.	Publications	56
TII.	CLI	MATOLOGICAL DATA	57
over	bу	CONNECTICUT AIR QUALITY SUMMARY - 1976	

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I. INTRODUCTION

This summary of 1976 ambient air quality levels in Connecticut is a compilation of all air pollutant measurements made at permanent Department of Environmental Protection (DEP) monitoring sites in the state.

A. Total Suspended Particulate Matter and Sulfur Dioxide Trends

Figures 1 and 2 show the long-term trends of Set I pollutants (particulate matter and sulfur dioxide) concentrations in Connecticut. In previous Air Quality Summaries data from all the Total Suspended Particulate (TSP) monitoring sites were included in developing the trend chart regardless of whether there were enough data for valid annual geometric means. In this year's summary the trend chart for TSP is based on those sites which had valid annual geometric means. Although this approach reduced the site population size in each year, the frequency distribution of geometric means did not change to any great extent.

The addition of TSP levels in 1976 to the trend chart shows a leveling off of the particulate air quality. It should be noted that the 1976 percentages include the annual geometric means calculated for the eight background TSP sites where lo-vol samplers are in operation (see Special Purpose Monitoring Section).

Figure 2 shows the sulfur dioxide trend. The most significant change between 1975 and 1976 is the increase in the number of sites with valid annual averages. This expansion in the monitoring network is responsible for the apparent improvement in the sulfur dioxide trend from 1975 to 1976. However, it can be seen from the trend chart that there has been a long-term improvement in sulfur dioxide levels. The secondary annual standard was not exceeded at any of the twelve sites in 1976 and only 5 sites have levels between 40 and 60 $\mu \rm g/m^3$.

FIGURE 1

TOTAL SUSPENDED PARTICULATE MATTER TREND

Percent of all sites in each concentration range

(1) Primary Annual Standard 75 $\mu g/m^3$ (2) Secondary Annual Standard 60 $\mu g/m^3$

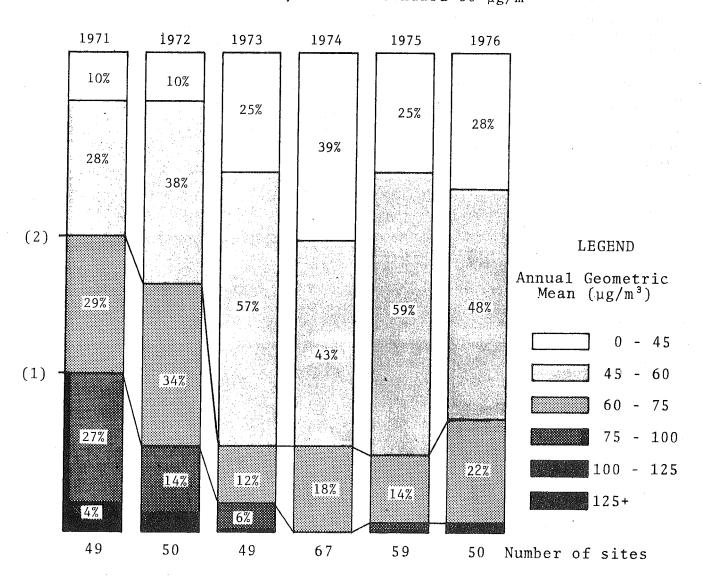


FIGURE 2
SULFUR DIOXIDE TREND

Percent of all sites in each concentration range

- (1) Primary Annual Standard 80 $\mu g/m^3$ (2) Secondary Annual Standard 60 $\mu g/m^3$
- 1971 1972 1973 1974 1975 1976 8% 20% 18% 22% 27% (2) ~ 38% 58% 37% LEGEND 60% Annual Arithmetic (1) -55% Mean $(\mu g/m^3)$ 31% 78% 0 - 40 27% 40 - 60 60 - 80 42% 15% 9% 80 - 100 20% 18% 100 + 9% 11 13 9 5 11 12 Number of sites

B. Air Monitoring Network

A computerized Air Monitoring network consisting of an IBM System 7 computer and 12 telemetered monitoring sites was put into full operation in 1975. Presently, up to 12 measurement parameters from each site are transmitted via telephone lines to the System 7 unit located in the DEP Hartford office. The data are then compiled into 24-hour summaries twice daily. The telemetered sites are located in the towns of Bridgeport, Danbury, Derby, Enfield, Greenwich, Groton, Hartford, New Britain, New Haven, Stamford, and Waterbury. The twelfth telemetered site is scheduled to become operational in 1977 in the town of Middletown.

Measured parameters include the pollutants sulfur dioxide, particulates (COHS), carbon monoxide, ozone, and meteorological data consisting of wind speed and wind direction, wind horizontal sigma, temperature, dew point, precipitation, barometric pressure and solar radiation.

The real-time capabilities of the System 7 telemetry network have enabled the Air Monitoring Unit to report the Air Quality Index for 12 towns on a daily basis while keeping a close watch for high pollution levels which may occur during adverse weather conditions throughout the year.

The complete monitoring network used in 1976 consists of:

- 63 Total Suspended Particulate sites
- 16 Sulfur Dioxide sites
- 12 Ozone sites
- 43 Nitrogen Dioxide sites
- 12 Carbon Monoxide sites

In April, 1976 a major revision to the network was completed which reduced the total number of monitoring sites. A complete description of all permanent air monitoring sites in Connecticut operated by DEP in 1976 is available from the Department of Environmental Protection, Air Compliance, State Office Building, Hartford, Connecticut 06115.

C. Air Quality Standards

Table 1 lists analysis methods and National Ambient Air Quality Standards (NAAQS) for each pollutant. The NAAQS were established by the U.S. Environmental Protection Agency (EPA) and are divided into two categories: primary, established to protect the public health; and secondary, established to protect plants and animals and to prevent economic damage.

Each standard specifies a concentration and an exposure time developed from studies of the effect of various levels of the different pollutants.

ASSESSMENT OF AMBIENT AIR OUALITY TABLE 1

			•		
	SAMPI, ING	OF ANALYSIS	NATIONAL	AMBIENT AIR	STANDARDS
POLLUTANT	PERIOD	REDUCTION	BASE	STANDARD	SECONDARY
				mdd gm/Bn	mg/m³ ppm
Total Suspended Particulates	24-Hours Every Sixth	24-Hour Average	Annual Geometric Mean	7.5	*09
			24-nour Concentration ²	260	150
Sulfur Oxides (Measured as Sulfur Dioxide)	Continuous 1	1-Hour Average	Annual Arithmetic Mean	80.03	60† .02
			Concentration ²	365 .14	260⁺ .10
			Concentration ²		1300 .5
Nitrogen Dioxide	24-Hours Every Sixth Day	24-Hour Average	Annual Arithmetic Mean	100 .05	Same as Primary
Photochemical Oxidants (Ozone)	Continuous	1-Hours Arrended	1 11 A 2		
		r-mour werage	ı-nour Average-	160 .08	Same as Primary
Hydrocarbons	Continuous¹	1-Hour Average	3-Hour Average ² (6-9 AM)	160** .24	Same as Primary
				mg/m³ ppm	mg/m³ ppm
Carbon Monoxide	Continuous¹	1-Hour Average	8-Hour Average ² 1-Hour Average ²	10 9 40 35	Same as Primary Same as Primary

 $^{1}\mathrm{EPA}$ assessment criteria requires 75% of possible data to compute valid averages. $^{2}\mathrm{Not}$ to be exceeded more than once per year.

*A guide to be used in assessing implementation plans to achieve the 24-hour standard. **For use as a guide in devising implementation plans to achieve oxidant standards. †Secondary Standard applies to State of Connecticut only. Units: $\mu g/m^3 = Micrograms$ per cubic meter; $mg/m^3 = Milligrams$ per cubic meter;

ppm = Parts per million

D. Pollutant Standards Index

The Pollutant Standards Index (PSI) is a daily air quality index recommended for common use in state and local agencies by the U.S. Environmental Protection Agency. Connecticut switched to reporting the PSI on a 7-day a week basis on November 15, 1976. The PSI incorporates five pollutants - carbon monoxide, sulfur dioxide, total suspended particulates, photochemical oxidants and nitrogen dioxide. The index converts each air pollutant concentration into a normalized number where the National Ambient Air Quality Standard for each pollutant corresponds to PSI = 100 and the Significant Harm Level corresponds to PSI = 500. Figure 3 shows the breakdown of index values for the commonly reported pollutants in Connecticut. Each day the pollutant with the highest PSI value of all the pollutants being monitored is reported, along with the dimensionless PSI number, and a descriptor word to characterize the daily air quality.

This information is available to the public each afternoon from the Connecticut Lung Association in East Hartford, and is transmitted to the news media by the National Weather Service with the afternoon forecast over the news service wires.

P.S.I. Value -500 400 -300 -200 -10050 1000+HAZARDOUS UNHEALTH-FUL VERY UN-HEALTHFUL OZONE MODERATE G005 µg/m³ 160 -800-400-807 1200-FIGURE 3 POLLUTANT STANDARDS INDEX P.S.I. Value -500 -300 -200 50 2100-HAZARDOUS +400 -100SULFUR DIOXIDE UNHEALTH-FUL VERY UN-HEALTHFUL MODERATE G00D µg/m³ 2620-1600 -365-80-800-P.S.I. Value -500 -300 875-HAZARDOUS \to 400 -200 50 -100PARTICULATES UNHEALTH-FUL VERY UN-HEALTHFUL MODERATE G00D µg/m³ 625-375-75-1000-260-

E. Quality Assurance

A vigorous and comprehensive Quality Assurance Program for air quality data encompasses a multitude of activities:

· Personnel training

· Site selection, evaluation and review

• Equipment evaluation, selection and modification when applicable

 Purchasing and inventory control of consumable supplies

 Instrument preventive maintenance, operation and calibration

 Calibration and traceability of working standards

· Sample collection and analysis

 Data recording, documentation, reduction, validation and reporting

· Interagency cross-checks

· Interlaboratory and instrument audits

The development of the above activities is an ongoing process in which detailed procedures are issued, constantly reviewed, updated and improved.

Data quality is a direct function of the implementation of these activities. In addition, with the advancement of instrument technology, personnel experience and improved quality control procedures for the operation, maintenance and calibration of monitoring equipment, the quality of data should improve from year to year until these advancements are balanced by other factors such as instrument degradation due to aging and personnel turnover (this balance has not yet been reached).

It is essential that data quality be assessed from an impartial source (EPA) that periodically performs quantitative audits on monitoring instruments, calibration systems, and laboratory functions. The results of Connecticut's DEP performance are summarized here in an effort to quantify the degree of data accuracy.

Particulates:

-Sample Weights

The EPA procedure of auditing the weighing of samples is no longer being performed because of a negative bias which exists at each subsequent weighing due to the loss of particulate matter during the sample shipping process.

-Flow Rates

The second parameter required to determine particulate concentrations is the sampling flow rate. Connecticut participated in

9 EPA audits, each of which contained 5 different flow rates for a total of 45 data points.

There were 4 data points which were outside the acceptable range defined by EPA, all of which were reported high. However, it should be noted that the unacceptable points were below the normal operating range of hi-vols. All audited values in the range of instrument operation were acceptable.

Sulfur Dioxide:

-Continuous Monitors

Nine instruments were audited by EPA during 1976 for a total of 54 data points. Performing a linear regression analysis on all these data points (CT vs EPA), produced a linear relationship of Y = 1.002 X + 0.0048 where Y = Connecticut's values and X = EPA's. The 95% confidence band for concentrations between 0 and 0.3 ppm was essentially constant and amounted to $\pm .038$ ppm. Put another way, any values between 0 and .3 ppm, probably deviated by a factor of no more than -.0332 to +.0428 ppm.

At the end of the year, a comparison was made between our primary calibration standard, 2 field calibration units and EPA, Region I's field calibrator. All results were within 5.5% of one another, well within the range of acceptability.

Ozone:

Instruments at 8 sites were audited by EPA during 1976 for a total of 46 data points. Performing a linear regression analysis on all these data points (CT vs EPA), produced a linear relationship of Y = 1.012 X - .0001 where Y = Connecticut's value and X = EPA's. The 95% confidence band for concentrations between 0 and 0.4 ppm was essentially constant and amounted to $\pm 0.029 \text{ ppm}$. Put another way, any value between 0 and 0.4 ppm, probably deviated by a factor of no more than -0.029 to +0.029 ppm, indicating excellent performance of the instruments and calibrations.

Nitrogen Dioxide:

-Bubblers

Fifteen EPA reagent samples were analyzed at the Environmental Chemistry Laboratory of the Connecticut Health Department to determine the accuracy of the analytical procedures. Two values were barely outside EPA's acceptable range and were 4.3% below and 6.1% above the mean expected values.

Carbon Monoxide:

Eight monitors were audited in 1976 per EPA's procedures (DEP personnel sample from 3 tanks of CO in nitrogen, where the CO concentration of each tank was unknown to them). Results indicate that at the upper end of the instruments' range (38.8 ppm input), the average reading was 0.5 ppm high with a standard deviation of 2.7 ppm. At midrange (18 ppm input) the average difference was 0.2 ppm with a standard deviation of 1.9 ppm. At the low end (7 ppm input), the average difference was -0.1 ppm ±1.3 ppm.

In the future, audits will be performed by EPA personnel, in conjunction with SO2 and O3 audits.

II. TOTAL SUSPENDED PARTICULATES

Conclusions:

In general, measured total suspended particulate (TSP) levels in Connecticut showed no significant change in air quality in 1976 as compared to 1975 (see Figure 1).

In 1976, 16 sites showed lower annual geometric means than in 1975, with 7 of these decreases being greater than 5 $\mu g/m^3$. The geometric means at 18 sites showed increases in 1976 over 1975, 6 of which increased more than 5 $\mu g/m^3$. When determining compliance with either the primary or secondary annual NAAQS for TSP the federal EPA recommends that only sites with at least 5 observations in each quarter of the year be evaluated. Using this criterion the primary annual standard was exceeded in Waterbury at site 123 while the secondary annual standard was exceeded at 12 sites in 1976, 5 more than in 1975 (many by a very slight margin).

Table 4 presents 1st and 2nd high 24-hour concentrations recorded at each site. There was no violation of the primary 24-hour standard recorded in 1976. Measured values exceeding the secondary 24-hour standard were recorded at 12 sites in 1976, 2 less than in 1975.

Discussion of Data:

Table 3 is the product of a computer program listing all monitoring sites used by the DEP. The data for each site includes the number of samples taken (generally, a maximum of 61 samples per year), the geometric mean, 95% confidence limits about the mean, the standard geometric deviation and statistical prediction of the number of days in each year the 24-hour primary and secondary NAAQS would have been exceeded if sampling had been conducted every day. This analysis, as were the national ambient standards, is based on the assumption that the particulate data are lognormally distributed.

Because manpower and economic limitations dictate that sampling of particulate matter occurs once every sixth day instead of every day, a degree of uncertainty as to whether the air quality at a site has either met or exceeded the national standards is introduced. This uncertainty can be quantified by determining 95% confidence limits about each of the annual geometric means. For example, (see Table 3), in Ansonia at site 003 in 1976, 61 samples were taken and a geometric mean of 59.8 $\mu g/m^3$ was calculated. However, the columns labeled "95-PCT-LIMITS" show the lower and upper limits for a 95% confidence interval of 53 and 68 $\mu g/m^3$, respectively. This means that if any other set of 61 samples were taken in 1976 at this site there is a 95% chance the geometric means would fall between these limits. Since the national secondary standard for particulates is within this interval, one cannot be 95% confident that the secondary standard was met in 1976.

In Table 2 all 1976 monitoring sites are examined for compliance with the standard using the statistical confidence limit criteria. The table shows that Waterbury 123 exceeded the primary annual standard with 95% confidence. It is uncertain whether the standard was either achieved or exceeded at 3 other sites. The table also shows that the secondary standard was exceeded with 95% confidence at 3 sites. Whether the secondary standard was exceeded is uncertain at 23 other sites. Comparing this to the results using the actual measured levels in the conclusions above, the 95% confidence method includes the same number of sites exceeding the primary standard and 9 less sites exceeding the secondary standard.

Table 4 presents the second high 24-hour concentrations recorded at each site. Although no violations of the primary 24-hour standard were measured, the statistical projections from Table III indicate that 3 sites having valid geometric means would have violated the standard had samples been collected every day. Violations of the secondary 24-hour standard were measured at 12 sites in 1976 but again, if samples were collected every day, statistical projections indicate that 24 additional sites would have recorded violations of the secondary standard.

Further TSP Work:

In 1977 the Air Monitoring Section will operate high volume samplers every third day at four locations. See also the Special Purpose Monitoring section of this report for discussion of Lo-Vol TSP sampling.

Sample Collection and Analysis:

Total suspended particulate levels are obtained from High Volume Samplers. These "Hi-Vols" resemble vacuum cleaners in their operation, with an 8" x 10" piece of fiberglass filter paper replacing the vacuum bag. The samplers operate every sixth day from midnight to midnight.

The matter collected on the filters is analyzed for weight and chemical composition. The flow through the filter is measured before and after sampling and the volume of air which has passed through the filter in 24 hours is calculated. The weight in micrograms (μg) divided by the volume of air in cubic meters (m^3) yields the pollutant concentration for the day, in micrograms per cubic meter. The chemical composition of the suspended particulate matter is determined as follows. A standardized strip of each of the Hi-Vol filters collected in each quarter year is cut-out and composited into one sample. This procedure is repeated three times so that for each site, three quarterly composited samples are made. One sample is digested in benzene and the residue is weighed. The weight of this residue represents the organic material in the sample and the result is reported as the benzene soluble fraction of the TSP in $\mu g/m^3$. Another sample is dissolved in water, re-fluxed and the resulting solution

is analyzed to determine the water soluble fraction of the TSP using wet chemistry techniques. Results are reported for each individual constituent of the water soluble fraction in $\mu g/m^3$. The last composited sample is digested in acid and the resulting solution is analyzed for the different metals in the TSP using an atomic absorption spectrophotometer. Results are reported for each individual metal in $\mu g/m^3$.

TABLE 2

CONFIDENCE OF TSP ANNUAL GEOMETRIC MEANS

Standard	Uncertain Whether Standard Has Been Achieved Or Exceeded		Derby 123 Greenwich 001 Greenwich 003 Greenwich 008		Norwalk 005 01d Saybrook 001 Stamford 003/123 Stamford 007 Stratford: 005 Torrington 123	_
Secondary	95% Confident Standard Has Been Exceeded (>60)	Bridgeport 123 Hartford 003 Waterbury 123				
tandard	Uncertain Whether Standard Has Been Achieved Or Exceeded	Bridgeport 123 Hartford 003 Torrington 123				
Primary Standard	95% Confident Standard Has Been Exceeded (>75)	Waterbury 123				

						TABLE 3				
CCNAEC	CCNNECTICUT DEFARTMENT	FARTN W	u. Ca	ENVIRONMENTA	L PRCTE	CT10.1	PAGE	1 AIR	CCMPLIANCE	MONITORING
PCLLUT	PCLLUTANIPARTICULATES	TICUL	ATES						DISTRIBUTION-	LOGNERMAL
TCWN N	NAME	SITE	YEAR	SAMPLES	GEON MEAN	95-PCT- LUWER	-LIMITS UPPER	STD GECM DEV	PREDICTED DAYS OVER 150 UG/M3	PRESICTED DAYS OVER 260 UG/M3
AN SONI A	ø	03	1976	61	59.8	53	68	1.715	16	-
BERLIN		01	1976	13	38.0	59	55	1.538		
BRIDGEPORT	PORT	01	1976	61	54.3	64	61	1.595	īΛ	
BRIDGEPORT	PORT	123	1976	90	68.4	61	11	1.638	20	-1
BRISTOL	. 1	01	1976	55	58 8	51	89	1.743	16	pmd
PRISTGL	. t	04	1976	64	9*09	<i>8</i> 3	69	1.658	13	.
S BURLINGTON	STON	01	1976		24.3	77	41	1.791		·
DANBURY	_	123	1976	90	Ç 23°€	47	09	1.671	ю	
DERBY		123	1976	58	53.9	48	61	1.634	_	
EAST H	HARTFORD	01	1976		35.4	21	09	2.212	13	2
EAST H	HARTFORD	0.5	1976	53	41.02	36	14	1.680	7	
EAST W	WINDSOR	01	1976	13	0.69	54	88	1.512	10	
ENFIELD		123	1976	56	43.2	38	65	1.638	. 7	
GREENWICH	НО1	10	1976	58	5.4.94	49	61	1.567	4	
GREENWICH	ICH	02	1976	16	54.6	7 7	99	1.502	~	
GREENWICH	ICH	03	1976	54	55.8	50	63	1.580	rU	
GREENWICH	LCH 1CH	04	1976	57	40.3	35	94	1.755	4	

CONNECTICUT	DEPARTMENT	MENT OF	ENVIRONMENTAL	PRCT	ECTION	PAGE	2 AIR	CCMPLIANCE	MCNITORING
PCLLUTANTPARTICULATES	AKTICU	LATES		·			· · · · · · · · · · · · · · · · · · ·	DISTRIBUTION	LGGNORMAL
TOWN NAME	SITE	YEAR	SAMPLES	GEOM MEAN	95-PCT- LOWER	-LIMITS UPPER	STD GECM DEV	PREDICTED DAYS OVER 150 US/M3	PREDICTED DAYS OVER 260 HG/W3
GREENWICH	6.0	1976	57	55.2	64	62	.65)))))
GRETON	123	1976	58	44.07	41	64	1.506		
HADDAM	0 5	1976	58	35.4	31	40	1.641	 1	
HARTFORD	02	1976	59	48 5	43	54	1.619	4	
HARTFORD	03	1976	58	73.5	19	81	1.496	13	
HARTFORD	04	1976	13	53°6	77	78	1.624	10	
-16 HARTFORD	05	1976	12	55.8	5 5	7.1	1.481	2	
HARTFCRD	123	1976	19	47°7	40	57	1.455		
MANCHESTER	01	1976	55	39.8	35	45	1.604	, , , , , , , , , , , , , , , , , , , 	
MANSFIELD	01	1976	14	40.5	31	54	1.646	, 2	
MERIDEN	02	1976	51	51.8	46	53 8	1.560	ı m	
MERIDEN	05	1976	9	62.5	54	73	1.917		ſĽ
MIDDLETOWN	03	1976	09	58.4	52	65	1,591	. 60	
MILFORD	Ŏ	1976	09	50.6	45	56	1.566	m	
MILFORD	. 02	1976	58	52.5	14	58	1 .538	2	
MORRIS	01	1976	. 21	35.0	25	20		2	
NAUGATUCK	01	1976	09	54.5	48	62	1.685	10	
NEW BRITAIN	03	1976	25	L. +99	25	73	1 0 6 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	. 20	· 📥

CCNNECTICUT DE	DEPARTMENT	ENT OF	ENVIRONMENTAL	TAL PRETECTION	ICM	PAGE	3 AIR	R COMPLIANCE	MCNITORING
PCLLUTANIPARTICULATE	TICUL	ATES						DISTRIBUTION-	LEGNORMAL
TCWN NAME	SITE	YEAR	SAMPLES	GEOM MEAN	95+PCT- LOWER	-LIMITS UPPER	STD GEGM DEV	PREDICTED DAYS OVER 150 UG/M3	PREDICTED DAYS OVER 260 UG/M3
NEW ERITAIN	04	1976	13	48.7	37	64	1.575	2	
NEW BRITAIN	123	1976	51	56.7	51	63	1.607	7	
NEW HAVEN	01	1976	58	58•J	51	99	I •699	13	
NEW HAVEN	0.2	1976	58	60.3	54	67	1.551	7	
NEW HAVEN	03	1976	15	49.2	41	. 65	1.396		
NEW HAVEN	05	1976	19	53.9	41	69	1.760	13	~
NORTH CANAAN	01	1976	59	39.6	34	46	1.794	4	
NORWALK	0.1	1976	10	68.3	52	06	1.470	တ	
NORWALK	0.5	1976	59	58°7	55	99	1.620	10	
NORWICH	01	1976	59	49.6	4.5	55	1.489	-	
CLD SAYBROCK	01	1976	58	63.8	57	7.1	1.569	10	
PUTNAM	02	1976	16	63°5	47	86	1.773	. 24	2
STAMFCRD	01	1976	11	68.9	50	94	1.609	20	1
STAMFORD	40	1976	, m	45.9	34	19	1.545	1	
STAMFORD	0.7	1976	58	57.5	51	49	1.577	1	
STAMFORD 03/	123	1976	61	9.09	55	19	1.538	7	
STRATFORD	01	1976	47	47.2	42	54	1.595	2	
STRATFORD	05	1976	09	0.09	54	19	1.567	∞.	
		,							

-17-

CCNNECTICUT DEPARTMENT OF ENVIRONMENTAL	DEPARTI	MENT OF	ENVIRONMEN	TAL PROTECTION	NOI	PAGE	4 AIR	R COMPLIANCE	MONITORING	
POLLUTANTPARTICULATES	ARTICUL	ATES						DISTRIBUTIONLOGNORMAL	LOGNORMAL	
TOWN NAME	SITE	YEAR	SAMPLES	GEOM MEAN	95-PCT- LOWER	95-PCT-LIMITS LOWER UPPER	STD GEOM DEV	PREDICTED DAYS OVER 150 UG/M3	PREDICTED DAYS OVER 260 UG/⊮3	
TORRINGTON	123	1976	57	67.7	59	77	1.702	24	7	
VOLUNTOWN	01	1976	.12	22.7	18	59	1.497			
MALLINGFORD	01	1976	09	58.4	52	65	1.608	ဆ		
WATERBURY	05	1976	09	60.1	54	67	1.625	10		
WATERBURY	03	1976	13	65.0	47	89	1.711	20	7	
WATERBURY	123	1976	09	86.5	76	86	1.689	58	P~~	
WATERFORD	01	1976	57	34.3	30.	39	1.633			
" WILLIMANTIC	01	1976	2	54.7	4.5	99	1.377			
WINCHESTER	Ö	1976	13	68.7	14	100	1.886	42	P	

TABLE 4
Total Suspended Particulates
1976 - Connecticut
24-Hour Concentration

		Da	ate 2nd	150 260
Site		High	High	0 100 200 300 400
				(micrograms per cubic meter)
Ansonia	003	2/24	1/7	197
Ber1in	001	1/13		66 (3 months)
Bridgeport	001	4/18	6/11	159
Bridgeport	123	6/17	2/25	
Bristol	001	9/15	8/28	200
Bristol	004	4/18	5/24	150 133
Burlington	001	3/25	·	55 (3 months)
Danbury	123	6/11	2/24	163 127
Derby	123	6/11	3/25	153 138
E. Hartford	001	3/1		93 (3 months)
E. Hartford	002	6/11	4/18	150 104
E. Windsor	001	2/24		(3 months)
Enfield	123	12/20	6/11	143
Greenwich	001	6/11	4/18	151
Greenwich	002	3/19		108 (3 months)
			ı	SECONDARY PRIMARY

		D	ate 2nd	150 260
Site		High	High	
				(micrograms per cubic meter)
Greenwich	003	3/25	5/6	126
Greenwich	004	3/25	6/11	145 137
Greenwich	008	6/11	4/19	177
Groton	123	4/18	6/11	105
Haddam	002	6/11	12/20	128
Hartford	002	6/11	2/25	140
Hartford	003	6/11	1/13	190 161
Hartford	004	2/24		121 (3 months)
Hartford	005	2/24		98 (3 months)
Hartford	123	11/26	,	100 (3 months)
Morris	001	2/24		107 (3 months)
Manchester	001	6/11	3/25	139
Mansfield	001	2/24		86 (3 months)
Meriden	002	6/11	1/13	143
Meriden	005	3/31	6/11	438
•			. '	SECONDARY PRIMARY

		D	ate 2nd	150	260
Site		High		100 2	00 300 400
Middletown	003	6/11	4/18	(mrcrograms p	er cubic meter)
Milford	001	6/11	4/18	 125)
Milford	002	6/11	4/18	136	
Naugatuck	001	3/19	6/11		
New Britain	003	6/11	9/15	214	
New Britain	004	1/13		100 (3 months)	
New Britain	123	2/25	6/11	149	
New Haven	001	6/11	8/28		
New Haven	002	6/11	4/18	167 132	
New Haven	003	2/24		80 (3 months)	
New Haven	005	1/13			
N. Canaan	001	6/11	12/14	186	
Norwalk	001	2/24		 (3 months)	
Norwalk	005	6/11	1/7	147 144	
Norwich	001	6/11	1/7	150	
01d Saybrook	001	6/11	2/24	178	
				SECONDARY	PRIMARY

		D	ate	1.50
Site		High	2nd High	150 260 <u>Q</u> 100 200 300 400
	, , 1			(micrograms per cubic meter)
Putnam	002	3/19		159 (3 months)
Stamford	001	2/24		131 (3 months)
Stamford	003	4/18		138 (5 months)
Stamford	004	1/7		88 (3 months)
Stamford	007	12/8	6/11	144
Stamford	123	12/20	11/26	142 115
Stratford	001	7/17	4/18	140 134
Stratford	005	4/18	2/24	135 134
Torrington	123	3/25	3/19	
Voluntown	001	3/19		46- (3 months)
Wallingford	001	2/24	3/19	169 158
Waterbury	002	6/11	9/15	159 135
Waterbury	003	2/24	:	175 (3 months)
Waterbury	123	2/26	3/25	281
Waterford	001	6/11	4/18	155 92
Willimantic	001	3/19		85 (3 months)
Winchester	001	2/24		137 (3 months)
				SECONDARY PRIMARY

III. SULFUR DIOXIDE

Conclusions:

At no monitoring site in Connecticut was the primary or secondary annual sulfur dioxide (SO₂) standard exceeded in 1976.

The primary 24-hour ambient standard for SO2 was not exceeded in Connecticut during 1976. Milford 002 recorded a violation of the secondary 24-hour ambient standard for SO2.

There was no violation of the 3-hour SO₂ standard recorded at monitoring sites in Connecticut in 1976.

Discussion of Data:

A total of 16 continuous SO₂ monitors recorded data in 14 towns during 1976. Eleven of these sites telemetered the data to the central computer in Hartford on a real-time basis. Sufficient data for valid annual means were recorded at 12 of these 16 sites in the network, up from 5 in 1975.

Method of Measurement:

The Air Monitoring Unit uses several types of instruments to continuously measure sulfur dioxide levels. The coulometric method is employed by Philips instruments, the flame photometric method by Bendix instruments.

Philips monitoring instruments were used at the following sites in 1976:

Bridgeport	001	Mil:	ford	002
Meriden	002	New	Haven	004
Middletown	003			

Bendix instruments were used at the following sites in 1976:

Bridgeport	123	Hartford	123
Danbury	123	New Britain	123
Derby	123	New Haven	123
Enfield	123	Stamford	123
Greenwich	004	Waterbury	123
Groton	123		٠

Connecticut also used modified West-Gaeke sulfur dioxide bubblers at 26 sites, however, the Department regards all SO2 bubbler data to date as invalid due to problems associated with the collection, storage and transport of bubbler samples (see section on Special Studies for further discussion of bubbler data.)

TABLE 5 ANNUAL ARITHMETIC AVERAGES OF SULFUR DIOXIDE AT SITES WITH CONTINUOUS MONITORS

Primary NAAQS 80 μg/m³

Town		Site Name	1976	1975	1974	1973	1972	1971
Bridgeport	001	City Hall	47	63	42	44	62	76
Bridgeport	002	Fairfield Avenue Fire House			51	31	54	
Bridgeport	003	McKinley School		47	49	50	50	-
Bridgeport	123	Hallett Street	50	1	-			
Danbury	123	Western Conn. State College	37	31	1	MA 155	MA 5-4	en in
Derby	123	Dziadik Street	37	1				-
Enfield	123	Kosciuszko Junior H.S.	26	$(42)^2$			-	
Greenwich	001	Town Hall Annex			37	53	45	62
Greenwich	004	Bruce Golf Course	26	1	(29)2	29	33	43
Greenwich	800	Cos Cob Pumping Station			48	55	43	71
Groton	123	Fort Griswold State Park	29	(29)3		denis desar	-	em 1
Hartford	003	Public Library			48	69	61	91
Hartford	123	State Office Building	42	1		from from		
Meriden	002	Stoddard Building	(42)²	-		Acad Anna		
Middletown	003	City Hall	1	en ma	、			
Milford	002	Devon Community Center	(68)²	50	31	$(25)^2$		
New Britain		City Hall		1		(80)3	120	96
New Britain		Lake Street	31	1				
New Haven	004	Community Service Building	1	50	40	54	79	84
New Haven	800	Agricultural Station				38	41	51
New Haven	123	State Street	50	1			*** ***	
Norwalk	005	Health Department			44	50	62	65
Stamford	123	Health Department	45	(50)²	1	(78)²	90	119
Torrington	123	Franklin Avenue		l		60% mad	-	
Waterbury	001	City Hall			(56)4	84	93	103
Waterbury	123	Bank Street	31	1			pro 1000	-

¹Insufficient data for valid annual average or estimate (less than 6 months) ²Estimate based on partial data (6 to 9 months)

³Based upon questionable data ⁴September - December data missing

TABLE 6 Sulfur Dioxide 1976 - Connecticut 24-Hour Concentration

		D	ate		
Site	•	High	2nd High	0 100 200 365 0 100 200 300	400
		mign	111811	(micrograms per cubic meter)	400
Bridgeport	001	2/4	1/21	213 200	•
Bridgeport	123	1/21	2/4	248	
Danbury	123	1/21	2/26	157	
Derby	123	9/11	9/12	167	
Enfield	123	1/11	12/26	71 71	
Greenwich	004	1/21	1/10	200	
Groton	123	2/4	12/24	100 87	,
Hartford	123	1/11	4/13	187	
Meriden	002	2/4	1/21	159	
Middletown	003	6/16	•	67 (2 months)	
Milford	002	11/16	6/14		
New Britain	123		12/5	129	
New Haven	004	2/4			
New Haven	123	1/11	1/21	236	
Stamford	123	1/8	1/21	203	
Waterbury	123	1/7	1/21	136	

TABLE 7
SULFUR DIXOIDE
1976 - CONNECTICUT
3-HOUR CONCENTRATION

	Site		Date		0 100 200 300 400 500		009
	Bridgeport	001	2/4				
	Bridgeport	123	1/21				
	Danbury	123	5/14				
	Derby	123	1/21		187		
	Enfield	123	1/13		157		
	Greenwich	004	1/21				
- 27	Groton	123	2/25		167		
7 _	Hartford	123	1/13				
	Meriden	002	2/4		227		
	Middletown	003	7/22		100		
	Milford	002	11/15			1 1 1 1	
	New Britain	123	1/13	<u> </u>	253		
	New Haven	004	2/4				
	New Haven	123	2/4	·, · · · · · · · · · · · · · · · · · ·			
	Stamford	123	2/10			·	
	Waterbury	123	1/21		218		

Conclusions:

As in past years, Connecticut experienced very high concentrations of photochemical oxidants (measured as ozone) in the summer months of 1976. At each of the twelve monitoring sites, levels in excess of the NAAQS of 0.08 ppm were frequently recorded, with one-hour average concentrations occasionally exceeding 0.2

· 中心、翻 跨海中 轉列機區 医自己性的性病性的遗憾的影響

Discussion of Data:

Tables 8 & 9, which are summary comparisons of 1976, 1975, and 1974 ozone data, indicate no discernible downward trend in ambient oxidant concentrations. In fact, at many sites, 1976 second highs and frequencies exceeded those in 1975.

In order to gather information which will further the understanding of production and transport, as well as to provide realtime data for the daily Pollutant Standards Index, DEP operated in 1976 a state-wide ozone monitoring network consisting of four types of sites:

Urban: Bridgeport, Derby, Hartford, Middletown, New Haven

Advection from southwest: Danbury, Greenwich

Suburban: Enfield, Groton

Eastford, Hamden, Morris Rural:

Method of Measurement:

The Air Compliance Unit uses chemiluminescent instruments to measure levels of ozone which is the major constituent of photo-chemical oxidants in this area. These instruments measure and record instantaneous concentrations of ozone continuously by means of a fluorescent technique. Properly calibrated, these instruments have been shown to be remarkably reliable and stable.

公共企業法 网络中亚 法不允许的 法基础编制 医拉尔伯诺利氏神经原物

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电 超升级学

TABLE 8
APRIL TO SEPTEMBER OZONE SUMMARY THREE YEAR TREND

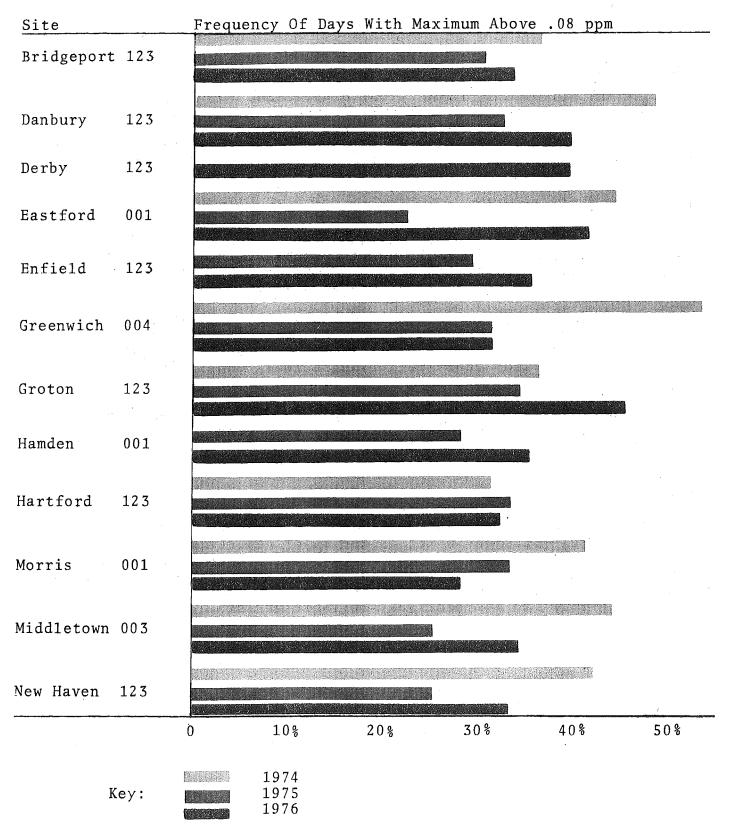


TABLE 9 OZONE THREE YEAR TREND

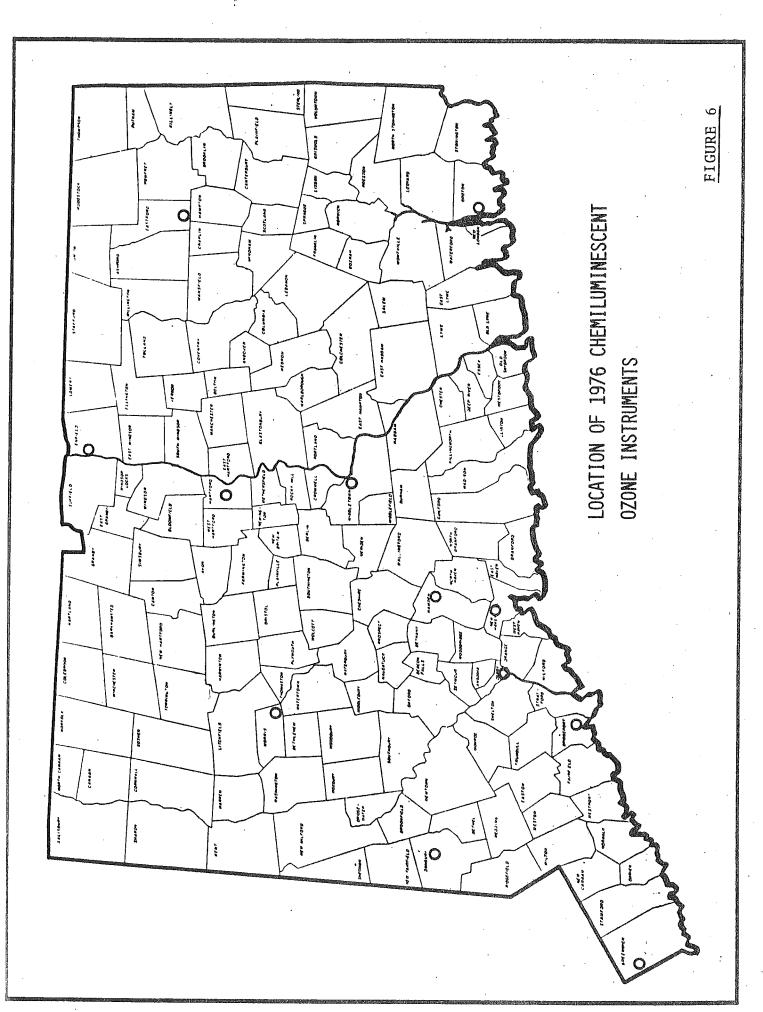
						Tabu	lation
			Gra	aph of			Second
<u>Site</u>		Second	High One	Hour Conc	entration	$\frac{\text{High}}{.250}$	High .230
Duidenant	0.01	2.40 M 10 M				.285	.250
Bridgeport	001	PERSONAL PROPERTY.		Bar and a second		.231	.228
						.270	.255
Danbury	123					. 225	.200
·						.241	.233
D1	127					.290	.280
Derby	123			<u></u>		. 250	. 200
			-			.197	193
Eastford	001					.253	.223
						.260	.230
						210	200
Enfield	123					.210 .241	.200 .218
	•		Ţ	-		. 441	. 210
						.240	.240
Greenwich	004					.185	.180
0.00						.250	.240
						244	0.00
					×	. 244	.228
Groton	123			en were transfer or a		.220 .235	.210
			•			. 233	. 444
1	001					.290	.270
Hamden	001			113		.240	. 240
		100 MICE 1000 TO 100 TO					
						.306	.221
Hartford	123					.200 .242	.190 .214
				-Primary	Standard	. 444	· 4.14
				-Pilmary	Stanuaru	.225	.174
Morris	001				•	.185	.155
				1		.195	.195
							~ A
	00#	(<u>1</u> 535)				. 365	. 324
Midd1etown	003	Lagrange and				.325	. 295
						.265	.255
						.302	.287
New Haven	123					.315	.310
				(1)		.274	.235
		0	.1ppm	.2ppm	.3ppm		
		PROGRAMMENT PROGRAMMENT AND ADMINISTRATION OF THE PROGRAMMENT AND ADMI					
	**		1974				
	Key:		1975				
		1.14	1976				

Table 10 1976 Ozone

		Apri1	Number May	of days June	with July	1-hour >.0 August	8 ppm Sept	Total
Bridgeport	123	5	3	11	17	15	9	60
Danbury	123	6	10	16	13	11	2	58
Derby	123	6 .	7	17	19	17	7 7	73
Eastford	001	9	9	15	17	16	9	75
Enfield	123	7	11	14	9	15	7	63
Greenwich	004	5	4	13	17	11	6	56
Groton	123	9	11	14	21	· · · · 17	10	82
Hamden	001	7	8	18	16	2	4	55
Hartford	123	5	9	13	12 ***	13	8 .	60
Morris	001	6	8	13	12	6	8	53
Middletown	003	3	7	13	13	13	9	58
New Haven	123	5	8	13	15	14	5	60

TABLE 11 1976 OZONE

	•	Maximu A	Maximum 1-hc A M	our concentration J J A	centr: J	}	(mdd)	High	Date	2nd High	Date
Bridgeport	123	.199	.123	.225	.194	.231	.220	. 231	8/12/14	. 228	8/13/14
Danbury	123	.233	.137	.206	.183	.241	.127	.241	8/13/16	.233	4/19/15
Derby	123	.210	.128	.280	.220	.290	.177	.290	8/12/15	.280	6/24/15
Eastford	001	.190	.130	.215	.230	.260	.145	. 260	8/12/18	.230	7/20/16
Enfield	123	.241	.135	.202	.215	.190	.155	.241	4/19/16	.218	4/19/15
Greenwich	004	.148	.128	.250	.176	.224	.160	.250	6/08/15	.240	6/28/14
Groton	123	.134	.110	.235	.222	.167	.180	.235	6/11/15	.222	7/20/14
Haddam	001	.217	.140	.240	.240	.120	.120	.240	6/24/16	.240	7/20/14
Hartford	123	.214	.114	.209	.242	.214	.165	.242	7/20/16	.214	4/19/15
Morris	001	.195	.125	.170	.153	.195	.145	. 195	4/19/16	.195	8/05/16
Middletown	003	.125	.117	.255	.240	.265	.138	.265	8/12/16	.255	6/24/16
New Haven	123	.159	.107	.274	.202	.235	.143	.274	6/24/15	.235	8/12/15
Date is read	ad as:		month/day/	hour of		occurrence					



V. NITROGEN DIOXIDE

Conclusions:

Nitrogen dioxide levels at all sampling sites in Connecticut were lower than the National Ambient Air Quality Standard of 100 $\mu\text{g/m}^3$, annual arithmetic mean.

Discussion of Data:

There were 43 nitrogen dioxide sites in 1976 as compared to 41 in 1975. The sites are distributed in a network which covers urban, residential and suburban locations.

The nitrogen dioxide data is presented in Table 12. The format is the same used to list the total suspended particulate data. Note that although the distribution of NO2 data is lognormal, the annual arithmetic mean is shown for direct comparison to the NAAQS for nitrogen dioxide. The 95 percent limits and standard deviation are also arithmetic calculations, but the geometric means and standard deviations were used to give accurate predictions of the number of days the levels of 100 $\mu g/m^3$ and $282~\mu g/m^3$ would be exceeded at each site if sampling had been conducted on a daily basis. Although there is no 24-hour NAAQS for NO2, the 282 $\mu g/m^3$ level was selected because at this level a 2nd stage air pollution alert is to be declared according to the State of Connecticut's Administrative Regulations for the Abatement of Air Pollution, while the 100 $\mu g/m^3$ level was selected as an indication of how many days per year the annual NAAQS may have been exceeded if sampling was performed daily.

Sample Collection and Analysis:

The Air Monitoring Unit uses gas bubblers employing the NASN Sodium Arsenite method. These instruments sample for twenty-four hours every sixth day, the same schedule as the suspended particulate instruments. The samples are later chemically analyzed in the laboratory.

CONNECTICUT DEPARTMENT	PARTM	0 H	ENVIRONMENTAL	. PRCTECTION	MOIJ	PAGE	1	AIR	COMPL IANCE	MONITORING
POLLUTANTNITROGEN	ROGEN	DICXIDE	111					[DISTRIBUTIONLOGNORMAL	-LOGNORMAL
TCWN NAME	SITE	YEAR	SAMPLES	MEAN	95-PCT- LOWER	-LIMITS UPPER	STD DEV	DEVIATION	PREDICTED DAYS OVER 100 UG/M3	PREDICTED DAYS OVER 282 UG/M3
BERLIN	01	1976	13	49.7	35	49	24.	24.304	24	-
BRIDGEPORT	01	1976	57	69.1	19	7.7	31.	.261	1	7
BRIDGEPORT	03	1976	£.	65.8	. 12	75	20.	20.216	5	
BRIDGEPORT	123	1976	59	70.3	63	7.7	29	29°970	19	
BRISTOL	01	1976	52	53.1	45	19	31	31,396	53	
BRISTOL	04	1976	14	prod 0 rood prod	7	16	7	7.770		
BURLINGTON	Ŏ	1976	6	9 • 6	m	11	ထိ	8.832		
COLCHESTER	Ö	1976	10	33.6	22	45	16.	16.076	m	,
DANBURY	r 2	1976	57	41.1	35	14	23	155	24	
DERBY	123	1976	56	52.1	41	64	46.	46.595	24	
EAST HARTFCRD	ō	1976	13	40.6	26	26	25	25.370	50	≠4
EAST HARTFORD	02	1976	ស	41.2	34	48	27.	27.462	50	
EAST WINDSOR	01	1976	13	60.2	77	76	26	26.740	29	
ENFIELD	123	1976	9	44.6	40	50	21.	365	P	
GREENWICH	01	1976	54	K. 60	64	82	36	327	1	2
GREENWICH	04	1976	57	53.9	48	09	26.	323	រោ កា	
GREENWICH	08	1976	54	35.9	32	40	5	15,819	~	

CONNECTICUT	DEPARTMENT	MENT OF	ENVIRONMENTAL	PRCTECTION	NOI.	PAGE	2	AIR COMPLIANCE	MONITORING
POLLUTANTNITROGEN	VITROGEA	DIOXID	DE					DISTRIBUTIONLOGNORMAL	LOGNORMAL
TCWN NAME	SITE	YEAR	SAMPLES	MEAN	95-PCT- LOWER	-LIMITS UPPER	STD DEVIATION	PREDICTED DAYS OVER N 100 UG/M3	PREDICTED DAYS OVER 282 UG/M3
GROTON	123	1976	58	41.6	38	45	14.171	*	
HARTFORD	02	1976	58	58.5	53	49	22.128	56	
HARTFORD	123	1976	29	65.6	53	72	27.472	ار ا	
LITCHFIELD	01	1976	13	38 8	30	. 48	15.094	4	
MANSFIELD	01	1976	. 11	35°1	24	747	17.433	2	
MERIDEN	05	1976	54	51°1	77	09	31.479	45	~
MIDDLETOWN	03	1976	13	61.3	47	75	23.869	33	
MILFORD	01	1976	13	56.6	37	77	33.575	67	©
NAUGATUCK	01	1976	en en	43.0	30	56	21.828	42	m
NEW BRITAIN	02	1976	16	59.6	46	73	25.908	is m	
NEW BRITAIN	123	1976	43	39.1	ታ ጠ	4	16.941	*	
NEW HAVEN	6	1976	55	61.9	61	75	28.160	®	
NEW HAVEN	123	1976	57	73.5	99	81	30.813	19	
NORWALK	05	1976	25	74.4	99	833	35.956	-	Ħ
NORWICH	01	1976	59	43.9	40	48	18,365	6 2	
OLD SAYBROOK	K 01	1976	р п рп	50°9	38	64	19.934	æ	
PUTNAM	02	1976	13	3402	22	46	19.858	7	
STAMFORD	07	1976	56	53,3	14	59	24.870	42	,

	CONNECTICUT DEPARTMENT OF ENVIRONMENTAL	EPARTM	ENT OF 1	ENVIRONMENTAL	PRCTECTION	NOI	PAGE	m	AIR COMPLIANCE	E MONITORING	
	POLLUTANTNITROGEN DICXIDE	TROGEN	DICXID	ш			u.		DISTRIBUTIO	DISTRIBUTIONLOGNORMAL	
	TOWN NAME	SITE	YEAR	SAMPLES	MEAN	95-PCT-LIMITS LOWER UPPER	LIMITS	STD DEVIATION	PREDICTED DAYS OVER IN 100 UG/M3	PREDICTED DAYS OVER 282 UG/M3	
	STAMFORD	123	1976	56	62.6	57	99	23 = 375	45		
	STRATFORD	02	1976	58	69.1	62	76	.27°554	ις Θ		
	TORRINGTON	123	1976	57	47.7	43	52	18,254	in.		
	VOLUNTOEN	Ö	1976	12	22°8	_	28	8 8 99		•	
	WATERBURY	05	1976	E	57.7	77	72	23.545	24		
7 (WATERBURY	03	1976	(**) pred	61.4	37	98	41 ,355	0	gazal)	
3	WATERBURY	123	1976	09	65.6	09	pared 	24 352	S		
	WILLIMANTIC	0	1976	01	618	30	53	16,208	œ		

VI. CARBON MONOXIDE

Conclusions:

The eight-hour NAAQS was exceeded in five urban centers in Connecticut (New Britain 002, New Haven 007, Norwalk 005, Stamford 123 and Waterbury 004) but no violations of the one-hour standard were recorded in 1976.

In general, levels recorded in 1976 were slightly lower than in 1975.

Discussion of Data:

The network in 1976 consisted of 12 carbon monoxide monitors, one less than in 1975. A review of the network, as part of the annual State Implementation Plan review, has resulted in several changes: a new site in Waterbury (004), replacing the former one (Waterbury 123); the elimination of sites Bridgeport 123, Groton 123 and Hartford 123, none of which were properly placed to record violations of the standards. Also underway is a relocation of the Stamford CO site. Improvements were made in the sampling probe exposure at Bridgeport 004, New Britain 002, and Hartford 009.

CO levels from the past several years were examined to determine whether the new car CO emission standards had resulted in a downward trend in measured levels, but no significant changes were evident.

Method of Measurement:

The DEP Air Monitoring Unit uses instruments employing non-dispersive infra red techniques to continuously measure carbon monoxide levels. The instantaneous concentrations are recorded on strip charts from which hourly averages are extracted. The instruments are fairly insensitive to sampling line length, but concentrations vary dramatically with inlet exposure and proximity to traffic lanes.

CARBON MONOXIDE ANNUAL SUMMARY

Site		Maximum 8-Hour Average	Timelof Maximum 8-Hour	2nd High 8-Hour Average	Time ¹ of 2nd High 8-Hour	Maximum 1-Hour Average	Time ² of Maximum 1-Hour	2nd High 1-Hour Average	Time ² of 2nd High 1-Hour
Bridgeport	004	10.3	9/2/16	9.6	12/20/10	20.0	12/20/09	16.2	12/15/19
Bridgeport	123*	7.3	1/8/01	6.7	2/17/07	15.0	1/7/23	13.0	1/7/08
Greenwich	001	10.6	12/15/23	9.3	1/26/08	19.0	12/15/20	16.0	9/13/08
Groton	123*	9.4	3/9/19	7.6	2/11/24	12.5	2/28/16	11.2	2/11/17
Hartford	600	7.9	7/23/16	7.9	9/14/13	15:7	12/7/09	12.0	12/6/09
Hartford	123*	10.4	1/13/17	6.9	1/16/19	22.5	1/13/17	21.5	1/16/17
New Britain	005*	15.0	1/13/21	13.4	1/13/13	21.5	1/13/09	21.0	1/13/17
New Haven	007	10.6	12/2/16	10.2	1/21/13	22.5	12/2/12	16.5	9/14/09
Norwalk	005	11.3	1/7/16	10.8	4/17/03	22.0	1/7/11	21.0	2/25/14
Stamford	123	12.1	8/12/09	11.2	8/12/17	19.6	1/24/24	19.0	12/15/19
Waterbury	004*	13.7	11/4/20	12.5	9/17/17	27.5	11/4/13	25.0	11/4/18
Waterbury	123*	8.4	1/26/22	6.4	1/7/14	17.6	1/7/09	14.0	1/26/18
*Dartiel Veer	رم د			_					

*Partial Year

month/day/hour (EST) specifying the end of ¹Time of 8-hour averages is reported as follows: the 8-hour average period.

month/day/hour (EST) specifying the end of $^2\mathrm{Time}$ of 1-hour averages is reported as follows: the 1-hour average period.

Note: All concentrations are in ppm.

		,									
20.0		19.0 1.3	i 1 1 t	15.7	F 1	1 I	22.5	, .	• •	22.0	1 . 4
8.5	i i i i	13.0 2.5	1 	8.5	. 1 1	i i	12.5	20.0	16.1 1.8	27.5	1 I 1 I
11.0		9.5	1 1 1 1	9.5	! i ! !	1 1	11.0	17.5	8.1	16.5	1 i 1 1
15.0	i i	16.0	i i	10.5	; i	! !	16.5	13.0 3.4	10.4	20.0	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
12.0 4.9	1 1	7.0	1 1 1 1	9.9	7.0	8.0	14.0	10.5	12.8	8.5	1 1
 	1 1, 1 1,	6.5	1 1	10.0	6.5	15.5	11.5	10.0	10.6	! ! !	3.7
9.5	; f	10.0	t I	10.0	5.0	14.5	10.5	9.5	8.1	i i i i	4.5
6.5	1 1 1 1	7.5	1 1 1 1	9.0	8.9	13.5	7.5	10.0	7.0	1 1	4.6
12.5]	15.0	9.1	1.8	11.0	13.5	14.5	15.5	11.0	l 1 l 1 .	8.0
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(All values are ppm)
Max is maximum 1-hour average
Avg is monthly average

VII. SPECIAL MONITORING STUDIES

In an effort to improve monitoring techniques and gain a better understanding of the ambient air quality in Connecticut, the Air Monitoring Unit conducted several special studies in 1976. Although a majority of these studies pertained to the air pollutants for which EPA has established national ambient standards, some work has been done concerning other recognized pollutants. The following sections will discuss these special studies.

ASBESTOS MONITORING

An ambient air asbestos survey was conducted during late 1975 and early 1976 to define the magnitude of the health hazard posed by airborne asbestos fibers in Connecticut to provide information for use in the establishment of an ambient air quality standard for asbestos at some future time. A newly developed low volume particulate sampler, which operates continuously for 30-days at an air sampling flow rate of 4 cfm, equipped with special membrane filters was used to collect ambient TSP samples for subsequent chrysotile asbestos electron microscopic determination by the Batelle-Columbus Laboratories.

Approximately 30 monitoring sites were selected; locations included "typical" urban sites removed from known sources of asbestos emissions, rural-background sites and stations contiguous to 4 industrial users of asbestos (i.e., manufacturers of friction products, insulated wire and cable, ammunition and molding compounds) and 3 toll plazas situated at various points along Interstate-95. Ambient chrysotile asbestos levels removed from asbestos emission sources in both urban and rural locations were below 10 $\eta g/m^3$. However, asbestos concentrations above 30 $\eta g/m^3$ were measured near each of the industrial users of asbestos. Furthermore, asbestos levels adjacent to the toll plazas were also elevated (in the 10 $\eta g/m^3$ to 25 $\eta g/m^3$ range), implicating asbestos emissions from vehicle brake lining decomposition as a significant source of airborne asbestos fibers.

TRANSPORT OF PARTICULATE MATTER

Several types of statistical techniques were used to ascertain whether Total Suspended Particulate (TSP) concentrations measured during 1971-1975 at some 80 sites distributed throughout Connecticut demonstrated higher values on days when the general air mass flow was from the New York City Area (NY-Complex) to Connecticut (i.e., southwesterly wind flow). Over 80% of Connecticut's monitoring sites exhibited significantly elevated TSP concentrations during southwesterly wind flow. Only 2 (<5%) locations failed to demonstrate a SW directional dependency according to any of the 3 statistical procedures used. Mean TSP concentrations were 15% to almost 200% higher during periods of persistent southwesterly wind flow as compared to days when the prevailing winds were from other directions. On the other hand, both northeasterly and northwesterly wind flow were associated with lower TSP levels for approximately one-third of the monitoring sites investigated.

On a short-term basis, the average impact of particulate transport during persistent SW wind flow varied from 13 to 60 $\mu g/m^3$, generally decreasing with distance from the NY-Complex. On a percentage basis, the short-term "transport effect" varied from 14 to 69% of "typical" observed daily TSP levels. The "transport effect" demonstrated a statistically significant inverse negative correlation with radial distance from the NY-Complex which further strengthens the transport hypothesis. On a long-term basis, the average impact of particulate transport during SW wind flow varied from 1 to 11 $\mu g/m^3$, again decreasing with increasing distance from the NY-Complex. On a percentage basis, the long-term "transport effect" varied from 1.5 to 17.7% of long-term (e.g., 1-year, 5-year) mean TSP levels.

It is strongly recommended that some sort of regional TSP control strategy be formulated to reduce the transport of suspended particulates into Connecticut.

SULFUR DIOXIDE BUBBLER MONITORING

A special study to improve the sample collection efficiency of the modified West-Gaeke bubbler method during the summer months was begun in 1976. Previously, it was observed that sample solutions containing SO2 decayed significantly at temperatures over 90°F (up to 40% loss of collected SO_2 when the solution was exposed to 105°F for 24 hours).

The approach taken was to put the SO2 bubbler unit in a small refrigerator where the solution temperature could be controlled at around 60°F. There were several problems encountered using this method, the primary concern being excessive condensation in the intake tubing during warm humid days. Once condensation forms the water acts as a scrubber absorbing whatever SO2 is present in the sample air. Condensation was greatly reduced by insulating the intake tubing and using a two-timer system, which turns on the refrigerator only after sampling has begun so as not to pre-cool the insulation (which would render it ineffective). This two-timer system also allows the refrigerator to remain on until the samples are retrieved by field technicians.

Analysis of the few samples taken before the cool weather set in last fall showed up to 60% improvement in the collection efficiency of the bubbler while sampling on warm sunny days. However, until further testing is done on this refrigerated system this summer, SO2 bubbler data collected during the summer months will be suspect due to possible thermal degradation.

Recent improvements by several instrument manufacturers employing thermoelectric cooling directly to the bubbler tubes look promising and will be investigated during the coming year.

HISTORICAL TOTAL SUSPENDED PARTICULATE DATA

A tabulation of total suspended particulate data from 1957 through 1976 has been prepared and may be purchased, at cost, from the Department of Environmental Protection.

PASSIVE SAMPLING ERROR

Some preliminary investigations by Connecticut DEP staff members and others had indicated that substantial particulate matter would be collected on a high volume sampler filter while the sampling motor was not operating (passive).

A standard high volume sampler filter is typically mounted a few days before the sampling day (high volume samples run every sixth day) and retrieved a few days after. The average field time for our filters is six days. In 1976, fourteen passive samples of seven days duration were analyzed. The average weight increase was .0157 grams.

Had these samples actually run as high volume (hi-vol) filters sometime during the seven day period with flows of 50 cfm for 24 hours (2038.8 m³/day) this weight would have added 7.7 $\mu g/m^3$ to the reported hi-vol measurement (the maximum would have been 15.2 $\mu g/m^3$). The measured hi-vol average for samples from a nearby site for the same time period was 55 $\mu g/m^3$, implying the passive sampling error results in an over estimate of 10 to 20%.

Inverted filters, placed along side the upright passive samples recorded very small weight gains. Microscopic evaluation of the filters indicated most of the particles on the inverted filters were very small. The staff has concluded, therefore, that the most important mechanism for passive particulate matter collection is settling of particulates.

Since neither settlable nor non-sampling-period matter should be included in a total suspended particulate matter sample, it appears that most hi-vol measurements have a positive bias. Corrective action requires either electronically operated covers or more frequent visits to the site by field personnel, neither of which is possible with current resource limitations. While the implications are being further considered, the existence of this error is being accounted for in Air Quality Maintenance Plan development.

LOW VOLUME TSP MEASUREMENTS

As discussed in the 1975 Air Quality Summary, a continuous 30-day low volume (i.e., lo-vol) sampler was developed and field tested. This low volume sampler, which is enclosed in a shelter similar to a hi-vol and uses the same glass fiber filter paper, operates at an air sampling flow rate approximately one-tenth that used by a standard hi-vol (i.e., 4 cfm as opposed to 40 to 60 cfm). The air flow through the lo-vol is measured by a temperature compensating dry gas meter. The lo-vol measurement is essentially an arithmetic average for the entire 30-day sampling interval.

In April, 1976 hi-vol samplers at eight monitoring locations were replaced with lo-vols to provide information concerning background TSP concentrations. The monthly lo-vol TSP values were adjusted to daily sample period geometric means.

The following table shows the monthly lo-vol sample concentrations and the adjusted annual geometric means. These geometric means can be directly compared to those obtained in previous years by hi-vol sampling at the respective sites.

Since the lo-vol sampler is not a reference or equivalent method, these data were not included in the routine discussion of the states TSP air quality.

Dates	Berlin	Burlington	Kent	Morris	Mansfield	Putnam	Voluntown	 Willimantic
April May June July August September October November December	X 38 43 32 42 29 26 28 31	32 39 27 59 44 21 25 X 9	X 27 34 X 43 25 18 25 20	61 43 50 31 35 26 20 22	47 51 52 41 40 36 31 45	75 42 48 35 35 43 38 46 51	47 67 61 37 34 28 15 21	52 40 49 40 48 50 48 51 48
Average	32	23	25	29	41	43	28	47

Note: Units in $\mu g/m^3$

CHEMICAL COMPOSITION OF TOTAL SUSPENDED PARTICULATE MATTER SAMPLES

Annual averages of sixteen components or characteristics of the particulate matter collected at each sampling location have been computed for the years 1971 through 1975 and are tabulated in Table 15. The abbreviations used in the table are defined below. All values shown are annual arithmetic means in micrograms per cubic meter except for pH, which is a non-dimensional measure of acidity.

BE - Berylium
CD - Cadmium
CR - Chromium
CU - Copper
FE - Iron
PB - Lead
MN - Manganese

MN - Manganese NI - Nickel ZN - Zinc V - Vanadium

NO4 - Total Nitrates

S04 - Total Sulfates

NH4 - Ammonium NA - Sodium pH - Acidity

BENZ - Total Benzene Soluables

CHEMICAL COMPOSITION OF TOTAL SUSPENDED PARTICULATE MATTER SAMPLES

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PUBLICATIONS

The following is a partial listing of technical papers and study reports dealing with various aspects of Connecticut air pollutant levels and air quality data.

- 1. "Particulate Matter Transport", Bruckman, December, 1976.
- 2. "An Investigation of Long-Range Transport of Ozone Across the Midwestern and Northeastern U.S.", Wolff, Lioy, Wight, Meyers, Cederwall, December, 1976.
- 3. "Transport of Ozone Associated with an Air Mass", Wolff, Lioy, Wight, Meyers, Cederwall, March, 1977.
- 4. "High Volume Sampling: Errors Incurred During Passive Sampling Exposure Periods", Bruckman, Rubino, September, 1976.
- 5. "Asbestos and Mesothelioma Incidence in Connecticut", Bruckman, Rubino, Christine, February, 1977.
- 6. "A Comparison of Low Volume and High Volume Particulate Samppling", Bruckman, Hyne, Keever, 1976.
- 7. "Data Validation and Monitoring Site Review", (part of the Air Quality Maintenance Planning Process), June 15, 1976.
- 8. "Air Quality Data Analysis", (part of the Air Quality Maintenance Planning Process), August 16, 1976.
- 9. "Monitored Asbestos Concentrations in Connecticut", Bruckman, Rubino, March, 1977.

VIII. CLIMATOLOGICAL DATA

Meteorology is often the most significant factor influencing short-term changes in air quality and also has an effect on longterm trends.

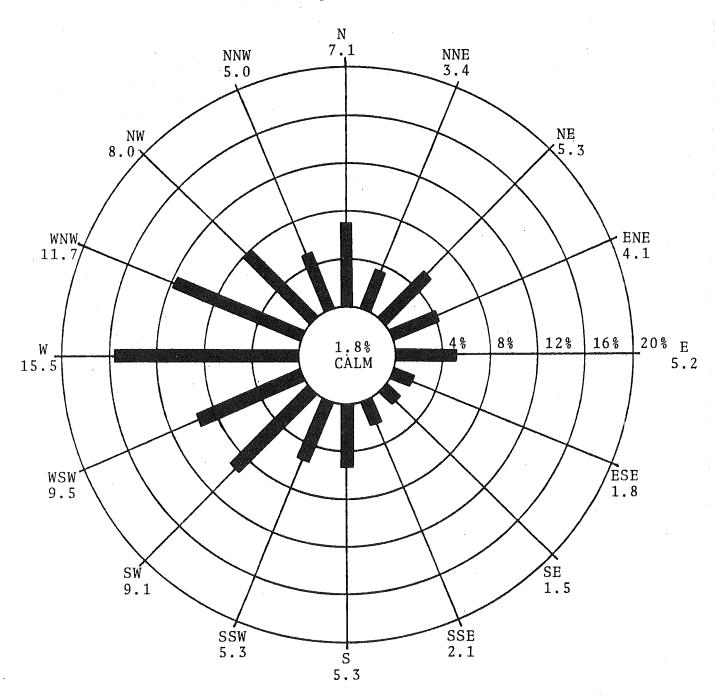
For example, motor vehicle emission control programs have a long-term effect on photochemical oxidant and carbon monoxide levels, but large year-to-year changes are more likely attributable to meteorological differences than to emission changes.

Windroses for Bradley Airport near Hartford and Sikorsky Airport in Stratford have been developed from 1976 National Weather Service surface observations and are shown in the following figures.

The Bradley Airport Windrose for 1976 shows some departures from normal: southerly winds occurred 18.8% of the time as compared to a normal of 15.4%; northerly 10.6% compared to normal 14.8%; northwesterly 12.5% compared to normal 9.7%.

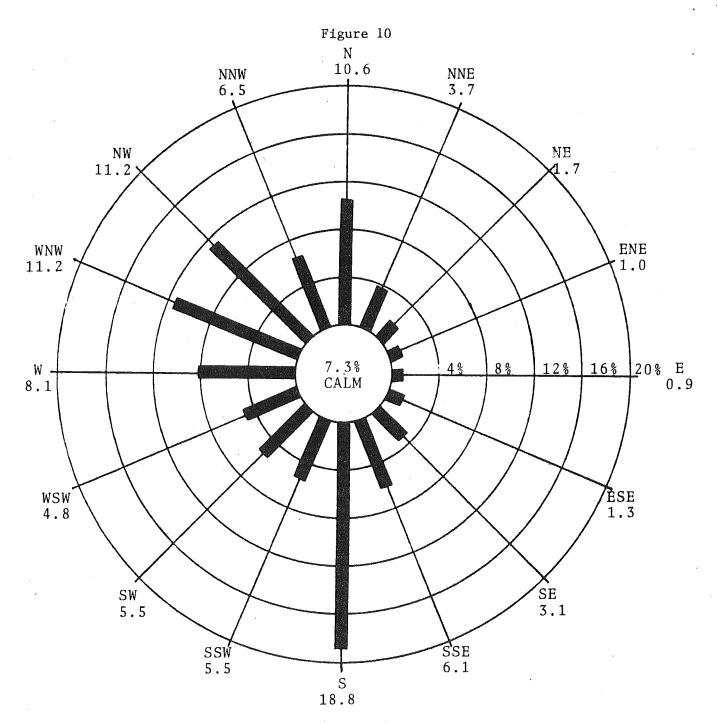
The Sikorsky Airport Windrose for 1976 also exhibits some abnormalities: westerly winds occurred 15.5% of the time in 1976 as compared to a normal of 9.8%; southwesterly winds occurred 9.1% of the time compared to a normal of 7.9%, west northwesterly 11.7% compared to 9.1% normally.





WIND ROSE SIKORSKY MEMORIAL AIRPORT ANNUAL 1976

WIND FREQUENCY APPEARS BENEATH EACH DIRECTIONAL ABBREVIATION (Calms are distributed over all directions)



WIND ROSE

BRADLEY INTERNATIONAL AIRPORT

ANNUAL 1976

WIND FREQUENCY APPEARS BENEATH EACH DIRECTIONAL ABBREVIATION

(Calms are distributed over all directions)