

DEPARTMENT OF ENVIRONMENTAL REGULATION

ST. JOHNS RIVER DISTRICT
3319 MAGUIRE BOULEVARD
SUITE 232
ORLANDO, FLORIDA 32803

PAID
AUG 12 1983



BOB GRAHAM
GOVERNOR
VICTORIA J. TSCHINKEL
SECRETARY
ALEX SENKEVICH
DISTRICT MANAGER

SAINT JOHNS RIVER DISTRICT

DER

APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCE AUG 24 1983

SOURCE TYPE: Blanket Spreading Operation New¹ Existing¹
APPLICATION TYPE: Construction Operation Modification
COMPANY NAME: DAVID "M" COMPANY COUNTY: SEMINOLE

BAQM

Identify the specific emission point source(s) addressed in this application (i.e. Line Spreaders #1, #2, #3)
Kiln No. 4 with Venturi Scrubber; Peaking Unit No. 2, Gas Fired) Pestooning and Mixing area

SOURCE LOCATION: Street 201 Valentine Way City Longwood
UTM: East 17-470106 North 3177160
Latitude 28° 43' 26" N Longitude 81° 18' 18" W

APPLICANT NAME AND TITLE: Wayne L. Brady, General Manager
APPLICANT ADDRESS: 201 Valentine Way, Longwood, Florida 32750

SECTION I: STATEMENTS BY APPLICANT AND ENGINEER

A. APPLICANT

I am the undersigned owner or authorized representative* of David "M" Company
I certify that the statements made in this application for a construction permit are true, correct and complete to the best of my knowledge and belief. Further I agree to maintain and operate the pollution control source and pollution control facilities in such a manner as to comply with the provision of Chapter 403, Florida Statutes, and all the rules and regulations of the department and revisions thereof. I also understand that a permit, if granted by the department, will be non-transferable and I will promptly notify the department upon sale or legal transfer of the permitted establishment.

*Attach letter of authorization

Signed: [Signature]
Wayne L. Brady, General Manager
Name and Title (Please Type)

Date: 8/ /83 Telephone No. 305/321-0945

B. PROFESSIONAL ENGINEER REGISTERED IN FLORIDA (where required by Chapter 471, F.S.)

This is to certify that the engineering features of this pollution control project have been designed/examined by me and found to be in conformity with modern engineering principles applicable to the treatment and disposal of pollutants characterized in the permit application. There is reasonable assurance, in my professional judgment, that

¹ See Florida Administrative Code Rule 17-2.100(57) and (104)

STATE OF FLORIDA
DEPARTMENT OF ENVIRONMENTAL REGULATION

Nº 74416

RECEIPT FOR APPLICATION FEES AND MISCELLANEOUS REVENUE

Received from Senclair & Valentine Co. Date Aug 12, 1983

Address 201 Valentine Way, Longwood Dollars \$ 1,000.00

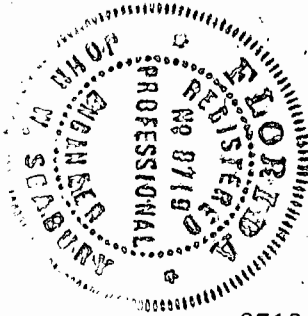
Applicant Name & Address _____

Source of Revenue David "M" Co / Blanket Spreading

Revenue Code 001001 Ch # 1588 Application Number AC59-73966

By K Tulloch

the pollution control facilities, when properly maintained and operated, will discharge an effluent that complies with all applicable statutes of the State of Florida and the rules and regulations of the department. It is also agreed that the undersigned will furnish, if authorized by the owner, the applicant a set of instructions for the proper maintenance and operation of the pollution control facilities and, if applicable, pollution sources.



Signed John W. Seabury

John W. Seabury

Name (Please Type)

Seabury-Bottorf Associates, Inc.

Company Name (Please Type)

3702 Silver Star Rd., Orlando, Florida 32808

Mailing Address (Please Type)

Florida Registration No. 8719 Date: August 11, 1983 Telephone No. 305/298-0846

SECTION II: GENERAL PROJECT INFORMATION

A. Describe the nature and extent of the project. Refer to pollution control equipment, and expected improvements in source performance as a result of installation. State whether the project will result in full compliance. Attach additional sheet if necessary.

The project will involve the installation of a lithographic printing blanker manufacturing facility and will include the following sources which are part of a common exhaust system. Spreaders #1, #2, #3, Festooning area, and Mixing area. Please refer to emission points 1-5 on the following process layout diagrams to ascertain particular emission points and exhaust ventilation configurations.

B. Schedule of project covered in this application (Construction Permit Application Only)

Start of Construction upon arrival Completion of Construction 1 month post approval

C. Costs of pollution control system(s): (Note: Show breakdown of estimated costs only for individual components/units of the project serving pollution control purposes. Information on actual costs shall be furnished with the application for operation permit.)

Not applicable

D. Indicate any previous DER permits, orders and notices associated with the emission point, including permit issuance and expiration dates.

None known

E. Requested permitted equipment operating time: hrs/day 16 ; days/wk 5 ; wks/yr 50 ;
if power plant, hrs/yr N.A. ; if seasonal, describe: _____

F. If this is a new source or major modification, answer the following questions.
(Yes or No)

1. Is this source in a non-attainment area for a particular pollutant? No
 - a. If yes, has "offset" been applied? No
 - b. If yes, has "Lowest Achievable Emission Rate" been applied? No
 - c. If yes, list non-attainment pollutants. _____ No
 2. Does best available control technology (BACT) apply to this source?
If yes, see Section VI. Yes
 3. Does the State "Prevention of Significant Deterioration" (PSD)
requirement apply to this source? If yes, see Sections VI and VII. No
 4. Do "Standards of Performance for New Stationary Sources" (NSPS)
apply to this source? No
 5. Do "National Emission Standards for Hazardous Air Pollutants"
(NESHAP) apply to this source? No
- H. Do "Reasonably Available Control Technology" (RACT) requirements apply
to this source? No
- a. If yes, for what pollutants? N.A.
 - b. If yes, in addition to the information required in this form,
any information requested in Rule 17-2.650 must be submitted.

Attach all supportive information related to any answer of "Yes". Attach any justifi-
cation for any answer of "No" that might be considered questionable.

F. 2. (Yes) Florida Air Pollution Regulation 17-2.500(5)(c) Application of best
available control technology to pollutants subject to NSR requirements as set forth
in 17.2.5(2)(f).

SECTION III: AIR POLLUTION SOURCES & CONTROL DEVICES (Other than Incinerators)

A. Raw Materials and Chemicals Used in your Process, if applicable:

Description	Contaminants		Utilization Rate - lbs/hr	Relate to Flow Diagram
	Type	% Wt		
Rubber	None Known	-	Variable	Used to formulate cements 2 Mixing (2), Spreading (3, 4, 5), Festooning (6) (At step #6 99%+ of V.O.C. has been flashed off)
Rubber Cement	V.O.C.	30-70%	Variable	
	MEK			
	Toluene			

*The process utilizes rubber cements which are mixtures of various rubber compounds i.e. (nitrile, etc) and hydrocarbon diluents such as toluene or MEK. VOC content of cement may vary from 30 to 70% of

B. Process Rate, if applicable: (See Section V, Item 1) coating weight depending upon type.

- Total Process Input Rate (lbs/hr): Variable
- Product Weight (lbs/hr): Variable

C. Airborne Contaminants Emitted: (Information in this table must be submitted for each emission point, use additional sheets as necessary)

Name of Contaminant	Emission ¹		Allowed Emission Rate per Rule 17-2	Allowable ³ Emission lbs/hr	Potential ⁴ Emission		Relate to Flow Diagram
	Maximum lbs/hr	Actual T/yr			lbs/yr	T/yr	
VOC - MEK/	103	84	BACT	To be determined	420,000	210	1
" Toluene	5	4	BACT	To be determined	20,000	10	3
"	94.5	75.6	BACT	To be determined	378,000	189	4
"	94.5	75.6	BACT	To be determined	378,000	189	5

* Potential emissions are based upon a two shift (16 hour) operation, five days per week and 250 days per year. Actual emissions are based upon a one shift (8 hour) operation, five days per week, and 250 days per year.

¹See Section V, Item 2.

²Reference applicable emission standards and units (e.g. Rule 17-2.600(5)(b)2, Table II, E. (1) - 0.1 pounds per million BTU heat input)

³Calculated from operating rate and applicable standard.

⁴Emission, if source operated without control (See Section V, Item 3).

D. Control Devices: (See Section V, Item 4)

Name and Type (Model & Serial No.)	Contaminant	Efficiency	Range of Particles Size Collected (in microns) (If applicable)	Basis for Efficiency (Section V Item 2)

E. Fuels

Type (Be Specific)	Consumption*		Maximum Heat Input (MMBTU/hr)
	avg/hr	max./hr	

*Units: Natural Gas--MMCF/hr; Fuel Oil--gallons/hr; Coal, wood, refuse, other--lbs/hr.

Fuel Analysis:

Percent Sulfur: N.A. Percent Ash: N.A.
 Density: N.A. lbs/gal Typical Percent Nitrogen: N.A.
 Heat Capacity: N.A. BTU/lb N.A. BTU/gal
 Other Fuel Contaminants (which may cause air pollution): N.A.

F. If applicable, indicate the percent of fuel used for space heating.

Annual Average N.A. Maximum N.A.

G. Indicate liquid or solid wastes generated and method of disposal.

Rubber cement waste is generated by the process and consists of rubber and varying amounts of rubber mixed with VOC. The waste is packaged in 55 gallon open head drums and disposed of in a secure hazardous waste land disposal site in Emille, Alabama.

H. Emission Stack Geometry and Flow Characteristics (Provide data for each stack): (See following page)

Stack Height: _____ ft. Stack Diameter: _____ ft.

Gas Flow Rate: _____ ACFM _____ DSCFM Gas Exit Temperature: _____ °F.

Water Vapor Content: _____ % Velocity: _____ FPS

SECTION IV: INCINERATOR INFORMATION

Not Applicable

Type of Waste	Type 0 (Plastics)	Type I (Rubbish)	Type II (Refuse)	Type III (Garbage)	Type IV (Pathological)	Type V (Liq. & Gas By-prod.)	Type VI (Solid By-prod.)
Actual lb/hr Incinerated							
Uncontrolled (lb/hr)							

Description of Waste _____

Total Weight Incinerated (lb/hr) _____ Design Capacity (lb/hr) _____

Approximate Number of Hours of Operation per day _____ day/wk _____ wks/yr. _____

Manufacturer _____

Date Constructed _____ Model No. _____

	Volume (ft. ³)	Heat Release (BTU/hr)	Fuel		Temperature (°F)
			Type	BTU/hr	
Primary Chamber					
Secondary Chamber					

Stack Height: _____ ft. Stack Diameter: _____ Stack Temp. _____

Gas Flow Rate: _____ ACFM _____ DSCFM* Velocity: _____ FPS

*If 50 or more tons per day design capacity, submit the emissions rate in grains per standard cubic foot dry gas corrected to 50% excess air.

Type of pollution control devices: Cyclone Wet Scrubber Afterburner
 Other (specify) _____

H. Emission stack geometry and flow characteristics: From estimates

Emission Point $\diamond 1$ (SCE process floor diagram):

Stack Height: 18 ft. Stack Dimensions: 4.5 x 4.5 square (ft.)
Gas Flow Rate: 12,600 ACFM 9960 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content Ambient (variable)% Velocity: 600 FPS

Emission Point $\diamond 3$ (See process flow diagram):

Stack Height: 12 ft. Stack Dimensions: 1.5 x 1.5 square (ft.)
Gas Flow Rate: 1575 ACFM 1410 DSCFM Gas Exit Temp.: 95 (°F)
Water Vapor Content Ambient (variable)% Velocity: 700 FPS

Emission Point $\diamond 4$ (See process flow diagram):

Stack Height: 12.5 ft. Stack Dimensions: 2(1.33 x 1.33 outlets)(ft.)
Gas Flow Rate: 1415 ACFM 1120 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content: Ambient (variable)% Velocity: 800 FPS

Emission Point $\diamond 5$ (See process flow diagram):

Stack Height: 12.5 ft. Stack Dimensions: 2(1.33 x 1.33 outlets)(ft.)
Gas Flow Rate: 1415 ACFM 1120 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content: Ambient (variable)% Velocity: 800 FPS

Brief description of operating characteristics of control devices: Not Applicable

Ultimate disposal of any effluent other than that emitted from the stack (scrubber water, ash, etc.):

Not Applicable

NOTE: Items 2, 3, 4, 6, 7, 8, and 10 in Section V must be included where applicable.

SECTION V: SUPPLEMENTAL REQUIREMENTS

Please provide the following supplements where required for this application.

1. Total process input rate and product weight -- show derivation [Rule 17-2.100(127)]
2. To a construction application, attach basis of emission estimate (e.g., design calculations, design drawings, pertinent manufacturer's test data, etc.) and attach proposed methods (e.g., FR Part 60 Methods 1, 2, 3, 4, 5) to show proof of compliance with applicable standards. To an operation application, attach test results or methods used to show proof of compliance. Information provided when applying for an operation permit from a construction permit shall be indicative of the time at which the test was made.
NOT APPLICABLE
3. Attach basis of potential discharge (e.g., emission factor, that is, AP42 test).
SEE APPENDIX 1 ATTACHED
4. With construction permit application, include design details for all air pollution control systems (e.g., for baghouse include cloth to air ratio; for scrubber include cross-section sketch, design pressure drop, etc.)
SEE APPENDIX 1 ATTACHED
5. With construction permit application, attach derivation of control device(s) efficiency. Include test or design data. Items 2, 3 and 5 should be consistent: actual emissions = potential (1-efficiency).
NOT APPLICABLE
6. An 8 1/2" x 11" flow diagram which will, without revealing trade secrets, identify the individual operations and/or processes. Indicate where raw materials enter, where solid and liquid waste exit, where gaseous emissions and/or airborne particles are evolved and where finished products are obtained.
NOT APPLICABLE
7. An 8 1/2" x 11" plot plan showing the location of the establishment, and points of airborne emissions, in relation to the surrounding area, residences and other permanent structures and roadways (Example: Copy of relevant portion of USGS topographic map).
SEE APPENDIX 2
8. An 8 1/2" x 11" plot plan of facility showing the location of manufacturing processes and outlets for airborne emissions. Relate all flows to the flow diagram.
SEE APPENDIX 4
SEE APPENDIX 3

- 9. The appropriate application fee in accordance with Rule 17-4.05. The check should be made payable to the Department of Environmental Regulation. ATTACHED
- 10. With an application for operation permit, attach a Certificate of Completion of Construction indicating that the source was constructed as shown in the construction permit. NOT APPLICABLE

SECTION VI: BEST AVAILABLE CONTROL TECHNOLOGY

- A. Are standards of performance for new stationary sources pursuant to 40 C.F.R. Part 60 applicable to the source?
 Yes No

Contaminant	Rate or Concentration
N.A.	N.A.

- B. Has EPA declared the best available control technology for this class of sources (If yes, attach copy)
 Yes No

Contaminant	Rate or Concentration
BACT has not been declared for this class of sources.	

- C. What emission levels do you propose as best available control technology?

Contaminant	Rate or Concentration
VOLATILE ORGANIC COMPOUNDS	AVERAGE COATING VOC CONTENT OF 5.4 lbs./gal. VOC/gal. OF COATING DELIVERED TO COATING APPLICATOR.

- D. Describe the existing control and treatment technology (if any).
- | | |
|---------------------------|--------------------------|
| 1. Control Device/System: | 2. Operating Principles: |
| 3. Efficiency:* | 4. Capital Costs: |

*Explain method of determining

Brief description of operating characteristics of control devices: _____

Ultimate disposal of any effluent other than that emitted from the stack (scrubber water, ash, etc.):

NOTE: Items 2, 3, 4, 6, 7, 8, and 10 in Section V must be included where applicable.

SECTION V: SUPPLEMENTAL REQUIREMENTS

Please provide the following supplements where required for this application.

1. Total process input rate and product weight -- show derivation [Rule 17-2.100(127)]
2. To a construction application, attach basis of emission estimate (e.g., design calculations, design drawings, pertinent manufacturer's test data, etc.) and attach proposed methods (e.g., FR Part 60 Methods 1, 2, 3, 4, 5) to show proof of compliance with applicable standards. To an operation application, attach test results or methods used to show proof of compliance. Information provided when applying for an operation permit from a construction permit shall be indicative of the time at which the test was made.
3. Attach basis of potential discharge (e.g., emission factor, that is, AP42 test).
4. With construction permit application, include design details for all air pollution control systems (e.g., for baghouse include cloth to air ratio; for scrubber include cross-section sketch, design pressure drop, etc.)
5. With construction permit application, attach derivation of control device(s) efficiency. Include test or design data. Items 2, 3 and 5 should be consistent: actual emissions = potential (1-efficiency).
6. An 8 1/2" x 11" flow diagram which will, without revealing trade secrets, identify the individual operations and/or processes. Indicate where raw materials enter, where solid and liquid waste exit, where gaseous emissions and/or airborne particles are evolved and where finished products are obtained.
7. An 8 1/2" x 11" plot plan showing the location of the establishment, and points of airborne emissions, in relation to the surrounding area, residences and other permanent structures and roadways (Example: Copy of relevant portion of USGS topographic map).
8. An 8 1/2" x 11" plot plan of facility showing the location of manufacturing processes and outlets for airborne emissions. Relate all flows to the flow diagram.

5. Useful Life:

6. Operating Costs:

7. Energy:

8. Maintenance Costs:

9. Emissions:

Contaminant	Rate or Concentration

10. Stack Parameters

- a. Height: _____ ft.
- b. Diameter: _____ ft.
- c. Flow Rate: _____ ACFM
- d. Temperature: _____ °F.
- e. Velocity: _____ FPS

E. Describe the control and treatment technology available (As many types as applicable, use additional pages if necessary).

1. LOW SOLVENT TECHNOLOGY

- a. Control Device: FORMULATE COATING TO GIVEN SPECIFICATION
- b. Operating Principles: CONTROL VOC EMISSION BY FORMULATION
- c. Efficiency:¹ NOT APPLICABLE
- d. Capital Cost: ESTIMATE, \$10,000.00
- e. Useful Life: CONTINUING
- f. Operating Cost: \$20,000.00
- g. Energy:² NOT APPLICABLE
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE
- j. Applicability to manufacturing processes: APPLICABLE
- k. Ability to construct with control device, install in available space, and operate within proposed levels: ABLE TO INITIATE

2. CATALYTIC INCINERATION

- a. Control Device: CATALYTIC INCINERATOR
- b. Operating Principles: CATALYZED INCINERATION
- c. Efficiency:¹ 90% +
- d. Capital Cost: \$500,000.00
- e. Useful Life: 10-15 YEARS
- f. Operating Cost: NOT AVAILABLE AT THIS WRITING
- g. Energy:² NOT AVAILABLE FROM BUDGET
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE

¹ Explain method of determining efficiency.

² Energy to be reported in units of electrical power - KWH design rate.

- j. Applicability to manufacturing processes: APPLICABLE WITH SOME PROCESS MODIFICATIONS
- k. Ability to construct with control device, install in available space, and operate within proposed levels: NO FORSEEABLE DIFFICULTIES

3. SOLVENT RECOVERY

- a. Control Device: CARBON ADSORBER
- b. Operating Principles: PHYSICAL ADSORPTION
- c. Efficiency:¹ 85-90%
- d. Capital Cost: \$600,000.00-\$700,000.00
- e. Useful Life: 10-15 YEARS
- f. Operating Cost: NOT AVAILABLE AT THIS WRITING
- g. Energy:² NOT AVAILABLE FROM BUDGET QUOTATION
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE
- j. Applicability to manufacturing processes: APPLICABLE WITH PROCESS MODIFICATIONS
- k. Ability to construct with control device, install in available space, and operate within proposed levels: NO FORSEEABLE MAJOR OBSTACLES TO INSTALLATION

4.

- a. Control Device:
- b. Operating Principles:
- c. Efficiency:¹
- d. Capital Costs:
- e. Useful Life:
- f. Operating Cost:
- g. Energy:²
- h. Maintenance Cost:
- i. Availability of construction materials and process chemicals:
- j. Applicability to manufacturing processes:
- k. Ability to construct with control device, install in available space, and operate within proposed levels:

f. Describe the control technology selected: LOW SOLVENT TECHNOLOGY

- 1. Control Device: FORMULATE TO GIVEN SPECIFICATION
- 2. Efficiency:¹ NOT APPLICABLE
- 3. Capital Cost: \$10,000.00
- 4. Useful Life: CONTINUING
- 5. Operating Cost: \$20,000.00
- 6. Energy:² NOT APPLICABLE
- 7. Maintenance Cost: NOT APPLICABLE
- 8. Manufacturer: NOT APPLICABLE
- 9. Other locations where employed on similar processes: NONE KNOWN

a. (1) Company: NOT APPLICABLE

(2) Mailing Address:

(3) City:

(4) State:

¹Explain method of determining efficiency.

²Energy to be reported in unite of electrical power - KWH design rate.

(5) Environmental Manager:

(6) Telephone No.:

(7) Emissions:¹

Contaminant

Rate or Concentration

(8) Process Rate:¹

b. (1) Company:

(2) Mailing Address:

(3) City:

(4) State:

(5) Environmental Manager:

(6) Telephone No.:

(7) Emissions:¹

Contaminant

Rate or Concentration

(8) Process Rate:¹

10. Reason for selection and description of systems:

¹Applicant must provide this information when available. Should this information not be available, applicant must state the reason(s) why.

SECTION VII - PREVENTION OF SIGNIFICANT DETERIORATION

NOT APPLICABLE

A. Company Monitored Data

1. _____ no. sites _____ TSP _____ () SO₂ _____ Wind spd/dir

Period of Monitoring _____ / _____ / _____ to _____ / _____ / _____
month day year month day year

Other data recorded _____

Attach all data or statistical summaries to this application.

*Specify bubbler (B) or continuous (C).

2. Instrumentation, Field and Laboratory

- a. Was instrumentation EPA referenced or its equivalent? Yes No
- b. Was instrumentation calibrated in accordance with Department procedures?
 Yes No Unknown

B. Meteorological Data Used for Air Quality Modeling

- 1. _____ Year(s) of data from _____ / _____ / _____ to _____ / _____ / _____
month day year month day year
- 2. Surface data obtained from (location) _____
- 3. Upper air (mixing height) data obtained from (location) _____
- 4. Stability wind rose (STAR) data obtained from (location) _____

C. Computer Models Used

- 1. _____ Modified? If yes, attach description.
- 2. _____ Modified? If yes, attach description.
- 3. _____ Modified? If yes, attach description.
- 4. _____ Modified? If yes, attach description.

Attach copies of all final model runs showing input data, receptor locations, and principle output tables.

D. Applicant's Maximum Allowable Emission Data

Pollutant	Emission Rate
TSP	_____ grams/sec
SO ²	_____ grams/sec

E. Emission Data Used in Modeling

Attach list of emission sources. Emission data required is source name, description of point source (on NEDS point number), UTM coordinates, stack data, allowable emissions, and normal operating time.

F. Attach all other information supportive to the PSD review.

G. Discuss the social and economic impact of the selected technology versus other applicable technologies (i.e., jobs, payroll, production, taxes, energy, etc.). Include assessment of the environmental impact of the sources.

H. Attach scientific, engineering, and technical material, reports, publications, journals, and other competent relevant information describing the theory and application of the requested best available control technology.

VOLATILE ORGANIC COMPOUND
EMISSION CALCULATIONS

APPENDIX I

Maximum Potential Emissions

Spreader #1

Maximum VOC loading in lbs./min. during spreading: 4.3 lbs./min. = 258 lbs./hr

Actual spreading time: 15% of operating time (from time study)

Potential operating parameters: 16 hours/day
250 days/year
5 days/week

Use Calculation:

16 hrs./day x .15 = 2.4 hrs./day in spread mode

2.4 hrs./day x $\frac{60 \text{ min.}}{1 \text{ hr.}}$ x 4.3 lbs./min. during spreading = 619.20 lbs./day emission
(maximum potential)

619.20 lbs./day x 250 days/yr. = 154,800 lbs./yr. = 77.4 tons/yr.

Spreaders #2 and #3

Maximum VOC loading in lbs./min. during spreading: 14 lbs./min. = 840 lbs./hr

Actual spreading time: 15% of operating time (from time study)

Potential operating parameters: 16 hours/day per spreader
250 days/year per spreader
5 days/week per spreader

Use Calculation:

16 hrs./day x .15 = 2.4 hrs./day in spread mode

2.4 hrs./day x $\frac{60 \text{ min.}}{1 \text{ hr.}}$ x 14 lbs./min. = 2016 lbs./day emission-maximum potential

2016 lbs./day x 250 days/yr. = 504,000 lbs./yr. = 252 tons/yr. per spreader

2 spreaders x 252 tons/year = 504 tons/yr. for spreaders #2 and #3 - Maximum potential

Festooning Area

The festooning area routinely flashes of one pass worth of solvent per day/blanket. Four blankets would be placed into festooning each day on a two shift operation. Process records indicate that a maximum of 10 lbs. of solvent per blanket enters the festooning area. The blankets leave nearly solvent free. Therefore maximum potential solvent emissions would approximate.

4 blankets/day x 250 days/yr. x 10 lbs. solvent/blanket = 10,000 lbs./yr. = 5 tons/yr.

Mixing Area

Emissions from the mixing area can be expected to approximate 12 tons/yr. maximum potential by estimate. This would include both local and general room exhausts. 10 T/yr. can be associated with local mixing tank exhaust and 2 T/yr. can be associated with general room exhaust.

Emission Point Allocation

Emission Point 1 :

Emission point 1 captures 25% of the VOC exhaust from spreaders #2 and #3, 100% of the VOC exhaust emissions from spreader #1, 100% of the festooning area VOC exhaust and 100% of general room exhaust from the mixing area.

$$\text{Emission Point 1 Total} = (\text{spreader (\# 1 : 77.4 T/yr.)} + (\text{spreaders (2 and 3) } 504 \text{ T/yr.} \times .25) + (\text{festooning } 5 \text{ tons/yr.}) + (\text{mixing general } 2.0 \text{ T/yr.}) = \boxed{210 \text{ T/yr.}}$$

Emission Point 3 :

Emission Point 3 captures 100% of the VOC exhaust from the mixing room tanks. This emission can be expected to approximate a maximum potential emission of 10 tons/yr.

Emission Point 4 :

Emission point 4 captures 75% of spreader #2's exhaust. Therefore maximum potential emission from emission point 4 would be

$$252 \text{ T/yr.} \times .75 = \boxed{189 \text{ tons/yr.}}$$

Emission Point 5 :

Emission point 5 captures 75% of spreader #3's exhaust. Therefore maximum potential emission from emission point 5 would be

$$252 \text{ T/yr.} \times .75 = \boxed{189 \text{ tons/yr.}}$$

Maximum Lbs./Hr. Calculation

Emission Point 1 :

- a) Spreader #1 maximum spreading time/hr. = 9.0 min./hr.
Spreader #1 maximum solvent use/min. = 4.3 lbs./min.
Spreader #1 emission point collection % = 100%
 $9.0 \text{ min./hr.} \times 4.3 \text{ lbs./min} \times 1.0 = \boxed{38.7 \text{ lbs./hr.}}$
- b) Spreader #2 maximum spreading time/hrs. = 9.0 min./hr.
Spreader #2 maximum solvent use/min. = 14.0 lbs./min.
Spreader #2 emission point collection % = 25%
 $9.0 \text{ min./hr.} \times 14.0 \times .25 = \boxed{31.5 \text{ lbs./hr.}}$
- c) Spreader #3 maximum spreading time/hr. = 9.0 min./hr.
Spreader #3 maximum solvent use/min. = 14.0 lbs./min.
Spreader #3 emission point collection % = 25%
 $9.0 \text{ min./hr.} \times 14.0 \times .25 = \boxed{31.5 \text{ lbs./hr.}}$

d) Mixing general exhaust 2 T/yr. = 4000 lbs./hr. ÷ 250 days/yr. = 16
 16 lbs./day ÷ 16 hrs./day = 1 lb./hr.

TOTAL MAXIMUM LBS./HR. = a + b + c + d
 = 103 lbs./hr.

Emission Point 3 :

Total annual emission = 10 T/yr. = 20,000 lbs./yr.

20,000 lbs./yr. ÷ 250 days/yr. = 80 lbs./day ÷ 16 hrs./day = 5.0 lbs./hr.

Emission Point 4 :

Spreader #2

Maximum spreading time/hr. = 9.0 min./hr.

Spreader #3

Maximum solvent use/min. = 14.0 lbs./min.

Spreader #2

Emission point collection % = 75%

9.0 min./hr. x 14.0 lbs./min. x .75 = 94.5 lbs./hr.

Emission Point 5 :

Spreader #3 maximum spreading time/hr. = 9.0 min./hr.

Spreader #3 solvent use/min. = 14.0 lbs./min.

Spreader #3 emission point collection % = 75%

9.0 min./hr. x 14.0 lbs./min. x .75 = 94.5 lbs./hr.

Actual Tons/Yr.

The facility currently operates at approximately 40% of its emitting potential. Therefore, potential T/yr. are multiplied by .40 to obtain actual emissions.

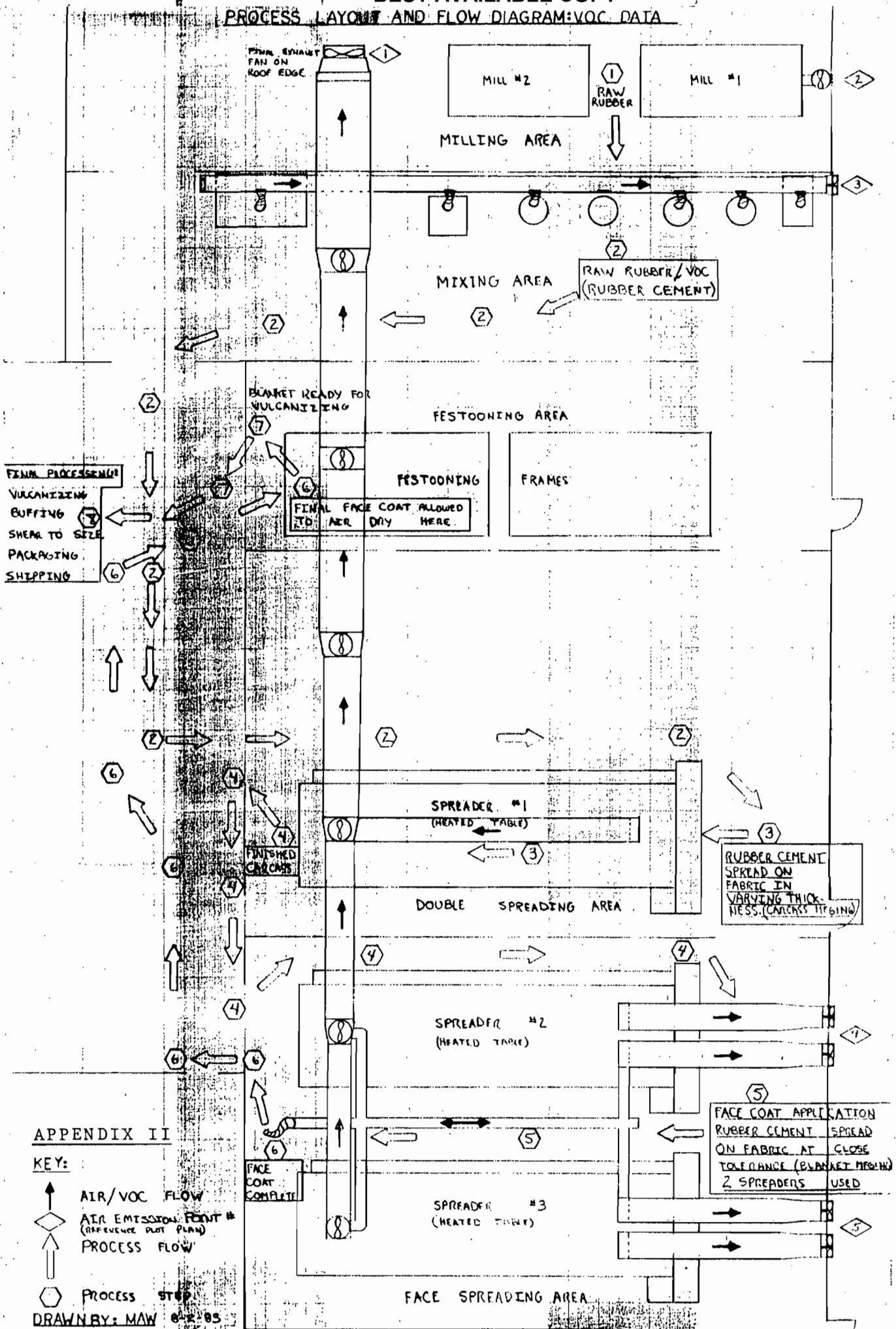
<u>Emission Point</u>	<u>Potential Emissions</u>	x	<u>Actual Emissions</u> .40	
1	210 T/yr.		84 T/yr.	
3	10 T/yr.		4 T/yr.	
4	189 T/yr.		75.6 T/yr.	
5	189 T/yr.		75.6 T/yr.	TOTAL ACTUAL TONS/YR. = 239.2

Total VOC/yr. figures correlate well with 1982/83 purchasing records for VOC raw materials. Current 1983 use projections would approximate 240 T/yr.

APPENDIX II CONT.
PROCESS FLOW EXPLANATION

- Step ① Rubber is milled into sheets for use in cement making. No calculation of emissions is included in this application since only particulate emissions would be regulated and this source is a minor one which will be dealt with separately from VOC emissions.
- Step ② Rubber is combined with VOC (MEK/Toluene) and mixed to the desired viscosity for spreading.
- Step ③ Rubber cement is spread on cotton fabric in repeated passes of varying thicknesses utilizing a knife type applicator. This step forms the blanket carcass. VOC flashes off due to steam heated coils and is captured by overhead hoods.
- Steps ④ & ⑤ Carcass is finished and moved in roll form to face spreaders 2 and 3 where top or face coats are applied to the blanket, solvent flashes off as in Step 3. The final pass is not dried, rather it is talced and recoiled with interleaving paper at which point it moves to Step 6.
- Step ⑥ The blanket is then unrolled and hung open in the festooning area where the final pass dries. The blanket is virtually solvent free after 16 hours of air drying time.
- Step ⑦ The blanket is once again rolled up and is put into a vulcanizer which is actually an autoclave which puts the final cure on the blanket.
- Step ⑧ Following final cure the blanket may be buffed or sheared and then packaged for shipping.

PROCESS LAYOUT AND FLOW DIAGRAM: VOC DATA



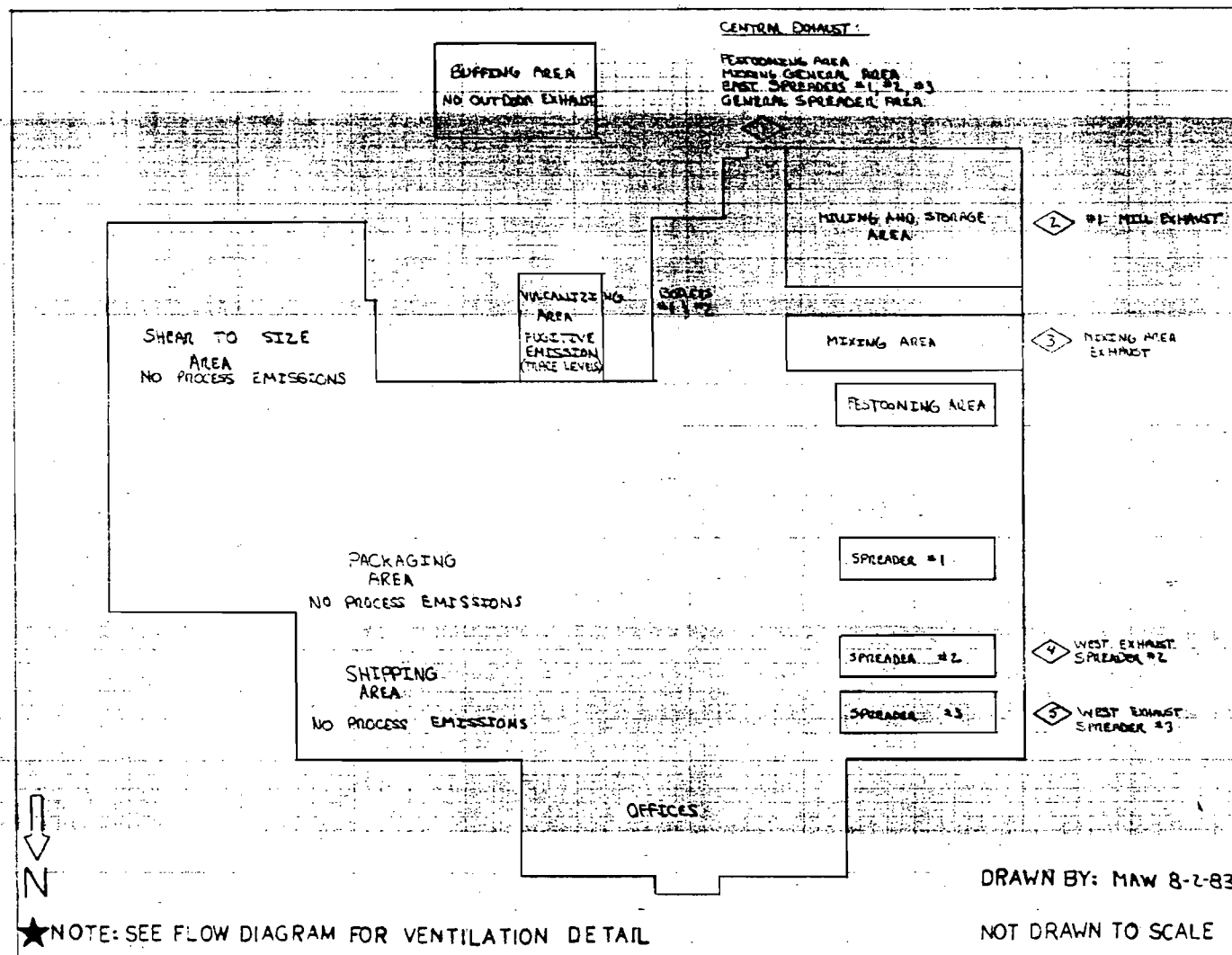
APPENDIX II

KEY:

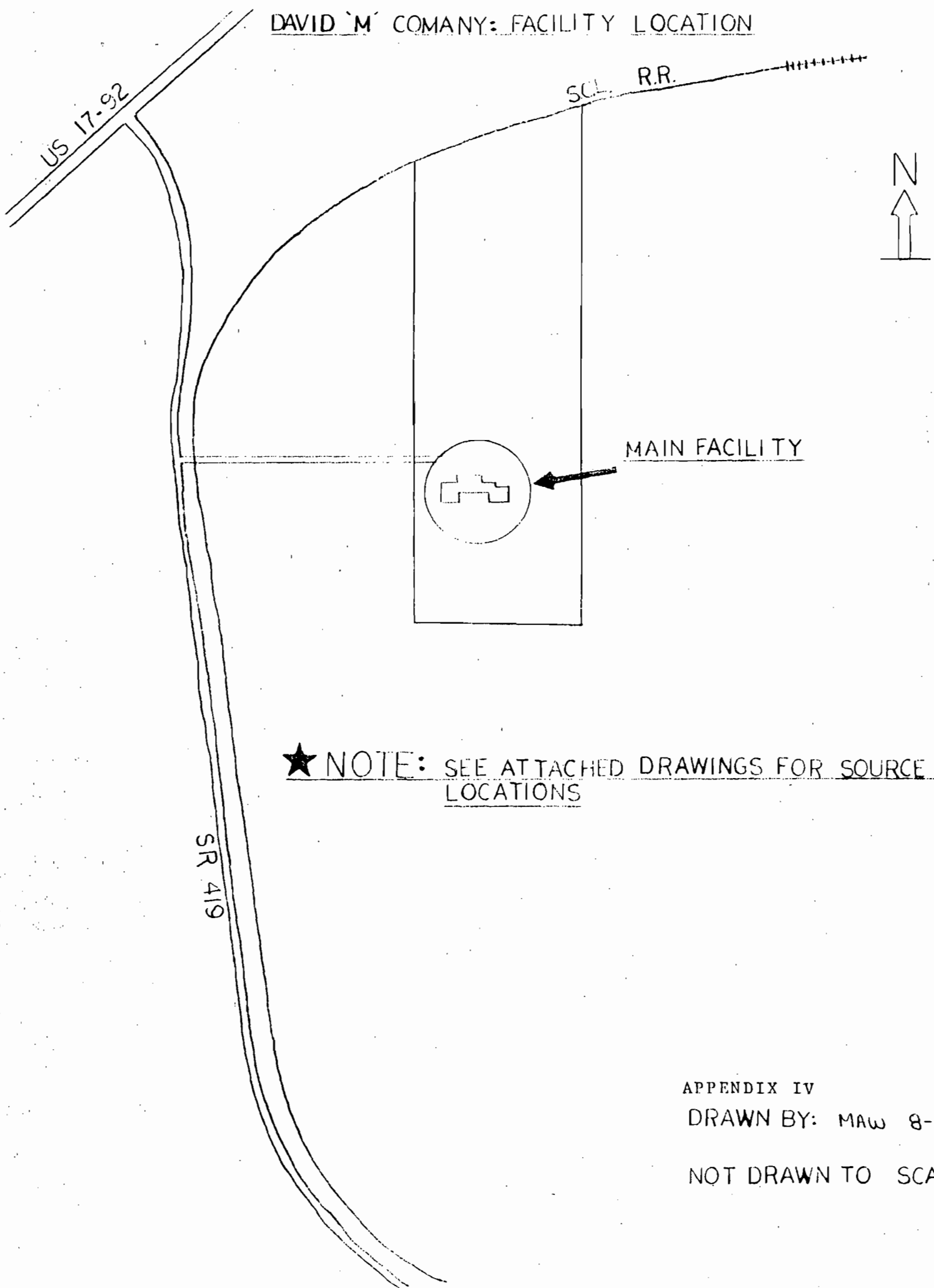
- ↑ AIR/VOC FLOW
 - ◇ AIR EMISSION POINT # (REFERENCE PLAT PLAN)
 - ↑ PROCESS FLOW
 - PROCESS STEP
- DRAWN BY: MAW 8-2-85

Best Available Copy

APPENDIX III
PROCESS LAYOUT AND EMISSION POINTS: \diamond



DAVID 'M' COMANY: FACILITY LOCATION



★ NOTE: SEE ATTACHED DRAWINGS FOR SOURCE LOCATIONS

APPENDIX IV
DRAWN BY: MAW 8-2-83
NOT DRAWN TO SCALE



DAVID "M" COMPANY

APPENDIX 1 CONTINUED
 SEABURY-BOTTORF ASSOCIATES, INC.
 CONSULTING ENGINEERS
 ORLANDO, FLORIDA

DAVID "M" COMPANY
 LONGWOOD, FLORIDA

DES	OWN J.S.	409-
-----	----------	------

DER
DEC 21 1984
BAQM

Ten UOP Plaza
Des Plaines, Illinois 60016-6195
312-391-2000

December 17, 1984

1-2-84
Copies to
Carol Furshman
& Charles Collins

Mr. Willard Hanks
Department of Environmental Regulation
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, Florida 32301-8241

Dear Mr. Hanks:

Those of us representing David M Company at the December 12, 1984 meeting feel it was quite productive. We appreciate your desire to review the implications of our changes in the permit application but also wish to reemphasize to you that the control technology does not change. We are forecasting this increased production to start in the fourth quarter 1985, and we feel it would have been counter-productive to have allowed the original permit application to go through and be followed immediately by a request for permission to install additional equipment.

In our meeting we discussed control of emissions from our organic liquid storage tanks. A letter addressed to Michael Ware dated September 27, 1984 from C. H. Fancy of the DER, directed us to investigate control methods and submit to your office by January 2, 1985 the results of our investigation rather than proceeding to install a control system. Per your instructions at our meeting on December 12, we will now take no further action on this phase of the VOC control pending direction from you after you have reviewed the revised permit application. I would like to ask you to review the calculations in that application indicating a total emission from the storage tanks of less than 150 pounds per year, and to consider whether a means of control is really necessary. It is a very small amount when compared to the other fugitive emissions.

You stated that, because of our revisions to the permit application, it will probably be the end of January 1985 before you can complete the preliminary determination and technical evaluation of this proposal, followed by a 30-day period for posting public notice of the intent to install this system. Consequently, you indicated approval of the application could be as late as March 15 of 1985. As we discussed, no allowance was made in our schedule for this delay, so it is not possible for us to meet the August 1, 1985 deadline for completion of installation. Sutcliffe Speakman estimates a minimum of five months for delivery after a purchase order is issued, plus eight weeks for installation. In our original schedule we had asked for a period from August 1, 1985 to December 1, 1985 for operational shakedown, debugging and compliance testing. Per our agreement with you on December 12 we are willing to shorten

Mr. Willard Hanks
December 17, 1984

that period. Even though the August 1 date will be no longer valid, we will make every effort to complete final compliance testing by December 1, 1985. On that basis, the estimated compliance schedule has been revised and is attached.

As we told you, an expansion of the Orlando plant is being planned and construction will closely parallel the solvent recovery installation. Since the two are adjacent, I had intended to pour the foundation slab for the solvent recovery system along with the new building footers. You indicated no advance approval can be issued for only this portion of the VOC control system. Since, however, construction of the slab will not alter our current emissions, and will result in substantial cost savings by allowing us to combine it with construction of the footings, per our original plans we intend to proceed with this construction prior to DER approval, recognizing, of course, that we do so at our own risk. We will do this unless there is an objection from the DER.

We sincerely appreciate the openness with which you received our presentation and hope for your help in expediting the approval of our permit application.

Sincerely,



R. H. Baddeley
Project Engineer

RHB/djc

cc: John Gamble
Wayne Brady
Nick Feagler
Tim Westman

October 19, 1983 - Revised December 13, 1984

ESTIMATED
OVERALL COMPLIANCE SCHEDULE
DAVID M COMPANY VOC CONTROL PROJECT

<u>PHASE</u>	<u>ORIGINAL</u> <u>DATE</u>	<u>REVISED</u> <u>DATE</u>
Data Accumulation	January 1, 1984	
Spreader Enclosure - (1) For Temperature Evaluation	January 15, 1984	
Process Modifications and Development	April 1, 1984	
Methods Comparison/Analysis - Thermal Incineration vs. Recovery	May 15, 1984	
Final Specification Development	July 1, 1984	
Final Specification/Quotations and Analysis	Spetember 1, 1984	
Appropriation Request Development and Approval	December 1, 1984	
Oven Enclosure and Exhaust System Installation	March 1, 1985	May 17, 1985
Solvent Recovery Delivery and Installation	August 1, 1985	October 18, 1985
Operational Shakedown, Debugging and Compliance Testing	December 1, 1985	December 1, 1985

STATE OF FLORIDA
DEPARTMENT OF ENVIRONMENTAL REGULATION

To be filed: DAVID M Co
cc to: _____

O&C Case No. 84-0044

MEETING DOCUMENTATION

Attendees: (Attach list) *

Location: TALLAHASSEE Date: 12/12/84 Time: 3:00 PM

Subject: Permitting Enforcement Other _____

Meeting requested by: John Gamble, Signal VOP Group, (312) 371-2440

Meeting Objectives: SUBMIT REVISED APPLICATION FOR PERMIT TO CONSTRUCT AIR POLLUTION CONTROL EQUIPMENT FOR A lithographic printing blanket manufacturing facility and new process equipment

Notes:

* Robert H. Baddeley, Prof. Eng., (312) 391-3552 (Chief Spokesman)
John H. Gamble, Director, " (Contact for questions)
HERB MYCROFT, Tech. Dir., David M. Co. (FIN. MGR.)
Ian Parry, Sutcliffe-Craftshow tech. eng. (301) 337/2800 (Control equip. rep.)

Application noted above submitted. Appears complete.

Because of schedule in consent order, Company told OK to expand building and pour cement slab for carbon absorption unit - at their risk.

Told Co. Preliminary Determination will be out in January, 1985.

Told Co. they could not increase production and emissions prior to receiving permit to construct.

Agreed that new mixing equipment won't increase production or emissions.

Explained permitting procedure/schedule.

Reviewed consent order schedule. Told Co. FINAL DATE (12/1/85)
IS DATE THEY MUST MEET OR REVISE CONSENT ORDER.

(over)



DEF
DAVID M COMPANY

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

BAQM

December 11, 1984

Mr. William Thomas
State of Florida
Department of Environmental Regulation
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-6241

Dear Mr. Thomas:

Enclosed you will find a revised construction permit application for the David 'M' Company. The application contains some significant modifications which bear some explanation. You may remember from our discussions in Tallahassee in October of 1983 that we expected a real business recovery during 1984-85. That recovery has indeed occurred and as such has necessitated that we look at additional capacity earlier than we had planned. The application evidences our proposal to add one spreader, several additional mixers and a new festooning area. Since we had not yet sized our solvent recovery system, we felt that it would be prudent to include our expansion plans in our original construction permit in order to prevent a nearly immediate request for a permit addition. You will note that very little else has changed operationally; coating VOC content, methods of application, and etc. remain appreciably the same. You will notice changes reflecting a better knowledge of the process than that which was evident at the time of our previous application.

We believe this revised permit application represents a fair and environmentally sound proposal which will serve to mutually benefit all parties involved. We sincerely hope that we have provided sufficient data for your approval process to proceed. In the event that we have not, please do not hesitate to contact me directly.

Sincerely,

Wayne L. Brady
General Manager
& Vice President

Enclosure

No. 0157029

RECEIPT FOR CERTIFIED MAIL

NO INSURANCE COVERAGE PROVIDED—
NOT FOR INTERNATIONAL MAIL
(See Reverse)

SENT TO			
Mr. Michael A. Ware			
STREET AND NO.			
P.O., STATE AND ZIP CODE			
POSTAGE	\$		
CONSULT POSTMASTER FOR FEES	CERTIFIED FEE	¢	
	SPECIAL DELIVERY	¢	
	RESTRICTED DELIVERY	¢	
	OPTIONAL SERVICES	RETURN RECEIPT SERVICE	¢
		SHOW TO WHOM AND DATE DELIVERED	¢
		SHOW TO WHOM, DATE, AND ADDRESS OF DELIVERY	¢
SHOW TO WHOM AND DATE DELIVERED WITH RESTRICTED DELIVERY		¢	
SHOW TO WHOM, DATE AND ADDRESS OF DELIVERY WITH RESTRICTED DELIVERY	¢		
TOTAL POSTAGE AND FEES	\$		
POSTMARK OR DATE			
10/1/84			

PS Form 3800, Apr. 1976

PS Form 3811, Jan. 1978

RETURN RECEIPT, REGISTERED, INSURED AND CERTIFIED MAIL

SENDER: Complete items 1, 2, and 3. Add your address in the "RETURN TO" space on reverse.

1. The following service is requested (check one.)

- Show to whom and date delivered..... ¢
- Show to whom, date and address of delivery..... ¢
- RESTRICTED DELIVERY
- Show to whom and date delivered..... ¢
- RESTRICTED DELIVERY.
- Show to whom, date, and address of delivery. \$ _____

(CONSULT POSTMASTER FOR FEES)

2. ARTICLE ADDRESSED TO:

Mr. Michael A. Ware
201 Valentine Way
Longwood, Florida 32750

3. ARTICLE DESCRIPTION:

REGISTERED NO.	CERTIFIED NO.	INSURED NO.
	0157029	

(Always obtain signature of addressee or agent)

I have received the article described above.


SIGNATURE Addressee Authorized agent

4. DATE OF DELIVERY: 10/3/84

5. ADDRESS (Complete only if requested)

6. UNABLE TO DELIVER BECAUSE:

CLERK'S INITIALS



STATE OF FLORIDA

DEPARTMENT OF ENVIRONMENTAL REGULATION

TWIN TOWERS OFFICE BUILDING
2600 BLAIR STONE ROAD
TALLAHASSEE, FLORIDA 32301-8241



BOB GRAHAM
GOVERNOR
VICTORIA J. TSCHINKEL
SECRETARY

September 27, 1984

Mr. Michael A. Ware
David M Company
201 Valentine Way
Longwood, Florida 32750

Dear Mr. Ware:

The department has reviewed your August 31, 1984, letter that summarizes the problems David M Company encountered in installing a vapor recovery system on the organic liquid storage tanks. We believe your company should investigate other means of controlling the volatile organic compound emissions from your storage tanks before selecting an air pollution control system for them. Among the systems we suggest you consider are:

1. A carbon absorption system, such as the ones used at dry cleaning establishments, or other type of controls to treat the vapors displaced when the storage tanks are filled.
2. Connecting the vent lines from the storage tanks to the air pollution control system selected for the coating production line.
3. Contact other industrial solvent distributors and gasoline carriers to see if trucks that are able to use the Coaxial Vapor Recovery System initially considered by your company are available.

There may other ways to control the emissions from these storage tanks and the department does not wish to restrict your company from investigating alternate means of control. Based on their regulations, the South Coast Air Quality Management District of El Monte, California have more experience in controlling emissions from small storage tanks than Florida and may be able to advise you of alternate control system that would be applicable to your situation.

The department does see some benefit to using the Coaxial Vapor Recovery System at your plant, even with the limitations described in your August 31, 1984, letter, but believes more

Mr. Michael A. Ware
Page Two
September 27, 1984

effective systems are available. Instead of installing the Coaxial Vapor Recovery System at this time to comply with the Consent Order No. 84-0044, the department request you use the remained of this year to investigate alternate means of control and submit a written report, summarizing the results of your investigation, to this office by January 2, 1985. We understand that the pressure/vacuum vents are being installed on the tanks on schedule.

If you have any questions on the legal matters on the Consent Order, please contact Nancy Wright at (904)488-9730. Questions on technical matters should be addressed to Willard Hanks at (904)488-1344.

Sincerely,



C. H. Fancy, P.E.
Deputy Chief
Bureau of Air Quality
Management

CHF/WH/s

cc: William Darling
Nancy Wright

9-18-84

Copies to Charles Collins and

Nancy Wright

MICHAEL A. WARE
Environmental Coordinator



Wheelabrator-Frye Inc.
Graphic Supplies Division
Chemicals and Coatings Group
2010 Indiana Street
Racine, Wisconsin 53405

11-6-84
Has
Left
Comments

Tel: (414) 554-1030



DAVID M COMPANY

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

August 31, 1984

Mr. Willard Hanks
Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-8241

File Reference: OGC Case No. 84-0044

Dear Mr. Hanks:

This correspondence shall serve as confirmation of our conversation of August 31, 1984, during the course of which we discussed evaporative emission controls for VOC Storage Tanks at our facility, required by item nine of our joint consent agreement.

We are encountering several problems in designing and implementing a storage tank Vapor Recovery System at our facility and would appreciate your assistance in this matter. The following obstacles to installation of such a system (which would displace vapors created by filling back into the tanker making the delivery) exist.

1. Standard vapor recovery equipment currently available is designed for service station use and as such is chemically resistant to gasoline only and would not be suitable for use with industrial solvents like Toluene and Methyl Ethyl Ketone. Such material incompatibilities would likely compromise the long term effectiveness of the control system. While we consider this problem to be significant, it is likely that o-rings, gaskets, valve seats and other components could be special ordered and received within 3-6 months time and therefore a system could possibly be custom made and installed for capture of VOC's at our facility. We perceive item 2 below to be a more significant problem than item 1 above.
2. We have contacted the three companies which currently sell and deliver our facility industrial solvents. None of these three companies have delivery equipment set up for vapor recovery at this time and more importantly, do not have facilities equipped to empty recovered vapors from a tanker when a facility is able to displace them back to the tanker. Our least frequent supplier informs us

RECEIVED
AUG 31 1984
M. J. HANKS

that it may be possible to route all shipments via a gasoline carrier who would be set up for evaporative exhaust emission controls, however, we are unaware of what would happen to the vapors after leaving our site since our vendors do not have provisions for vapor recovery at their facilities. Our surmial is that the vapors may find their way into the atmosphere even though much capital, time and effort has been expended for recovery activities at our location.

The aforementioned problems and questions have presented us with somewhat of a dilemma as to whether controls are appropriate and/or feasible in this case. We would sincerely appreciate your assistance in developing a proper course of action in this case. Pursuant to our recent conversation, we will not be required to comply with the 90 day installation period established by the original consent agreement for installation of Vapor Recovery equipment and we will await a reply from your office prior to proceeding with any installation of vapor recovery equipment.

In the event that you should require further information and/or assistance in resolving these questions, please feel free to contact me directly.

Sincerely,

Michael A. Ware

Michael A. Ware

cc: W. Brady
J. Spangler
H. Mycroft
B. Baddeley
T. Westman
J. Gamble

~~Class, Bill~~ 9/5
Incompleteness
response. District
& Nancy Knight
copied. Please
return for file
Patty



201 VALENTINE WAY
LONGWOOD, FLORIDA 32750



RETURN RECEIPT
REQUESTED

CERTIFIED
P 211 559 780
MAIL

Mr. Willard Hanks
Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-8241

DER
SEP 4 1984
BAQM



Wheelabrator-Frye Inc. Chemicals & Coatings Group

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

September 6, 1984

Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-8241

DER
SEP 11 1984
BAQM

Subject: Case #84-044

Dear Compliance Evaluation Specialist:

This report is being submitted in compliance with consent order (reference Case #84-044). This is our second report which has been prepared in recognition of the September 1st date referenced in our overall compliance schedule codified as Exhibit #2.

Please be advised that we have concluded our review of the alternative methods of VOC control for our Orlando facility, namely incineration and solvent recovery. Due to economic reasons, we have ruled out incineration and are now concentrating on investigations on solvent recovery systems. A set of initial parameters and specifications for a bid have been prepared and are attached as an addendum to this report for your evaluation.

These specifications have been submitted to Vara International, Inc., Vero Beach, Florida and Sutcliffe Speakman, Inc. of Lutherville, Maryland for consideration. We expect to have quotations returned to us by the end of September and will proceed to select one of these companies to be the primary supplier of the necessary pollution control equipment. Upon receipt of final quotations from the afore-mentioned suppliers, we will proceed to develop formal plans and specifications which will be forwarded you for approval as soon as they become available.

Because of the magnitude of the cost of this project, the appropriation of funds for same must be approved by our corporate management. With this consideration in mind, we plan to have specifications and estimated costs complete and the appropriation request submitted to our corporate management on or before November 5th. We will continue to apprise you of our progress as the project moves forward.

Department of Environmental Regulation

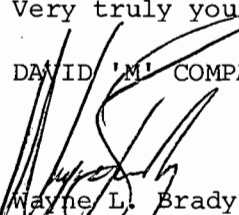
September 6, 1984

Page 2

In the event that you should have further questions and/or require further information, please do not hesitate to contact me directly.

Very truly yours,

DAVID 'M' COMPANY



Wayne L. Brady

Vice President and General Mgr.

WLB:bjw

Attach.

cc: M. A. Ware
R. H. Baddeley
J. Spangler
H. Mycroft
T. G. Westman
J. H. Gamble

July 9, 1984

CERTIFIED MAIL NO. 7236065

BEFORE THE STATE OF FLORIDA
DEPARTMENT OF ENVIRONMENTAL REGULATION

STATE OF FLORIDA DEPARTMENT OF ENVIRONMENTAL REGULATION,	:	IN THE OFFICE OF THE
	:	ST. JOHNS RIVER DISTRICT
	:	
Complainant,	:	
	:	OGC Case No. 84-0044
vs.	:	
	:	
DAVID "M" COMPANY	:	
	:	
	:	
Respondent.	:	

RECEIVED

JUL 9 1984

Dept. of Environmental Regulation
Office of General Counsel

CONSENT ORDER

This Consent Order is made and entered into between the State of Florida Department of Environmental Regulation ("Department") and David "M" Company, 201 Valentine Way, Longwood, Seminole County, Florida 32750, ("Respondent").

The Department finds and Respondent admits the following:

1. Respondent owns and operates a coating production line facility located at 201 Valentine Way, Longwood, Florida 32750. Section 27, Township 20 south, Range 30 east, Seminole County, Exhibit 1. This production line began operations in 1978 and manufactures a fabric coated material used in the off-set printing process. Coating of the material involves using volatile organic compounds (VOC's), which are sources of air pollution regulated by the Department.
2. Respondent also owns and operates two oil fired steam boilers at this same plant site used in conjunction with the fabric coating process.
3. Respondent constructed and began operation of the coating line without first obtaining a Department permit authorizing such construction or operation. This violates Florida Administrative Code Rules 17-2.210 and 17-4.03 and Section 403.087 and 403.161, Florida Statutes.

4. The Department issued a Warning Notice to the Respondent on June 3, 1983 in which the Respondent was to meet with the Department to resolve the violation of operating the two boilers without a permit. Permits were issued for the boilers, one on January 5, 1984, and the other on January 6, 1984.

5. An informal conference was held between the Respondent and the Department on July 28, 1983. In that meeting it was recommended to the Respondent that he should enter a Consent Order to address the failure to control VOC emissions from the coating line and the violations for constructing and operating the coating line and the two boilers without a permit.

6. The Respondent submitted an application for construction of the pollution source at the plant, specifically the VOC emission source, and the Department received the application on August 12, 1983.

THEREFORE, having reached a resolution of the matter, pursuant to Florida Administrative Code Rule 17-1.58, the Department and the Respondent mutually agree and it is:

ORDERED:

7. Respondent shall adopt and implement the compliance schedule proposed to the Department entitled: Estimated Overall Compliance Schedule David "M" Company VOC Control Project, dated October 19, 1983, attached and incorporated as Exhibit 2. With respect to the VOC emission violations covered in this Consent Order, compliance will be met as per the Respondent's schedule, Exhibit 2, and consistent with Department rules.

8. Respondent shall deliver a report, in writing to the Department within fifteen days of each deadline date noted in Exhibit 2. The report shall specify what steps have been made toward compliance with the applicable deadline, whether the deadline has been met, and the reasons for any delays. This report and all other reports, notices and applications required by this Order shall be mailed to:

Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, Florida 32301-8241

9. Respondent shall install Department approved VOC air pollution control devices on all VOC storage and holding tanks owned by the Respondent within ninety days of the effective date of this Consent Order. Notice shall be provided by the Respondent in writing within that ninety day period to the Department stating the type, name and manufacturer of the device(s). Any efficiency values shall be also noted. A brief sketch shall be attached showing the location of the control equipment and its connections to the tanks. Any pipe, valve, vent, exhaust, drain or other means for VOC's to be released into the ambient atmosphere shall be noted.

10. Respondent shall limit the operations at the plant site which involve the emissions of VOC's to only sixteen hours per day. This provision shall be effective as of the date of this Consent Order, and shall remain in force until replaced by the hours of operation as specified in a construction permit as issued by the Department.

11. Respondent shall have received and installed all pollution abatement and control equipment for control of VOC's on the coating line by August 1, 1985, as stated in Exhibit 2.

12. Respondent shall pay a settlement fee as agreed upon relating to matters in this Consent Order to the Department's Pollution Recovery Fund with either a money order, cashier's check or certified check in the amount of \$10,000.00. Payment shall be made within sixty (60) days of this Consent Order. The check or money order shall be mailed to the Department of Environmental Regulation, St. Johns River District, 3319 Maguire Boulevard, Suite 232, Orlando, Florida 32803.

13. Respondent shall allow authorized representatives of the Department access to the property at reasonable times for purposes of determining compliance with this Order and the rules of the Department.

14. The Department hereby expressly reserves the right to initiate appropriate legal action to prevent or prohibit the future violation of applicable statutes, or the rules promulgated thereunder.

15. Entry of this Consent Order does not relieve Respondent of the need to comply with applicable federal, state, or local laws, regulations, or ordinances. The entry of this Consent Order does not abrogate the rights of substantially affected persons who are not parties to this Order, pursuant to Chapter 120, Florida Statutes.

16. The terms and conditions set forth in the Consent Order may be enforced in a court of competent jurisdiction pursuant to Sections 120.69 and 403.121, Florida Statutes. Failure to comply with terms of this Consent Order shall constitute a violation of Section 403.161(1)(b), Florida Statutes.

17. Respondent is fully aware a violation of the terms of this Consent Order may subject Respondent to judicial imposition of damage, civil penalties of up to \$10,000 per offense, and criminal penalties.

18. This Consent Order shall take effect upon the date of filing and acknowledgement by the Clerk of the Department and shall constitute final agency action by the Department pursuant to Section 120.69, Florida Statutes and Florida Administrative Code Rule 17-1.58(3).

Best Available Copy

FOR THE RESPONDENT:

June 13, 1984
Date

Wayne Brady
WAYNE BRADY
President and General
Manager
David "M" Company
201 Valentine Way
Longwood, Florida 32750

DONE AND ORDERED this 15 day of June,
1984, in Orlando, Florida.

Galifian
A. ALEXANDER, P.E.
District Manager
State of Florida Department
of Environmental Regulation
3319 Maguire Boulevard
Suite 232
Orlando, Florida 32803
Telephone: (305) 894-7555

FILING AND ACKNOWLEDGEMENT

FILED, on this date, pursuant to §120.52 (9),
Florida Statutes, with the designated Depart-
ment Clerk, receipt of which is hereby acknow-
ledged.

Angela B. Boston 6/15/84
Clerk Date

Best Available Copy

11461463

BOOK PAGE
SEMINOLE COUNTY
FLORIDA

This Warranty Deed made and executed the 29th day of September A D 1977 by

HOOVER BALL AND BEARING COMPANY
a corporation existing under the laws of Michigan and having its principal place of
business at Post Office Box 10033, Ann Arbor, Michigan 48106
hereinafter called the grantor to

SINCLAIR & VALENTINE COMPANY, INC., a Delaware corporation,
whose postoffice address is Liberty Lane, Hampton, New Hampshire 03842

hereinafter called the grantee

Witnesseth: That the grantor for and in consideration of the sum of \$ 10.00 and other
valuable considerations, receipt whereof is hereby acknowledged by these presents does grant, bargain, sell,
also, remise, release, convey and confirm unto the grantee, all that certain land situate in
County Florida viz:

Lot 6, SHUMAN'S ADDITION TO EUREKA HAMMOCK, according to the plat
thereof as recorded in Plat Book 2, Page 51, of the Public Records of
Seminole County, Florida.

Subject to reservations in that certain Deed from the State of
Florida through the Trustees of the Internal Improvement Fund of
the State of Florida to T. J. Williams and R. W. Williams dated
May 21, 1946 and recorded on June 26, 1946 in Deed Book 132, Page
336, which reservations have been partially released pursuant to
those three certain Quitclaim Deeds dated February 7, 1973 and
recorded on February 22, 1973 in Official Records Book 969, Pages
1103, 1104 and 1105, respectively, Public Records of Seminole
County, Florida;

Together with all the tenements, hereditaments and appurtenances thereto belonging or in any
wise appertaining.

To Have and to Hold, the same in fee simple forever.

And the grantor hereby covenants with said grantee that it is lawfully seized of said land in fee
simple; that it has good right and lawful authority to sell and convey said land; that it hereby fully war-
rants the title to said land and will defend the same against the lawful claims of all persons whomsoever;
and that said land is free of all encumbrances except taxes accruing subsequent to
December 31, 1976, the reservations described above, and that
certain Easement executed by Hoover Ball and Bearing, Inc. in
favor of Florida Power Corporation dated May 8, 1973 and recorded
June 4, 1973, in Official Records Book 982, Page 695, of the
Public Records of Seminole County, Florida.

In Witness Whereof the grantor has caused these presents to
be executed in its name and its corporate seal to be hereunto affixed, by its
proper officers thereunto duly authorized, the day and year first above written.

ATTEST: *[Signature]* HOOVER BALL AND BEARING COMPANY

Signed, sealed and delivered in the presence of:
[Signature] By *[Signature]* Vice President

STATE OF Michigan
COUNTY OF Washtenaw

I HEREBY CERTIFY that on this day before me, an officer duly authorized in the State and County aforesaid, the above-
signed and subscribed Robert E. Ressler and Douglas E. Peck,
respectively,
and known to me to be the Vice President and Secretary,

WITNESS my hand and official seal in the County and State last aforesaid the 29th day of September A D 1977.

This instrument prepared by:
[Signature]
HAYNE L. MEYER

DOCUMENTARY SURTAX FLORIDA
280.00
88.60
975.00

UNITED LIFE INSURANCE COMPANY
P O Box 751
Orlando Florida 32802

Return to:

GEORGE F. ... & GOSTER

DEPT ORLANDO
EXHIBIT NO. 1

077708

6 12 20 PM '77

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October 13, 1983

ESTIMATED
OVERALL COMPLIANCE SCHEDULE
DAVID M COMPANY VOC CONTROL PROJECT

PHASE

Data Accumulation	January 1, 1984
Spreader Enclosure - (1) For Temperature Evaluation	January 15, 1984
Process Modifications and Development	April 1, 1984
Methods Comparison/Analysis - Thermal Incineration Vs. Recovery	May 15, 1984
Final Specification Development	July 1, 1984
→ Final Specification/Quotations and Analysis	September 1, 1984
Appropriations Request Development and Approval	December 1, 1984
Oven Enclosure and Exhaust System Installation	March 1, 1985
Solvent Recovery Delivery and Installation	August 1, 1985
Operational Shakedown, Debugging and Compliance Testing	December 1, 1985

STATE OF FLORIDA
DEPARTMENT OF ENVIRONMENTAL REGULATION

No 81922

RECEIPT FOR APPLICATION FEES AND MISCELLANEOUS REVENUE

Received from Sinclair & Valentine Co. Date May 14, 1954
Address 76 W. University Ave. Des Moines Dollars \$ 10,000.00
Applicant Name & Address _____
Source of Revenue Consent Order Case #
Revenue Code 204001 06th 272 Application Number N/A
By K. L. Litch



DAVID M COMPANY

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

June 29, 1984

Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-8241

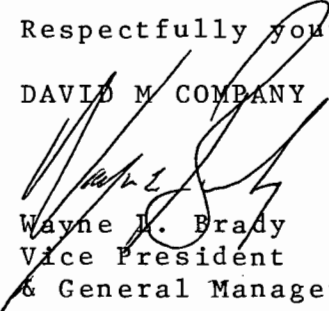
Case No. 84-0044

Gentlemen:

In accordance with the requirements stated in paragraph 8 of the Consent Order (Ref. Case No. 84-0044), entered into between the State of Florida Department of Environmental Regulation and David M Company, please find the attached report and exhibits.

Respectfully yours,

DAVID M COMPANY


Wayne L. Brady
Vice President
& General Manager

*They says they are enclosing report.
Evaluating VOC new vs incinerator
want to recover*

June 28, 1984

REPORT TO FLORIDA DEPT. OF ENVIRONMENTAL REGULATIONS

This report has been prepared in compliance with the Consent Order (Ref. Case No. 84-044, Par. 8). We have been working on this project according to the agreed schedule contained in the Consent Order. The following items refer to that schedule.

Item 1 - Data Accumulation

As background for determination of the capacity of the unit required to take care of the VOC emissions in this plant, we did extensive testing and recording of process data on our spreader lines. We recorded blanket size, pounds of cement applied per pass, time per pass, and then calculated the amount of solvent (toluene) per pass and per minute. The assumption made was that 100% of the solvent was flashed off in the spreader and it was this amount of VOC emitted per minute that we were particularly interested in. Copies of the data are attached as Exhibit I; the column headed "TOL - MIN" is the calculated emission in pounds per minute. A review of the data shows the wide fluctuation of this factor. We have been working on process revisions to reduce the peak to 4.5#/minute.

Item 2 - Spreader Enclosure

We have been aware that the enclosures on the three spreader lines would not be adequate to give us the capture efficiency required for the VOC control system. Because these lines are

equipped with steam coils to force dry the product, we were concerned that drying conditions would change with the revision of the enclosure and adversely affect the product quality. The enclosure on one spreader has been changed per drawings E0379-001 and -002 attached as Exhibit II. To date, this modification has not hurt product quality but, after reviewing the performance of this line, we have concluded the capture efficiency is not good enough. We have now contracted to have the "undercoil flooring" sections removed and panels added to the side of the structure. In this manner, the line will be totally enclosed with controlled air intake openings at each end.

Item 3 - Process Modifications and Development

Having accumulated much data in Phase I and enclosed one spreader (Phase II), David M Company then set out to better control run speed, thickness of spread and overall process control. In order to accomplish this end, tachometers and dial gauges have been added to our enclosed spreader. With these controls in place, we set out to evaluate: (1) our efficiency of capture with the newly-enclosed hood; (2) our ability to run at 40% of LEL and (3) the ability to continuously monitor fume generation in the enclosure via an infrared gas detection system. You will find attached as Exhibit III three example data sheets which summarize the type of information gathered by this phase of the project.

This information suggested that 40% of LEL operation and monitoring is feasible. In addition, it suggested that a significant amount of solvent was escaping underneath the spreader which

is currently unenclosed. As a result, the enclosure will be extended to the floor, yielding a totally enclosed spreader with a capture efficiency that we consider to be acceptable. The additional sheet metal panels will be installed the week of July 6th and additional diagnostic tests will be performed to assure that the enclosure lives up to our expectations. In addition, a permanent LEL detection system for this first unit is now on order and is expected to arrive and be installed in mid-August.

Item 4 - Methods Comparison

We have felt from the outset that our direction would be toward solvent recovery and have made an analysis of the comparative costs of incineration vs. recovery. However, no final decision has as yet been reached; and, therefore, we are leaving our options open, until such time as final costs of these two comparative systems are known.

We now have sufficient process data from which the air volume requirements of a control device may be calculated. In compliance with Exhibit II of the Consent Agreement, we will begin developing our final specifications for bids for the job on July 1, 1984. We are also expecting completion of same by September 1, 1984, in compliance with our overall compliance schedule. We would also expect to deliver the remaining information necessary to complete our permit application concurrently on September 1, 1984.

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DAVID M TOLUENE ANALYSIS

PAGE 22

DATE	LEN	WIDTH	DESCRIPTION	X TOL	LBS	WINS	SO YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN	
1/12/83	120	80.00	RUBBER 1492 1ST PASS	61.8	27.0	5.7	266.66	.101	16.586	.062	.139	2.927	63.15	63.15	
1/12/83	120	80.00	2ND PASS	61.8	21.0	9.7	266.66	.078	12.978	.048	.108	1.337	37.11	37.11	
1/12/83	120	80.00	3RD PASS	61.8	15.0	4.9	266.66	.056	9.270	.034	.077	1.091	33.46	33.46	
1/12/83	120	80.00	RUBBER 7002 1ST PASS	60.8	55.0	6.9	266.66	.206	33.440	.125	.278	4.845	52.17	52.17	
1/12/83	120	80.00	2ND PASS	60.8	47.0	4.5	266.66	.176	28.576	.107	.238	6.350	80.00	80.00	
1/12/83	120	80.00	3RD PASS	60.8	60.0	5.7	266.66	.225	36.480	.136	.304	6.400	63.15	63.15	
1/12/83	120	80.00	4TH PASS	60.8	51.0	6.9	266.66	.191	31.008	.116	.258	4.777	6.54	6.54	
1/12/83	120	80.00	5TH PASS	60.8	38.0	4.2	266.66	.142	23.104	.086	.192	5.500	85.71	85.71	
1/12/83	120	80.00	6TH PASS	60.8	49.0	4.3	266.66	.183	29.792	.111	.248	6.928	63.72	63.72	
1/12/83	120	80.00	7TH PASS	60.8	27.0	3.4	266.66	.101	16.416	.061	.136	4.828	105.88	105.88	
1/12/83	120	80.00	8TH PASS	60.8	39.0	4.2	266.66	.146	23.712	.088	.197	6.045	65.71	65.71	
1/12/83	120	80.00	9TH PASS	60.8	33.0	4.4	266.66	.123	20.064	.075	.167	4.560	81.81	81.81	
1/12/83	120	80.00	10TH PASS	60.8	36.0	4.7	266.66	.135	21.888	.082	.182	4.657	76.59	76.59	
1/12/83	120	80.00	11TH PASS	60.8	34.0	7.3	266.66	.127	20.672	.077	.172	2.831	49.31	49.31	
1/12/83	120	80.00	12TH PASS	60.8	28.0	3.4	266.66	.105	17.024	.063	.141	5.007	105.88	105.88	
1/12/83	120	80.00	13TH PASS	60.8	24.0	3.2	266.66	.090	14.592	.054	.121	4.560	112.50	112.50	
1/12/83	120	80.00	14TH PASS	60.8	24.0	4.0	266.66	.090	14.592	.054	.121	3.648	90.00	90.00	
1/12/83	120	80.00	RUBBER 2202 1ST PASS	70.5	40.0	11.9	266.66	.150	24.320	.091	.202	2.043	30.25	30.25	
1/12/83	120	80.00	RUBBER 2202 1ST PASS	70.5	15.0	7.9	266.66	.056	10.575	.039	.088	1.338	45.56	45.56	
1/12/83	120	80.00	2ND PASS	70.5	27.0	5.0	266.66	.101	19.035	.071	.158	3.807	72.00	72.00	
1/12/83	120	80.00	3RD PASS	70.5	25.0	4.7	266.66	.093	17.625	.066	.146	3.750	76.59	76.59	
1/12/83	120	80.00	4TH PASS	70.5	49.0	6.6	266.66	.193	34.545	.129	.287	5.234	54.54	54.54	
1/12/83	120	80.00	5TH PASS	70.5	39.0	5.3	266.66	.146	27.495	.103	.229	5.187	67.92	67.92	
1/12/83	120	80.00	6TH PASS	70.5	33.0	4.3	266.66	.123	23.265	.087	.193	5.410	83.72	83.72	
1/12/83	120	80.00	7TH PASS	70.5	32.0	4.7	266.66	.120	22.560	.084	.188	4.800	76.59	76.59	
1/12/83	120	80.00	8TH PASS	70.5	28.0	3.7	266.66	.105	19.740	.074	.164	5.335	97.29	97.29	
1/12/83	120	80.00	9TH PASS	70.5	22.0	3.5	266.66	.082	15.510	.058	.129	4.431	102.65	102.65	
1/12/83	120	80.00	10TH PASS	70.5	21.0	4.1	266.66	.078	14.805	.055	.123	3.610	87.80	87.80	
1/12/83	120	80.00	11TH PASS	70.5	26.0	4.4	266.66	.097	16.330	.060	.152	4.165	81.81	81.81	
1/12/83	120	80.00	12TH PASS	70.5	51.0	7.6	266.66	.191	35.955	.134	.299	4.730	47.36	47.36	
1/12/83	120	80.00	13TH PASS	70.5	16.0	3.7	266.66	.060	11.280	.042	.094	3.048	97.29	97.29	
1/12/83	120	80.00	14TH PASS	70.5	14.0	5.2	266.66	.052	9.870	.037	.082	1.898	69.23	69.23	
1/12/83	120	80.00	15TH PASS	70.5	16.0	3.9	266.66	.060	11.280	.042	.094	2.892	92.30	92.30	
1/12/83	120	80.00	TALI PASS	70.5	21.0	18.9	266.66	.078	14.805	.055	.123	.783	19.04	19.04	
TOTALS					1062.0	231.9	8799.78	.120	686.484						

TOTAL PASSES 33
 TOTAL PRODUCTION TIME 04:32:38
 AVERAGE LBS CEMENT PER PASS 32.161
 AVERAGE LBS CEMENT PER SQUARE YARD 3.982

MIN 3.2	MIN .034	.077	5.64	6.64
MAX 64.9	MAX .136	.304	6.928	112.50
AVG 7.0	AVG .077	.172	4.062	73.78

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RUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 21

DATE	LEN	WIDTH	DESCRIPTION	TOL	LBS	MINS	SO YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LVD	TOL MIN	FEET MIN	F/FEET MIN
1/07/83	100	60.00	RUBBER 1492 1ST PASS	62.5	37.0	5.0	166.66	.222	23.125	.138	.231	2.625	100.00	35.00
1/07/83	100	60.00	2ND PASS	62.5	18.0	4.2	166.66	.108	11.250	.067	.112	2.678	71.42	53.57
1/07/83	100	60.00	3RD PASS	62.5	12.0	3.7	166.66	.072	7.500	.045	.075	2.027	41.08	30.81
1/07/83	100	60.00	RUBBER 7002 1ST PASS	61.0	28.0	7.5	166.66	.168	17.080	.102	.170	2.277	40.00	30.00
1/07/83	100	60.00	2ND PASS	61.0	32.0	6.2	166.66	.192	19.520	.117	.195	3.148	48.38	36.29
1/07/83	100	60.00	3RD PASS	61.0	39.0	6.2	166.66	.234	23.790	.142	.237	3.837	48.38	36.29
1/07/83	100	60.00	4TH PASS	61.0	32.0	4.3	166.66	.192	19.520	.117	.195	3.539	49.76	32.32
1/07/83	100	60.00	5TH PASS	61.0	45.0	8.2	166.66	.270	27.450	.164	.274	3.347	36.58	27.43
1/07/83	100	60.00	6TH PASS	61.0	44.0	8.4	166.66	.204	20.740	.124	.207	3.840	38.55	31.66
1/07/83	100	60.00	7TH PASS	61.0	33.0	3.2	166.66	.198	20.130	.120	.201	6.290	93.75	70.31
1/07/83	100	60.00	8TH PASS	61.0	24.0	3.2	166.66	.144	14.640	.087	.146	3.575	33.75	30.31
1/07/83	100	60.00	9TH PASS	61.0	20.0	2.4	166.66	.120	12.200	.073	.122	5.083	125.00	93.75
1/07/83	100	60.00	RUBBER 6609 1ST PASS	71.7	24.0	5.3	166.66	.144	17.208	.103	.172	3.248	36.60	42.46
1/07/83	100	60.00	2ND PASS	71.7	39.0	5.8	166.66	.234	27.963	.167	.279	4.821	51.72	38.79
1/07/83	100	60.00	3RD PASS	71.7	29.0	4.6	166.66	.174	20.793	.124	.207	4.620	45.21	46.91
1/07/83	100	60.00	4TH PASS	71.7	43.0	8.0	166.66	.258	30.831	.184	.308	3.853	37.50	28.12
1/07/83	100	60.00	5TH PASS	71.7	31.0	4.7	166.66	.186	22.227	.133	.222	4.729	33.82	47.87
1/07/83	100	60.00	6TH PASS	71.7	37.0	5.8	166.66	.222	26.529	.159	.265	4.573	51.72	38.79
1/07/83	100	60.00	7TH PASS	71.7	47.0	8.1	166.66	.282	33.699	.202	.336	3.219	39.17	34.87
1/07/83	100	60.00	8TH PASS	71.7	35.0	4.9	166.66	.210	25.095	.150	.250	5.121	61.22	45.91
1/07/83	100	60.00	9TH PASS	71.7	51.0	4.0	166.66	.306	36.567	.219	.365	3.141	75.00	56.25
1/07/83	100	60.00	10TH PASS	71.7	17.0	6.1	166.66	.102	12.169	.073	.121	1.998	49.18	36.88
1/07/83	100	60.00	11TH PASS	71.7	32.0	4.7	166.66	.192	22.944	.137	.229	3.881	43.82	47.87
1/07/83	100	60.00	12TH PASS	71.7	26.0	3.5	166.66	.156	18.642	.111	.186	5.326	85.71	64.28
1/07/83	100	60.00	13TH PASS	71.7	23.0	2.7	166.66	.138	16.491	.098	.164	3.107	31.11	33.33
1/07/83	100	60.00	14TH PASS	71.7	24.0	2.7	166.66	.144	17.208	.103	.172	6.373	111.11	83.33
1/07/83	100	60.00	15TH PASS	71.7	20.0	2.6	166.66	.120	14.340	.086	.143	6.515	115.38	66.53
1/07/83	100	60.00	16TH PASS	71.7	35.0	3.2	166.66	.210	25.095	.150	.250	7.842	93.75	70.31
1/07/83	100	60.00	17TH PASS	71.7	20.0	2.5	166.66	.120	14.340	.086	.143	3.736	120.00	90.00
1/07/83	100	60.00	TALC PASS	71.7	15.0	2.7	166.66	.090	10.755	.064	.107	3.983	111.11	83.33
TOTALS					887.0	134.7	8833.14	.183	599.106					

TOTAL PASSES 29
 TOTAL PRODUCTION TIME 07:19:36
 AVERAGE LBS CEMENT PER PASS 30.586
 AVERAGE LBS CEMENT PER SQUARE YARD 5.322

MIN 2.4
 MAX 8.2
 AVG 4.6

MIN .045
 MAX .219
 AVG .123

MIN .075
 MAX .365
 AVG .206

MIN 1.998
 MAX 9.141
 AVG 4.767

MIN 36.56
 MAX 125.00
 AVG 72.74

MIN 27.43
 MAX 93.75
 AVG 54.55

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RUN ON 01/23/84 AT 14:11:22

DAVID W TOLUENE ANALYSIS

PAGE 14

DATE	LEN	WIDTH	DESCRIPTION	TOL	LBS	WINS	SO YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN
0/26/83	100	80.00	RUBBER 1402 1ST PASS	57.2	25.0	2.1	222.22	.112	14.300	.064	.143	3.487	73.17	73.17
0/26/83	100	80.00	2ND PASS	57.2	20.0	2.5	222.22	.090	11.440	.051	.114	4.576	120.00	120.00
0/26/83	100	80.00	3RD PASS	57.2	16.0	3.2	222.22	.072	9.152	.041	.091	2.860	83.75	83.75
0/26/83	100	80.00	RUBBER 7002 1ST PASS	58.5	64.0	10.4	222.22	.288	37.440	.168	.374	3.600	28.84	28.84
0/26/83	100	80.00	2ND PASS	58.5	47.0	5.5	222.22	.211	27.495	.123	.274	4.999	54.54	54.54
0/26/83	100	80.00	3RD PASS	58.5	77.0	5.5	222.22	.346	45.045	.202	.450	8.190	54.54	54.54
0/26/83	100	80.00	4TH PASS	58.5	10.0	5.3	222.22	.045	5.850	.026	.058	1.103	56.60	56.60
0/26/83	100	80.00	5TH PASS	58.5	33.0	4.5	222.22	.148	19.305	.086	.193	4.290	66.66	66.66
0/26/83	100	80.00	6TH PASS	58.5	50.0	4.4	222.22	.225	29.250	.131	.292	6.647	68.18	68.18
0/26/83	100	80.00	7TH PASS	58.5	37.0	4.0	222.22	.166	21.645	.097	.216	5.411	75.00	75.00
0/26/83	100	80.00	8TH PASS	58.5	27.0	2.8	222.22	.121	15.795	.071	.157	5.641	107.14	107.14
0/26/83	100	80.00	RUBBER 7002 1ST PASS	58.5	28.0	2.8	222.22	.126	16.380	.073	.163	5.850	107.14	107.14
0/26/83	100	80.00	2ND PASS	58.5	26.0	2.1	222.22	.117	15.210	.068	.152	7.242	142.85	142.85
0/26/83	100	80.00	RUBBER 6609 1ST PASS	69.6	53.0	6.2	222.22	.238	36.868	.165	.368	5.949	48.38	48.38
0/26/83	100	80.00	2ND PASS	69.6	32.0	3.7	222.22	.144	22.272	.100	.222	6.019	81.08	81.08
0/26/83	100	80.00	3RD PASS	69.6	31.0	3.1	222.22	.139	21.576	.097	.215	6.960	96.77	96.77
0/26/83	100	80.00	4TH PASS	69.6	27.0	2.7	222.22	.121	18.792	.084	.187	6.960	111.11	111.11
0/26/83	100	80.00	5TH PASS	69.6	28.0	3.1	222.22	.126	19.488	.087	.194	6.286	96.77	96.77
0/26/83	100	80.00	6TH PASS	69.6	18.0	2.3	222.22	.081	12.528	.056	.125	5.446	130.43	130.43
0/26/83	100	80.00	7TH PASS	69.6	16.0	3.3	222.22	.085	13.224	.059	.132	4.007	90.90	90.90
0/26/83	100	80.00	8TH PASS	69.6	18.0	2.5	222.22	.081	12.528	.056	.125	5.011	120.00	120.00
0/26/83	100	80.00	CTH PASS	69.6	15.0	2.7	222.22	.067	10.440	.046	.104	3.866	111.11	111.11
TOTALS					701.0	86.7	4888.84	.243	436.043					

TOTAL PASSES 22
 TOTAL PRODUCTION TIME 05:44:25
 AVERAGE LBS CEMENT PER PASS 31.863
 AVERAGE LBS CEMENT PER SQUARE YARD 3.154

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RUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 12

DATE	LEV	WIDTH	DESCRIPTION	X TOL	LBS	MINS	SO YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN
0/24/83	100	80.00	RUBBER 1497 1ST PASS	57.2	26.0	9.9	222.22	117	14.872	0.066	1.148	1.502	58.30	80.30
0/24/83	100	80.00	2ND PASS	57.2	29.0	7.2	222.22	130	16.588	0.074	1.165	2.303	41.66	41.66
0/24/83	100	80.00	3RD PASS	57.2	19.0	5.6	222.22	085	10.860	0.048	0.108	1.940	33.57	33.57
0/24/83	100	80.00	RUBBER 7002 1ST PASS	59.8	45.0	7.2	222.22	202	26.910	0.121	0.269	3.737	41.66	41.66
0/24/83	100	80.00	2ND PASS	59.8	42.0	5.2	222.22	189	25.116	0.113	0.251	4.830	57.69	57.69
0/24/83	100	80.00	3RD PASS	59.8	40.0	4.4	222.22	180	23.920	0.107	0.239	5.436	68.18	68.18
0/24/83	100	80.00	4TH PASS	59.8	48.0	4.9	222.22	216	28.704	0.129	0.287	6.857	81.22	81.22
0/24/83	100	80.00	5TH PASS	59.8	42.0	4.2	222.22	189	25.116	0.113	0.251	5.980	71.42	71.42
0/24/83	100	80.00	6TH PASS	58.2	44.0	3.3	222.22	198	25.608	0.115	0.256	7.760	90.90	90.90
0/24/83	100	80.00	7TH PASS	58.2	36.0	4.1	222.22	162	20.952	0.094	0.209	5.110	73.17	73.17
0/24/83	100	80.00	8TH PASS	58.2	27.0	3.4	222.22	121	15.714	0.070	0.157	4.621	68.23	68.23
0/24/83	100	80.00	9TH PASS	58.2	25.0	2.6	222.22	112	14.550	0.065	0.145	5.596	115.38	115.38
0/24/83	100	80.00	10TH PASS	58.2	21.0	2.7	222.22	094	12.222	0.054	0.122	4.526	111.11	111.11
0/24/83	100	80.00	RUBBER 7002 1ST PASS	58.2	21.0	3.5	222.22	094	12.222	0.054	0.122	3.492	85.71	85.71
0/24/83	100	80.00	2ND PASS	58.2	25.0	2.7	222.22	112	14.550	0.065	0.145	5.368	111.11	111.11
0/24/83	100	80.00	3RD PASS	58.2	44.0	2.9	222.22	198	25.608	0.115	0.256	6.830	103.44	103.44
0/24/83	100	80.00	RUBBER 5503 1ST PASS	65.4	26.0	65.0	222.22	117	17.004	0.076	0.170	4.261	4.61	4.61
0/24/83	100	80.00	2ND PASS	65.4	29.0	4.0	222.22	130	18.966	0.085	0.189	4.741	75.00	75.00
0/24/83	100	80.00	3RD PASS	65.4	36.0	6.8	222.22	162	23.544	0.105	0.235	5.462	84.81	84.81
0/24/83	100	80.00	4TH PASS	65.4	34.0	4.1	222.22	153	22.236	0.100	0.222	5.423	73.17	73.17
0/24/83	100	80.00	5TH PASS	65.4	29.0	3.4	222.22	130	18.966	0.085	0.189	5.578	88.23	88.23
0/24/83	100	80.00	6TH PASS	65.4	23.0	4.8	222.22	103	15.042	0.067	0.150	3.133	62.50	62.50
TOTALS					711.0	161.9	4898.84	145	420.278					

TOTAL PASSES 22
 TOTAL PRODUCTION TIME 05:35:40
 AVERAGE LBS CEMENT PER PASS 32.318
 AVERAGE LBS CEMENT PER SQUARE YARD 3.199

MIN 2.6	MIN 0.048
MAX 65.0	MAX 0.129
AVG 7.3	AVG 0.087

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JUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 9

DATE	LEN	WIDTH	DESCRIPTION	X TOL	LBS	WINS	SD YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LVD	TOL WIN	FEET WIN	F/FEET WIN	
9/26/83	98	78.50	RUBBER 7002 1ST PASS	67.5	36.0	3.7	213.68	168	24.300	.113	.247	3.626	43.88	43.05	
9/26/83	98	78.50	2ND PASS	67.5	21.0	4.6	213.68	.098	14.175	.066	.144	3.061	63.91	62.71	
9/26/83	98	78.50	3RD PASS	67.5	14.0	5.2	213.68	.065	9.450	.044	.096	2.250	70.00	68.68	
9/26/83	98	78.50	RUBBER 7002 1ST PASS	60.2	28.0	5.3	213.68	.131	16.856	.078	.172	3.180	55.47	54.43	
9/26/83	98	78.50	2ND PASS	60.2	33.0	5.4	213.68	.154	19.866	.092	.202	5.842	66.47	64.84	
9/26/83	98	78.50	3RD PASS	60.2	45.0	5.7	213.68	.210	27.090	.126	.276	4.752	51.57	50.61	
9/26/83	98	78.50	4TH PASS	60.2	37.0	5.3	213.68	.173	22.274	.104	.227	5.160	68.37	67.06	
9/26/83	98	78.50	5TH PASS	60.2	33.0	5.3	213.68	.154	19.866	.092	.202	6.020	89.09	87.41	
9/26/83	98	78.50	6TH PASS	60.2	49.0	5.2	213.68	.229	29.498	.138	.301	9.218	91.87	90.15	
9/26/83	98	78.50	7TH PASS	60.2	24.0	4.5	213.68	.112	14.448	.067	.147	3.210	65.33	64.10	
9/26/83	98	78.50	8TH PASS	60.2	55.5	5.4	213.68	.259	33.411	.156	.340	13.921	122.50	120.20	
9/26/83	98	78.50	9TH PASS	60.2	31.5	5.6	213.68	.147	18.963	.088	.193	3.386	52.50	51.51	
9/26/83	98	78.50	10TH PASS	60.2	36.0	5.4	213.68	.168	21.672	.101	.221	6.374	66.47	64.84	
9/26/83	98	78.50	11TH PASS	60.2	28.5	4.5	213.68	.133	17.157	.080	.175	3.812	65.33	64.10	
9/26/83	98	78.50	12TH PASS	60.2	27.0	5.6	213.68	.126	16.254	.076	.165	6.251	113.07	110.95	
9/26/83	98	78.50	13TH PASS	60.2	19.0	2.4	213.68	.088	11.438	.053	.116	4.765	122.50	120.20	
9/26/83	98	78.50	14TH PASS	60.2	22.0	2.5	213.68	.102	13.244	.061	.135	5.297	117.60	115.39	
9/26/83	98	78.50	RUBBER 7002 1ST PASS	58.7	21.5	2.5	213.68	.100	12.520	.059	.128	5.042	117.60	115.39	
9/26/83	98	78.50	2ND PASS	58.7	20.5	2.7	213.68	.095	12.033	.056	.122	4.456	108.88	106.84	
9/26/83	98	78.50	3RD PASS	58.7	19.0	2.7	213.68	.089	11.153	.052	.113	4.130	108.88	106.84	
9/26/83	98	78.50	4TH PASS	58.7	17.0	2.3	213.68	.079	9.979	.046	.101	4.338	127.82	125.42	
9/26/83	98	78.50	5TH PASS	58.7	17.0	2.3	213.68	.079	9.979	.046	.101	4.338	127.82	125.42	
9/26/83	98	78.50	6TH PASS	58.7	26.0	2.4	213.68	.121	15.262	.071	.155	4.359	122.50	120.20	
9/26/83	98	78.50	RUBBER 5503 1ST PASS	64.2	16.0	2.6	213.68	.082	12.198	.057	.124	4.691	113.07	110.95	
9/26/83	98	78.50	2ND PASS	64.2	18.0	2.7	213.68	.084	11.556	.054	.117	4.288	108.88	106.84	
9/26/83	98	78.50	3RD PASS	64.2	28.0	3.0	213.68	.131	17.976	.084	.183	5.992	96.00	96.16	
9/26/83	98	78.50	4TH PASS	64.2	23.0	2.7	213.68	.107	14.766	.069	.150	5.468	108.88	106.84	
9/26/83	98	78.50	5TH PASS	64.2	21.0	2.7	213.68	.098	13.482	.063	.137	4.953	108.88	106.84	
9/26/83	98	78.50	6TH PASS	64.2	18.0	2.9	213.68	.084	11.556	.054	.117	3.984	101.37	99.47	
9/26/83	98	78.50	7TH PASS	64.2	20.0	3.2	213.68	.093	12.840	.060	.131	4.612	91.87	90.15	
9/26/83	98	78.50	8TH PASS	64.2	19.0	2.4	213.68	.088	12.198	.057	.124	5.062	122.50	120.20	
9/26/83	98	78.50	9TH PASS	64.2	18.0	2.4	213.68	.084	11.556	.054	.117	4.215	122.50	120.20	
9/26/83	98	78.50	10TH PASS	64.2	16.0	2.3	213.68	.074	10.272	.048	.104	4.466	127.82	125.42	
9/26/83	98	78.50	11TH PASS	64.2	16.0	2.4	213.68	.074	10.272	.048	.104	4.280	122.50	120.20	
9/26/83	98	78.50	12TH PASS	64.2	16.0	2.7	213.68	.074	10.272	.048	.104	3.804	108.88	106.84	
9/26/83	98	78.50	13TH PASS	64.2	17.0	2.3	213.68	.079	10.914	.051	.111	4.745	127.82	125.42	
9/26/83	98	78.50	14TH PASS	64.2	16.0	2.5	213.68	.074	10.272	.048	.104	4.168	117.60	115.39	
9/26/83	98	78.50	TALC PASS	64.2	18.0	15.2	213.68	.084	11.556	.054	.117	4.760	19.34	18.97	
TOTALS					925.5	120.3	7906.16	.117	671.818						

TOTAL PASSES 37
 TOTAL PRODUCTION TIME 07:57:46
 AVERAGE LBS CEMENT PER PASS 25.023
 AVERAGE LBS CEMENT PER SQUARE YARD 4.331

MIN 2.3	MIN -.044
MAX 6.7	MAX .156
AVG 3.2	AVG -.071

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RUN ON 01/23/84 AT 14:11:22

DAVID W TOLUENE ANALYSIS

PAGE 7

DATE	LEN	WIDTH	DESCRIPTION	TOL	LBS	MIN	SD YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN	
09/22/83	100	81.75	RUBBER 1492 1ST PASS	57.8	36.0	4.2	227.08	158	20.808	.091	.208	3.356	88.38	89.44	
09/22/83	100	81.75	2ND PASS	57.8	26.0	4.8	227.08	114	15.028	.065	.150	3.130	62.50	63.86	
09/22/83	100	81.75	3RD PASS	57.8	18.0	5.1	227.08	679	10.404	.045	.104	2.040	58.82	60.11	
09/22/83	100	81.75	RUBBER 7002 1ST PASS	57.4	27.0	3.8	227.08	118	15.498	.068	.154	4.078	78.94	80.67	
09/22/83	100	81.75	2ND PASS	57.4	30.0	4.5	227.08	132	17.220	.075	.172	3.826	66.66	68.12	
09/22/83	100	81.75	3RD PASS	57.4	41.5	4.3	227.08	182	23.821	.104	.238	5.539	69.76	71.29	
09/22/83	100	81.75	4TH PASS	57.4	38.0	3.4	227.08	167	21.812	.096	.218	6.415	68.23	70.16	
09/22/83	100	81.75	5TH PASS	57.4	33.5	3.2	227.08	147	19.229	.084	.192	6.009	93.75	95.80	
09/22/83	100	81.75	6TH PASS	57.4	53.0	5.2	227.08	233	30.822	.133	.304	8.906	88.38	89.94	
09/22/83	100	81.75	7TH PASS	57.4	40.0	3.3	227.08	176	22.960	.101	.229	6.957	90.90	92.90	
09/22/83	100	81.75	8TH PASS	57.4	35.0	3.0	227.08	154	20.090	.088	.200	6.696	100.00	102.19	
09/22/83	100	81.75	9TH PASS	57.4	45.5	3.9	227.08	200	26.117	.115	.261	6.696	76.92	78.60	
09/22/83	100	81.75	10TH PASS	57.4	40.5	2.7	227.08	178	23.247	.102	.232	8.610	111.11	113.54	
09/22/83	100	81.75	11TH PASS	57.4	36.0	3.7	227.08	158	20.664	.090	.206	5.584	81.08	82.85	
09/22/83	100	81.75	12TH PASS	57.4	32.0	3.1	227.08	140	18.368	.080	.183	5.925	66.77	68.89	
09/22/83	100	81.75	RUBBER 7002 1ST PASS	54.5	58.5	6.1	227.08	257	31.982	.140	.318	5.226	49.18	50.25	
09/22/83	100	81.75	2ND PASS	54.5	46.0	4.1	227.08	202	25.070	.110	.250	6.114	73.17	74.77	
09/22/83	100	81.75	3RD PASS	54.5	41.5	3.7	227.08	182	22.617	.099	.226	6.112	81.08	82.85	
09/22/83	100	81.75	4TH PASS	54.5	29.0	3.5	227.08	127	15.805	.069	.158	4.515	65.71	67.59	
09/22/83	100	81.75	RUBBER 6607 1ST PASS	67.3	44.0	5.1	227.08	193	29.612	.130	.296	5.806	58.82	60.11	
09/22/83	100	81.75	2ND PASS	67.3	35.0	4.8	227.08	154	23.555	.103	.235	5.888	75.00	76.84	
09/22/83	100	81.75	3RD PASS	67.3	30.0	4.3	227.08	132	20.190	.088	.201	4.695	69.76	71.29	
09/22/83	100	81.75	RUBBER 6607 1ST PASS	65.8	21.0	3.0	227.08	92	13.818	.060	.136	4.606	100.00	102.19	
09/22/83	100	81.75	2ND PASS	65.8	26.5	3.8	227.08	125	18.753	.082	.187	4.935	78.94	80.67	
09/22/83	100	81.75	3RD PASS	65.8	14.0	2.7	227.08	661	9.212	.040	.092	3.911	111.11	113.54	
09/22/83	100	81.75	4TH PASS	65.8	51.5	5.5	227.08	226	33.887	.149	.338	5.161	54.54	55.78	
09/22/83	100	81.75	5TH PASS	65.8	44.0	4.1	227.08	193	28.952	.127	.289	7.861	73.17	74.77	
09/22/83	100	81.75	6TH PASS	65.8	35.0	3.8	227.08	154	23.030	.101	.230	6.060	78.94	80.67	
09/22/83	100	81.75	7TH PASS	65.8	17.0	3.3	227.08	874	11.186	.049	.111	2.601	69.76	71.29	
09/22/83	100	81.75	TALC PASS	65.8	24.0	17.0	227.08	105	15.792	.069	.157	.928	17.64	18.03	
TOTALS					1027.0	119.2	6585.32	-155	613.257						

TOTAL PASSES	29	MIN	2.7	MIN	.040	.092	2.040	48.38	89.44
TOTAL PRODUCTION TIME	07:42:55	MAX	6.2	MAX	.149	.338	8.610	111.11	113.54
AVERAGE LBS CEMENT PER PASS	35.413	AVG	4.1	AVG	.092	.211	6.274	76.94	78.62
AVERAGE LBS CEMENT PER SQUARE YARD	4.522								

*2% TOL CHANGED
 FROM 57.4% TO
 54.5% ON THIS
 PASS*

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RUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 6

DATE	LEN	WIDTH	DESCRIPTION	X TOL	LBS	MINS	50 YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN	
09/21/83	100	46.00	RUBBER 1492 1ST PASS	57.8	20.0	5.5	127.77	.156	8.560	.090	.215	1.778	66.15	26.53	
09/21/83	100	46.00	2ND PASS	57.8	11.5	4.1	127.77	.090	6.647	.052	.066	1.621	73.17	42.07	
09/21/83	100	46.00	3RD PASS	57.8	8.0	4.4	127.77	.062	4.624	.036	.046	1.050	80.18	39.20	
09/21/83	100	46.00	RUBBER 7002 1ST PASS	57.4	18.0	4.5	127.77	.140	10.332	.080	.103	2.296	66.66	38.33	
09/21/83	100	46.00	2ND PASS	57.4	22.0	2.9	127.77	.172	12.628	.098	.126	3.354	103.44	59.48	
09/21/83	100	46.00	3RD PASS	57.4	18.0	3.0	127.77	.140	10.332	.080	.103	3.444	100.00	57.50	
09/21/83	100	46.00	4TH PASS	57.4	13.0	2.4	127.77	.101	7.462	.058	.074	3.109	125.00	71.87	
09/21/83	100	46.00	RUBBER 7002 1ST PASS	57.4	42.0	5.0	127.77	.328	24.108	.188	.241	4.821	60.00	34.50	
09/21/83	100	46.00	2ND PASS	57.4	13.0	4.2	127.77	.101	7.462	.058	.074	3.554	62.50	35.93	
09/21/83	100	46.00	3RD PASS	57.4	9.0	4.0	127.77	.070	5.166	.040	.051	1.291	75.00	43.12	
09/21/83	100	46.00	4TH PASS	57.4	11.0	2.2	127.77	.086	6.314	.049	.063	2.070	136.36	78.40	
09/21/83	100	46.00	5TH PASS	57.4	9.0	3.4	127.77	.070	5.166	.040	.051	1.519	88.23	50.73	
09/21/83	100	46.00	6TH PASS	57.4	11.0	2.7	127.77	.086	6.314	.049	.063	2.338	111.11	63.88	
09/21/83	100	46.00	7TH PASS	57.4	9.0	1.9	127.77	.070	5.166	.040	.051	2.718	157.89	90.78	
09/21/83	100	46.00	8TH PASS	57.4	20.0	3.2	127.77	.156	11.480	.089	.114	3.567	93.75	63.00	
09/21/83	100	46.00	9TH PASS	57.4	12.0	2.8	127.77	.093	6.888	.053	.068	2.460	107.14	61.60	
09/21/83	100	46.00	10TH PASS	57.4	12.0	2.0	127.77	.093	6.888	.053	.068	3.444	158.00	86.25	
09/21/83	100	46.00	11TH PASS	57.4	11.0	2.3	127.77	.096	6.314	.049	.063	2.745	130.43	75.00	
09/21/83	100	46.00	12TH PASS	57.4	9.0	2.0	127.77	.070	5.166	.040	.051	2.583	158.00	86.25	
09/21/83	100	46.00	13TH PASS	57.4	10.0	2.0	127.77	.078	5.740	.044	.057	2.870	150.00	86.25	
09/21/83	100	46.00	14TH PASS	57.4	7.0	1.9	127.77	.054	4.018	.031	.040	2.114	157.89	90.78	
09/21/83	100	46.00	15TH PASS	57.4	8.0	1.8	127.77	.062	4.592	.035	.045	2.551	166.66	95.83	
09/21/83	100	46.00	16TH PASS	57.4	18.0	2.9	127.77	.140	10.332	.080	.103	3.562	103.44	59.48	
09/21/83	100	46.00	17TH PASS	57.4	14.0	2.3	127.77	.109	8.036	.062	.080	3.453	130.43	75.00	
09/21/83	100	46.00	18TH PASS	57.4	13.0	2.0	127.77	.101	7.462	.058	.074	3.731	158.00	86.25	
09/21/83	100	46.00	RUBBER 7002 1ST PASS	57.4	12.0	2.0	127.77	.093	6.888	.053	.068	3.444	158.00	86.25	
09/21/83	100	46.00	2ND PASS	57.4	12.0	2.0	127.77	.093	6.888	.053	.068	3.444	158.00	86.25	
09/21/83	100	46.00	3RD PASS	57.4	9.0	2.6	127.77	.070	5.166	.040	.051	1.985	115.38	66.34	
09/21/83	100	46.00	RUBBER 6607 1ST PASS	67.3	18.0	4.1	127.77	.140	12.114	.094	.121	2.954	73.17	42.07	
09/21/83	100	46.00	2ND PASS	67.3	12.0	2.7	127.77	.092	8.076	.062	.080	2.951	111.11	63.88	
09/21/83	100	46.00	3RD PASS	67.3	13.0	2.4	127.77	.101	8.749	.068	.087	3.645	125.00	71.87	
09/21/83	100	46.00	4TH PASS	67.3	26.0	3.5	127.77	.203	17.498	.136	.174	4.096	85.71	49.28	
09/21/83	100	46.00	5TH PASS	67.3	29.0	3.5	127.77	.226	19.517	.152	.195	6.576	85.71	49.28	
09/21/83	100	46.00	6TH PASS	67.3	14.0	2.9	127.77	.109	9.422	.073	.094	3.248	103.44	59.48	
09/21/83	100	46.00	7TH PASS	67.3	6.0	1.9	127.77	.046	4.638	.031	.040	2.125	157.89	90.78	
09/21/83	100	46.00	8TH PASS	67.3	10.0	2.4	127.77	.078	6.730	.052	.067	2.804	125.00	71.87	
09/21/83	100	46.00	9TH PASS	67.3	10.0	2.2	127.77	.078	6.730	.052	.067	3.059	136.36	78.40	
09/21/83	100	46.00	10TH PASS	67.3	9.5	2.5	127.77	.074	6.353	.050	.063	2.557	120.00	69.00	
09/21/83	100	46.00	11TH PASS	67.3	9.0	3.1	127.77	.070	6.057	.047	.060	1.953	84.77	65.64	
09/21/83	100	46.00	12TH PASS	67.3	4.5	1.0	127.77	.035	3.028	.023	.030	3.028	300.00	172.50	
09/21/83	100	46.00	TALC PASS	67.3	8.0	14.9	127.77	.062	8.384	.042	.053	3.361	20.13	11.57	
TOTALS					542.5	115.8	5110.60	.106	327.491						

TOTAL PASSES 40
 TOTAL PRODUCTION TIME 07:42:04
 AVERAGE LBS CEMENT PER PASS 13.562
 AVERAGE LBS CEMENT PER SQUARE YARD 4.245

MIN 1.0
 MAX 6.5
 AVG 2.6

MIN .023
 MAX .188
 AVG .063

MIN .030
 MAX .241
 AVG .081

MIN 1.050
 MAX 3.576
 AVG 2.892

MIN 46.15
 MAX 360.00
 AVG 117.47

MIN 26.53
 MAX 172.50
 AVG 67.54

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RUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 4

DATE	LEN	WIDTH	DESCRIPTION	% TOL	LBS	WINS	SO VDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN	
09/19/83	95	80.50	RUBBER 1492 1ST PASS	55.3	35.0	5.9	212.42	.164	29.355	.091	.203	2.260	68.30	82.60	
09/19/83	95	80.50	2ND PASS	55.3	22.0	4.3	212.42	.103	12.166	.057	.128	2.829	66.27	66.69	
09/19/83	95	80.50	3RD PASS	55.3	15.0	3.7	212.42	.070	6.295	.039	.087	2.241	77.02	77.50	
09/19/83	95	80.50	RUBBER 7002 1ST PASS	59.3	34.5	6.1	212.42	.162	20.458	.096	.215	3.353	46.72	47.01	
09/19/83	95	80.50	2ND PASS	59.3	51.0	5.4	212.42	.240	30.243	.142	.318	3.600	62.77	53.10	
09/19/83	95	80.50	3RD PASS	59.3	25.0	4.0	212.42	.117	14.825	.069	.156	3.706	71.25	71.69	
09/19/83	95	80.50	4TH PASS	59.3	30.5	4.3	212.42	.143	18.086	.085	.190	3.206	66.27	66.69	
09/19/83	95	80.50	5TH PASS	59.3	33.5	4.1	212.42	.157	19.865	.093	.209	4.845	69.51	69.94	
09/19/83	95	80.50	6TH PASS	59.3	36.5	3.2	212.42	.218	27.574	.129	.290	3.616	69.06	69.61	
09/19/83	95	80.50	7TH PASS	59.3	40.0	4.5	212.42	.188	23.720	.111	.249	5.271	63.33	63.72	
09/19/83	95	80.50	8TH PASS	59.3	49.0	3.9	212.42	.230	29.057	.136	.305	3.450	73.07	73.63	
09/19/83	95	80.50	9TH PASS	59.3	47.0	4.1	212.42	.221	27.871	.131	.293	5.797	69.51	69.94	
09/19/83	95	80.50	10TH PASS	59.3	41.0	3.2	212.42	.193	24.313	.114	.255	3.577	75.71	76.65	
09/19/83	95	80.50	11TH PASS	59.3	35.0	3.1	212.42	.164	20.755	.097	.218	6.695	91.93	92.50	
09/19/83	95	80.50	12TH PASS	59.3	27.5	2.9	212.42	.129	16.307	.076	.171	6.623	68.27	68.68	
09/19/83	95	80.50	RUBBER 7002 1ST PASS	59.3	19.5	2.6	212.42	.091	11.563	.054	.121	4.447	109.61	110.29	
09/19/83	95	80.50	2ND PASS	59.3	18.0	2.5	212.42	.084	10.674	.050	.112	3.269	114.00	114.70	
09/19/83	95	80.50	3RD PASS	59.3	16.5	2.5	212.42	.077	9.784	.046	.102	3.913	114.00	114.70	
09/19/83	95	80.50	RUBBER 6609 1ST PASS	68.2	36.5	4.4	212.42	.171	24.893	.117	.262	3.657	64.77	65.17	
09/19/83	95	80.50	2ND PASS	68.2	23.5	2.5	212.42	.110	16.527	.075	.168	6.410	114.00	114.70	
09/19/83	95	80.50	3RD PASS	68.2	27.5	2.1	212.42	.129	18.755	.088	.197	6.930	135.71	136.55	
09/19/83	95	80.50	4TH PASS	68.2	23.0	2.5	212.42	.108	15.456	.073	.165	6.274	114.00	114.70	
09/19/83	95	80.50	5TH PASS	68.2	22.0	2.2	212.42	.103	15.004	.070	.157	6.020	129.54	130.36	
09/19/83	95	80.50	RUBBER 6600 1ST PASS	68.2	44.0	4.7	212.42	.207	30.008	.141	.315	7.144	67.85	68.27	
09/19/83	95	80.50	2ND PASS	68.2	24.8	2.7	212.42	.112	16.368	.077	.172	6.062	105.55	106.21	
09/19/83	95	80.50	3RD PASS	68.2	21.0	2.9	212.42	.092	14.322	.067	.150	4.938	98.27	98.88	
09/19/83	95	80.50	4TH PASS	68.2	31.0	3.0	212.42	.145	21.142	.099	.222	3.047	95.00	95.59	
09/19/83	95	80.50	5TH PASS	68.2	18.5	2.5	212.42	.087	12.617	.059	.132	5.046	114.00	114.70	
09/19/83	95	80.50	6TH PASS	68.2	26.5	2.5	212.42	.124	18.073	.085	.190	3.229	114.00	114.70	
09/19/83	95	80.50	7TH PASS	68.2	20.5	3.7	212.42	.096	12.981	.065	.147	4.660	95.00	95.59	
09/19/83	95	80.50	RUBBER 6609 1ST PASS	68.2	18.0	3.8	212.42	.084	12.276	.057	.129	2.116	49.13	49.44	
09/19/83	95	80.50	TALC PASS	68.2	18.5	12.7	212.42	.087	12.617	.059	.132	.993	22.44	22.58	
TOTALS					922.5	189.5	6585.02	.140	574.863						

TOTAL PASSES 31
 TOTAL PRODUCTION TIME 06:49:30
 AVERAGE LBS CEMENT PER PASS 29.756
 AVERAGE LBS CEMENT PER SQUARE YARD 4.342

Handwritten: 4.6

Handwritten: 4.3

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RUN ON 01/23/84 AT 14:11:22

DAVID M TOLUENE ANALYSIS

PAGE 3

DATE	LEN	WIDTH	DESCRIPTION	TOL	LBS	WINS	SO YDS	CEM SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL WIN	FEET MIN	F/FEET MIN	
09/16/83	75	45.75	RUBBER 1492 1ST PASS	56.4	15.0	5.7	95.31	.157	3.650	.088	.112	1.484	39.47	22.57	
09/16/83	75	45.75	2ND PASS	56.4	10.0	4.3	95.31	.104	5.640	.059	.075	1.311	52.32	29.92	
09/16/83	75	45.75	3RD PASS	56.4	6.5	3.3	95.31	.068	3.666	.038	.048	1.110	68.18	38.99	
09/16/83	75	45.75	RUBBER 7032 1ST PASS	58.8	16.5	6.1	95.31	.173	9.702	.101	.129	1.590	36.88	21.09	
09/16/83	75	45.75	2ND PASS	58.8	14.5	5.9	95.31	.257	14.406	.151	.192	2.441	38.13	21.80	
09/16/83	75	45.75	3RD PASS	58.8	18.5	5.3	95.31	.194	10.878	.114	.145	2.052	42.45	24.27	
09/16/83	75	45.75	4TH PASS	58.8	15.5	3.2	95.31	.162	9.114	.095	.121	2.048	70.31	40.20	
09/16/83	75	45.75	5TH PASS	58.8	12.0	2.8	95.31	.125	7.056	.074	.094	2.520	80.35	45.95	
09/16/83	75	45.75	6TH PASS	58.8	10.5	2.6	95.31	.118	5.174	.064	.082	2.374	86.63	49.48	
09/16/83	75	45.75	7TH PASS	58.8	19.5	4.0	95.31	.204	11.466	.120	.152	2.866	56.25	32.16	
09/16/83	75	45.75	8TH PASS	58.8	14.0	3.2	95.31	.146	8.232	.086	.109	2.572	70.31	40.20	
09/16/83	75	45.75	9TH PASS	58.8	16.5	4.2	95.31	.173	9.702	.101	.129	2.310	53.57	30.63	
09/16/83	75	45.75	10TH PASS	58.8	13.5	3.9	95.31	.141	7.938	.083	.105	2.035	57.69	32.99	
09/16/83	75	45.75	11TH PASS	58.8	9.0	3.6	95.31	.094	5.292	.055	.070	1.470	62.50	35.74	
09/16/83	75	45.75	12TH PASS	58.8	2.0	2.2	95.31	.230	2.936	.135	.172	3.080	63.57	36.63	
09/16/83	75	45.75	13TH PASS	58.8	17.0	3.1	95.31	.178	9.996	.104	.133	3.224	72.58	41.50	
09/16/83	75	45.75	14TH PASS	58.8	18.5	2.5	95.31	.194	10.878	.114	.145	3.351	80.00	51.46	
09/16/83	75	45.75	15TH PASS	58.8	11.5	2.4	95.31	.120	6.762	.070	.090	2.817	93.75	53.61	
09/16/83	75	45.75	16TH PASS	58.8	8.5	3.9	95.31	.089	4.998	.052	.066	3.281	57.69	32.99	
09/16/83	75	45.75	17TH PASS	58.8	9.5	3.2	95.31	.079	5.586	.058	.074	1.745	70.31	40.20	
09/16/83	75	45.75	RUBBER 6607 1ST PASS	64.8	7.5	3.5	95.31	.078	4.860	.050	.064	3.368	64.28	36.76	
09/16/83	75	45.75	2ND PASS	64.8	17.5	4.5	95.31	.187	11.340	.118	.151	2.520	50.00	28.59	
09/16/83	75	45.75	3RD PASS	64.8	15.0	4.0	95.31	.157	9.720	.101	.129	2.438	86.25	32.16	
09/16/83	75	45.75	4TH PASS	64.8	14.0	3.8	95.31	.146	9.872	.095	.121	2.387	57.21	35.86	
09/16/83	75	45.75	5TH PASS	64.8	9.0	3.3	95.31	.094	5.832	.061	.077	3.767	88.18	36.99	
09/16/83	75	45.75	6TH PASS	64.8	8.0	5.6	95.31	.083	5.164	.054	.066	.925	48.17	22.97	
09/16/83	75	45.75	7TH PASS	64.8	7.0	4.5	95.31	.073	4.536	.047	.060	1.888	68.00	28.59	
09/16/83	75	45.75	8TH PASS	64.8	7.5	3.4	95.31	.078	4.860	.050	.064	1.429	66.17	37.84	
09/16/83	75	45.75	TALC PASS	64.8	8.0	10.8	95.31	.083	5.184	.054	.069	4.480	28.83	31.91	
TOTALS					374.0	110.0	2668.68	.140	224.286						

TOTAL PASSES	28	WIN	2.4	WIN	.038	.048	.025	36.88	21.09
TOTAL PRODUCTION TIME	06:36:37	MAX	6.1	MAX	.151	.192	3.351	93.75	53.61
AVERAGE LBS CEMENT PER PASS	13.357	AVG	3.9	AVG	.083	.106	2.119	60.96	34.86
AVERAGE LBS CEMENT PER SQUARE YARD	3.924								

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RUN ON 01/23/84 AT 14:11:22

DAVID V TOLUENE ANALYSIS

PAGE 25

DATE	LEN	WIDTH	DESCRIPTION	X TOL	LBS	MINS	SO YDS	CEN SYD	TOLUENE USED	TOL SYD	TOL LYD	TOL MIN	FEET MIN	F/FEET MIN
GRAND TOTALS					19281.0	2984.1	126091.55		11995.663					

GRAND TOTAL PASSES 708
GRAND TOTAL AVERAGE LBS CEMENT PER PASS 27.233
GRAND TOTAL AVERAGE LBS CEMENT PER SQUARE YARD 152

$$\text{AVG} = \frac{\text{TOL}}{\text{MIN}} = \frac{11995.663}{2984.1} = 4.02$$

$$\text{AVG} = \frac{\text{TOL}}{\text{PASS}} = \frac{11995.663}{708} = 16.94$$

May 30, 1984

TO: R. H. Baddeley

FROM: M. A. Ware

The following three pages is the data you requested regarding David 'M' Company in Orlando. A detailed report covering same will follow at a later date.

M. A. Ware

MAW:bjw

SUMMARY

Date: May 25, 1984
 Roll Length: 50 yards (resurface blanket)
 Roll Width: 79"
 # Plys: 3-ply

Product Code: 4-Black Special
 Operator: Mack

Magnehelic Setting: .07-.08
 Duct Temp: 135°F
 Flow ACFM: 5698
 Flow SCFM: 5076

Pass	Rubber	% VOC	Cement Lbs/Pass	Toluene Lbs/Pass	Min/ Pass	Avg Lbs Tol/ Min	Ft/Min Tach	Ft/Min Calc	ppm Meter	% LEL Meter	% LEL Calc
1	B 1280	58.5%	9.5	Mixture of Toluene, MEK & Monochloroben- zene	3.3	N.A.	40-42	45.5	N.A.	N.A.	N.A.
2	B 1280		10.0	"	3.2	N.A.	44-45	46.9	N.A.	N.A.	N.A.
3	7002 (1)	58.7%	11.0	6.5	3.5	1.9	39-41	43.0	1313	10.3	10.8
4	7002		26.0	15.3	4.4	3.5	29-30	34.1	2000	15.7	19.8
5	7002		34.0	20.0	5.1	3.9	24-26	29.4	2313	18.2	22.1
6	7002		30.5	17.9	4.4	4.1	30-31	34.1	2625	20.7	23.2
7	7002		21.5	12.6	3.2	3.9	41-42	46.9	2625	20.7	22.1
8	7002		17.5	10.3	3.3	3.1	41-42	45.5	2625	20.7	17.6
9	7002		14.0	8.2	2.5	3.3	54-55	60.0	2627	20.7	18.7
10	7002		26.5	15.6	4.4	3.5	30-31	34.1	2625	20.7	19.8
11	7002		17.5	10.3	3.1	3.3	41-42	48.4	2625	20.7	18.7
12	7002		14.0	8.2	1.8	4.6	72-73	83.3	3250	25.6	26.0
13	7002		11.5	6.8	1.8	3.8	72-75	83.3	2938	23.2	21.5
14	7002		12.0	7.0	2.2	3.2	72-75	68.2	2938	23.2	18.1
15	7002 (1) Talc Pass		11.5	6.8	7.2	.94	Not on Tach	20.8	1125	8.9	5.3

Avg Lbs Tol/Hr: 42.0
Avg Lbs Tol/Pass: 11.2
Total Production Time: 3:27:45

Avg Min/Pass: 3.6

Avg Lbs/Min: 3.3
Avg % LEL: 19.2%

SUMMARY

Date: May 24, 1984
 Roll Length: 100 yards
 Roll Width: 62"
 # Plys: One

Product Code: One Ply Red
 Operator: Mack

Magnehelic Setting: .07-.08
 Duct Temp: 135°F
 Flow ACFM (a 135°F): 5698
 Flow SCFM (a 70°F): 5076

<u>Pass</u>	<u>Rubber</u>	<u>% VOC</u>	<u>Cement Lbs/Pass</u>	<u>Toluene Lbs/Pass</u>	<u>Min/ Pass</u>	<u>Avg Lbs Tol/ Min</u>	<u>Ft/Min Tach</u>	<u>Ft/Min Calc</u>	<u>ppm Meter</u>	<u>% LEL Meter</u>	<u>% LEL Calc</u>
1	7002	Tol 59.7%	57.0	34.0	10.37	3.28	29-30	28.9	2125	16.7	18.6
2	7002		40.5	24.2	10.3	2.35	29-30	28.9	1687	13.3	13.3
3	7002		40.5	24.2	10.7	2.26	29-30	28.0	1625	12.8	12.8
4	7002		28.0	16.7	9.1	1.83	34-36	33.0	1594	12.6	10.4
5	7002		28.0	16.7	9.0	1.86	29-36	33.3	1625	12.8	10.5
6	7002		25.5	15.2	7.8	1.95	40-42	38.5	1500	11.8	11.0
7	7002		20.0	11.9	7.5	1.59	40-43	40.0	1438	11.3	9.0
8	7002 (1) & (2)	59.8%	35.5	21.2	8.8	2.41	40-41	34.1	1625	12.8	13.6
9	7002 (2)		31.0	9.9	5.8	1.71	55-56	52.0	2313	18.2	9.7
10	7002 (2)		31.0	18.5	5.6	3.30	54-55	53.6	2000	15.7	18.7
11	3303 (1) Face	69.7%	25.5	17.8	8.9	2.00	34-36	33.7	2125	16.7	11.3
12	3303 (1)		37.0	25.8	7.5	3.44	40-41	40.0	2438	19.2	17.9
13	3303 (1)		35.0	24.4	7.3	3.34	40-42	41.1	2313	18.2	18.4
14	3303 (1)		37.0	25.8	5.7	4.53	55-57	52.6	2314	18.2	25.7
15	3303 (1)		22.0	15.3	5.6	2.73	54-55	53.6	2314	18.2	15.5
16	3303 (1) Dry Pass		--	--	--	--	---	--	--	--	--
17	3303 (1) Talc Pass		27.0	18.8	13.6	1.38	Not on Tach Readout	--	1313	10.3	7.81

Avg Lbs Tol/Hr: 49.8
Avg Lbs Tol/Pass: 20.03
Total Production Time: 6:26:52

Avg Min/Pass: 8.35

Avg Lbs/Min: 2.5
Avg % LEL: 14.9%

SUMMARY

Date: May 23, 1984
 Roll Length: 100 yards
 Roll Width: 82"
 # Plys: 4-ply conv.

Product Code: 2 Green Spec.
 Operator: Mack

Magnehelic Setting: .07-.08
 Duct Temp: 135°F
 Flow ACFM (a 135°F): 5698
 Flow SCFM (a 70°F): 5076

Pass	Rubber	% VOC	Cement Lbs/Pass	Toluene Lbs/Pass	Min/ Pass	Avg Lbs Tol/ Min	Ft/Min Tach	Ft/Min Calc	ppm Meter	% LEL Meter	% LEL Calc
1	1492-Subface	Tol/MEK 52.3%	29.5	15.4	5.8	2.66	56-57	51.7	1625	12.8	15.0
2	1492		21.0	11.0	6.0	1.83	54-56	50.0	1500	11.8	10.4
3	7002 (1)	Tol 60.1%	36.5	21.9	6.3	3.48	45-50	47.6	2375	18.7	19.7
4	7002 (1)		43.5	26.1	7.3	3.58	40-43	41.0	2625	20.6	20.27
5	7002 (1)		37.0	22.2	4.9	4.53	64-73	61.2	2625	20.6	25.7
6	7002 (1)		53.5	32.1	7.2	4.46	40-42	41.7	2875	22.6	25.3
7	7002 (1)		45.5	27.3	6.2	4.40	54-55	48.4	3250	25.6	24.9
8	7002 (1)		64.0	38.5	7.5	5.13	40-43	40.0	3000	23.6	29.0
9	7002 (2)		47.5	28.5	7.3	3.90	41-42	41.0	3000	23.6	20.4
10	7002 (2)		51.5	31.0	5.8	5.34	53-56	51.7	3260	25.7	30.2
11	7002 (2)		50.5	30.4	7.4	4.10	40-42	40.5	3000	23.6	23.2
12	7002 (2)		50.0	30.1	4.9	6.14	62-67	61.2	3260	25.7	34.8
13	7002 (2)		37.0	22.2	4.3	5.16	70-75	69.7	3260	25.7	29.2
14	2202-Face Green	Tol 67.0%	40.5	27.1	11.6	2.34	24-25	25.8	2625	20.6	13.3
15	2202		40.5	27.1	9.8	2.77	29-30	30.6	2625	20.6	15.7
16	2202		45.5	30.4	7.1	4.28	40-43	42.2	3000	23.6	24.2
17	2202		36.5	24.5	5.7	4.30	54-56	52.6	3000	23.6	24.3
18	2202		54.0	36.2	8.4	4.31	35-37	35.7	3000	23.6	24.4
19	2202		38.0	25.5	7.1	3.59	40-42	42.2	3000	23.6	20.3
20	2202		23.0	15.4	6.8	2.26	40-42	44.1	2125	16.7	12.8
21	2202		28.0	18.8	7.0	2.68	40-42	42.8	(No Reading)		15.2
22	2202-Talc Pass		23.0	15.4	13.3	1.16	---	22.6	1312	10.33	6.57

Avg Lbs Tol/Hr: 73.1
Avg Lbs Tol/Pass: 25.32
Total Production Time: 7:37:00

Avg Min/Pass: 7.2

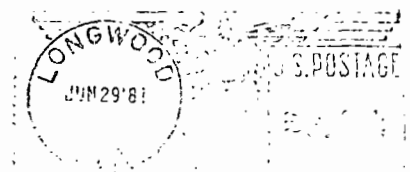
Avg Lbs/Min: 3.51
Avg % LEL: 20%

DAVID M COMPANY

201 VALENTINE WAY

LONGWOOD, FLORIDA 32750

Case No. 84-0044



CERTIFIED
P 211 559 779
MAIL

Department of Environmental Regulation
Bureau of Air Quality Management
Twin Towers Office Building
2600 Blair Stone Road
Tallahassee, FL 32301-8241

DER
JUL 2 1984
BAQM

P 408 530 377

RECEIPT FOR CERTIFIED MAIL

NO INSURANCE COVERAGE PROVIDED—
NOT FOR INTERNATIONAL MAIL

(See Reverse)

PS Form 3800, Feb. 1982

Sent to	
Mr. Michael A. Ware	
Street and No.	
P.O., State and ZIP Code	
Postage	\$
Certified Fee	
Special Delivery Fee	
Restricted Delivery Fee	
Return Receipt Showing to whom and Date Delivered	
Return Receipt Showing to whom, Date, and Address of Delivery	
TOTAL Postage and Fees	\$
Postmark or Date	
11/23/83	

PS Form 3811, Jan. 1979

● SENDER: Complete items 1, 2, and 3. Add your address in the "RETURN TO" space on reverse.

1. The following service is requested (check one.)

Show to whom and date delivered.....¢

Show to whom, date and address of delivery.....¢

RESTRICTED DELIVERY

Show to whom and date delivered.....¢

RESTRICTED DELIVERY.

Show to whom, date, and address of delivery. \$ _____

(CONSULT POSTMASTER FOR FEES)

2. ARTICLE ADDRESSED TO:

Mr. Michael A. Ware
2010 Indiana Street
Racine, Wisconsin 53405

3. ARTICLE DESCRIPTION:

REGISTERED NO.	CERTIFIED NO.	INSURED NO.
	P408530377	

(Always obtain signature of addressee or agent)

I have received the article described above.

SIGNATURE Addressee Authorized agent

4. DATE OF DELIVERY

11/25/83

5. ADDRESS (Complete only if requested)

6. UNABLE TO DELIVER BECAUSE:

CERK'S INITIALS

POSTMARK
RACINE WISCONSIN
NOV 24 1983

STAR GPO : 1979-300-459

DEPARTMENT OF ENVIRONMENTAL REGULATION

TWIN TOWERS OFFICE BUILDING
2600 BLAIR STONE ROAD
TALLAHASSEE, FLORIDA 32301-8241



BOB GRAHAM
GOVERNOR

VICTORIA J. TSCHINKEL
SECRETARY

November 22, 1983

Mr. Michael A. Ware
Graphics Division Environmental
Coordinator
Wheelabrator-Frye Inc.
2010 Indiana Street
Racine, Wisconsin 53405

Dear Mr. Ware:

The Department has reviewed your November 1, 1983 reply to our letter requesting additional information to complete the application for permit to construct the air pollution control devices required at David "M" Company. As noted in your letter, some of the requested information is not available at this time. Until we receive:

- 1) a description of the proposed control equipment selected for the storage tanks and the estimated emissions from the storage tanks;
- 2) the equipment specifications and talc emissions from the proposed baghouse;
- 3) the specifications on the control device selected for the manufacturing equipment and its design emissions of criteria pollutants,

your application will be incomplete and we will be unable to process it. Your Overall Compliance Schedule shows some of this information may not be available until September 1, 1984.

The Department finds it normal permitting procedure is not appropriate because of the delay involved in selecting, purchasing and installing the control equipment needed to bring the emissions from David "M" Company into compliance with state and federal regulations. Therefore, the Department's Bureau of Air Quality Management staff will meet with the Enforcement Section and draft a Consent Order to address this situation. We tentatively plan to have a copy of the proposed Order to you during December, 1983. After review of the proposed Order, the Department would expect your Company to accept it or meet with us to resolve any differences that may exist.

Mr. Michael A. Ware
Page Two
November 22, 1983

If you have any questions on this matter, please call
Willard Hanks (904/488-1344) or Tom Bessa (305/894-7955).

Sincerely,



C. H. Fancy, P.E.
Bureau of Air Quality
Management

CHF/WH/s

cc: Tom Bessa, St. Johns River District

INTEROFFICE MEMORANDUM

For Routing To District Offices And/Or To Other Than The Addressee		
To: _____	Loctn.: _____	
To: _____	Loctn.: _____	
To: _____	Loctn.: _____	
From: _____	Date: _____	
Reply Optional []	Reply Required []	Info. Only []
Date Due: _____	Date Due: _____	

TO: Nancy Wright
THRU: Bill Thomas *BT*
FROM: Willard Hanks *wmh*
DATE: November 14, 1983
SUBJ: David "M" Company

Michael Ware, representing David "M" Company, replied to BAQM's September 2, 1983 "incompleteness letter" on November 1, 1983. He was unable to answer all the questions in our September 2 letter. His reply included an Overall Compliance Schedule (attached) which shows some of the information required to complete the application will not be available until after September 1, 1984.

Their reply included expected emissions from the controlled plant, the minimum efficiency of the control device, and a schedule of when the control device will be installed and in operation. A permit to construct cannot be drafted without more information from the Company.

Under these circumstances, it appears a "Consent Order" is needed. The Consent Order should include:

1. A requirement for reporting the status of each step listed in the Overall Compliance Schedule.
2. DER's option to "speed up" the schedule, if justified.
3. A requirement that air pollution control devices be installed on the storage tanks ASAP. They are minor sources of VOCs, and the selection of control equipment is nearer to being completed. The control equipment for the tanks would be installed without a permit to construct in this circumstance.
4. Limit operations of the plant to 16 hr/day.
5. Set a deadline for the application for permit to construct to be complete, based on their proposed schedule.
6. Set a deadline for the equipment to be installed and in operation.
7. Set a deadline for the application for permit to operate to be submitted.

Memorandum
Page Two
November 14, 1983

A letter needs to be sent to the Company by November 30, 1983 to officially confirm their application is still incomplete. A proposed draft letter is attached.

WH/ks

cc: Tom Bessa, St. Johns River District 2 sent copy 11/22/83

October 19, 1983

ESTIMATED
OVERALL COMPLIANCE SCHEDULE
DAVID M COMPANY VOC CONTROL PROJECT

PHASE

Data Accumulation	January 1, 1984
Spreader Enclosure - (1) For Temperature Evaluation	January 15, 1984
Process Modifications and Development	April 1, 1984
Methods Comparison/Analysis - Thermal Incineration Vs. Recovery	May 15, 1984
Final Specification Development	July 1, 1984
Final Specification/Quotations and Analysis	September 1, 1984
Appropriations Request Development and Approval	December 1, 1984
Oven Enclosure and Exhaust System Installation	March 1, 1985
Solvent Recovery Delivery and Installation	August 1, 1985
Operational Shakedown, Debugging and Compliance Testing	December 1, 1985



Wheelabrator-Frye Inc.

Graphic Supplies Division

7520 University

Des Moines, Iowa 50311

Tel. (515) 274-4901

November 1, 1983

Mr. Willard Hanks
Engineer
State of Florida
Department of Environmental Regulation
2600 Blair Stone Road
Twin Towers Building
Tallahassee, FL 32301

DER
NOV 02 1983
BAQM

Dear Mr. Hanks:

I'd like to take this opportunity to thank you, Bill Thomas and Nancy Wright for the courtesies extended myself and Herb Mycroft during the course of our recent meeting in Tallahassee. In accordance with our conversations at that time, please find enclosed additional data gathered pursuant to Mr. C.H. Fancy's request for same dated September 2, 1983.

Should you require further information and/or assistance in processing David "M" Company's air pollution permit application, please don't hesitate to contact me directly.

Sincerely,

Michael A. Ware

Michael A. Ware
Graphics Div. Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

MAW;bjw
Encl.

cc: Wayne Brady
Jerry Spangler
Herb Mycroft
Bob Baddeley
J. H. Gamble
T. J. Lucas

SUPPLEMENT TO DAVID 'M' COMPANY
APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCES

Prepared By: Michael A. Ware

1. What will be the maximum production and operation time (hour/day) of the plant?

The maximum uncontrolled emissions of the plant will be 300 tons/year based upon a two shift, 16 hours/day operation. The potential emissions incorporating an end of pipe control device capable of delivering 90% removal efficiency would approximate 30 tons/year. The 300 ton/year two shift operation is based upon the following historical information:

<u>YEAR</u>	<u>OPERATING HOURS/DAY</u>	<u>ESTIMATED % NORMAL BUSINESS VOLUME</u>	<u>ANNUAL EMISSIONS</u>
1982	10	90%	271 Tons
1983	16	80%	255 Tons (Annualized Estimate)

2. What percent of the maximum production was the emission and raw material data in the application based upon?

The application was based upon 80% of the historically derived 300 tons/year uncontrolled emission number and reflected current economic conditions which approximate 70-80% of the usual demand for lithographic printing blankets. Should blanket demand return to 100% of the normal level, 300 tons uncontrolled and/or 30 tons controlled would approximate annual emissions without subtracting rubber cement containing VOC's which are disposed of at Emille, Alabama.

3. Will solvents other than MEK and Toluene be used at this plant?

Yes, small amounts of Naptha and Isopropyl Alcohol will be used at this facility for the purpose of manually cleaning blankets during inspection. The additional fugitive type emissions from the use of these materials will approximate

5,000 lbs/yr	Ispropyl Alcohol
2,000 lbs/yr	Naptha

4. How many gallons per year of each solvent will be consumed at the maximum production of the plant?

Approximately 75,000 gallons Toluene and 9,000 gallons MEK will be used at the maximum production of the plant. Were the use of MEK to be discontinued in order to facilitate efficient recovery of Toluene, approximately 83,000 gallons of Toluene would be used annually at the maximum production of the plant.

5. How will the solvents be shipped to the plant? Submit a drawing showing the storage area and piping to the mixing area.

The solvents will be shipped via bulk tanker and Appendix I is a conceptualization of the storage area and piping to the mixing area.

6. a) How many storage tanks will be built at the plant?

Five, two for #2 diesel fuel, two for Toluene storage and one for MEK storage.

- b) What air pollution control equipment will be used to minimize VOC emissions from the tanks?

A coaxial vapor recovery system of the type 1 or 2 nature will be installed to minimize VOC emissions during off loading. In addition, an OPW 523, or 523-S pressure vacuum vent will be installed on each tank vent line to minimize VOC emissions during storage. Appendix II contains example information on coaxial vapor recovery and OPW 523 and 523-S vents like those proposed for installation.

- c) What will be emissions be from the tanks? Submit emissions calculation.

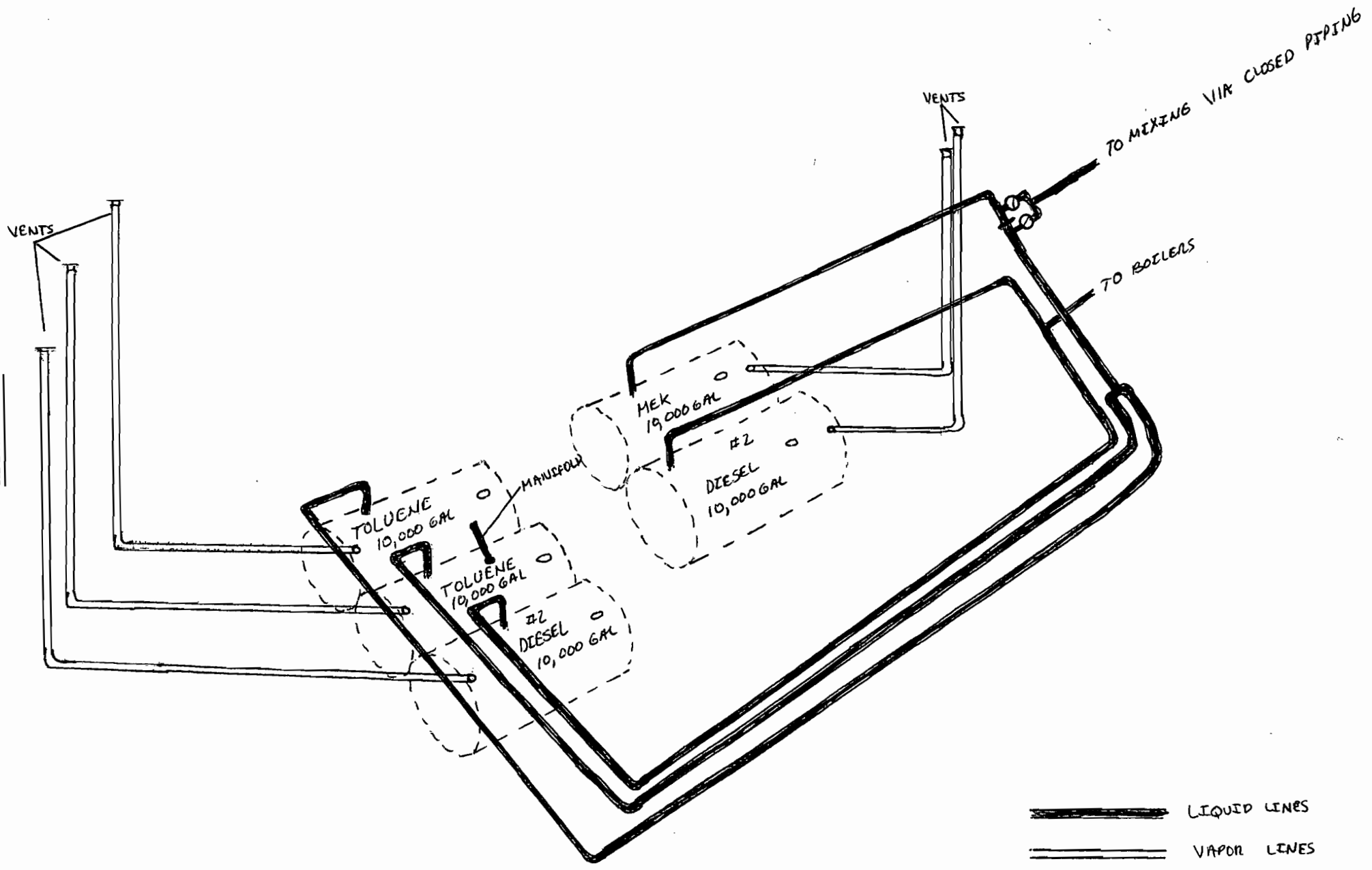
The equipment supplier has not as yet provided information concerning expected emissions from tanks equipped with coaxing recovery and pressure vents. We will endeavor to obtain this information as soon as possible.

7. What will be the mixing ratio of solvent to rubber in lbs/lb?

The chart below outlines the mixing ratio of solvent to rubber on a 100 lb. batch basis.

EXISTING SOLVENT STORAGE / DELIVERY SYSTEM

APPENDIX I



LIQUID LINES

VAPOR LINES

TANKS: UNDERGROUND
PLUMBING: UNDERGROUND TO PUMPS

COAXIAL

SECTION B

Coaxial Vapor Recovery Equipment FOR USE IN STAGE I AND STAGE II SYSTEMS

System C

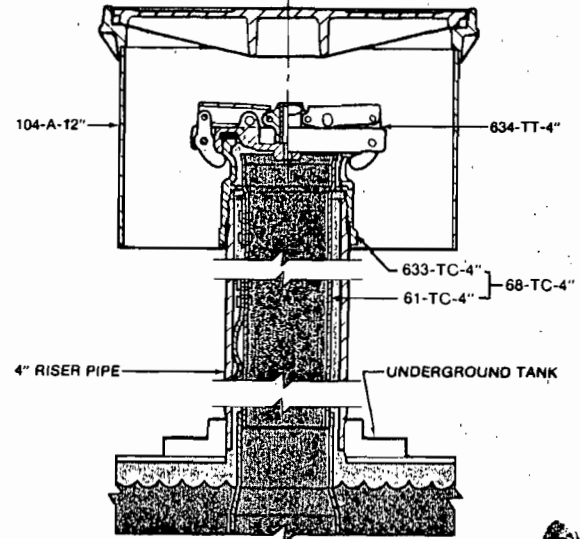
**TYPE 1 - 68-TC - 3" & 4" Series Fittings
(Non Poppeted)**

**TYPE 2 - 68-TCP - 3" & 4" Series Fittings
(Poppeted)**

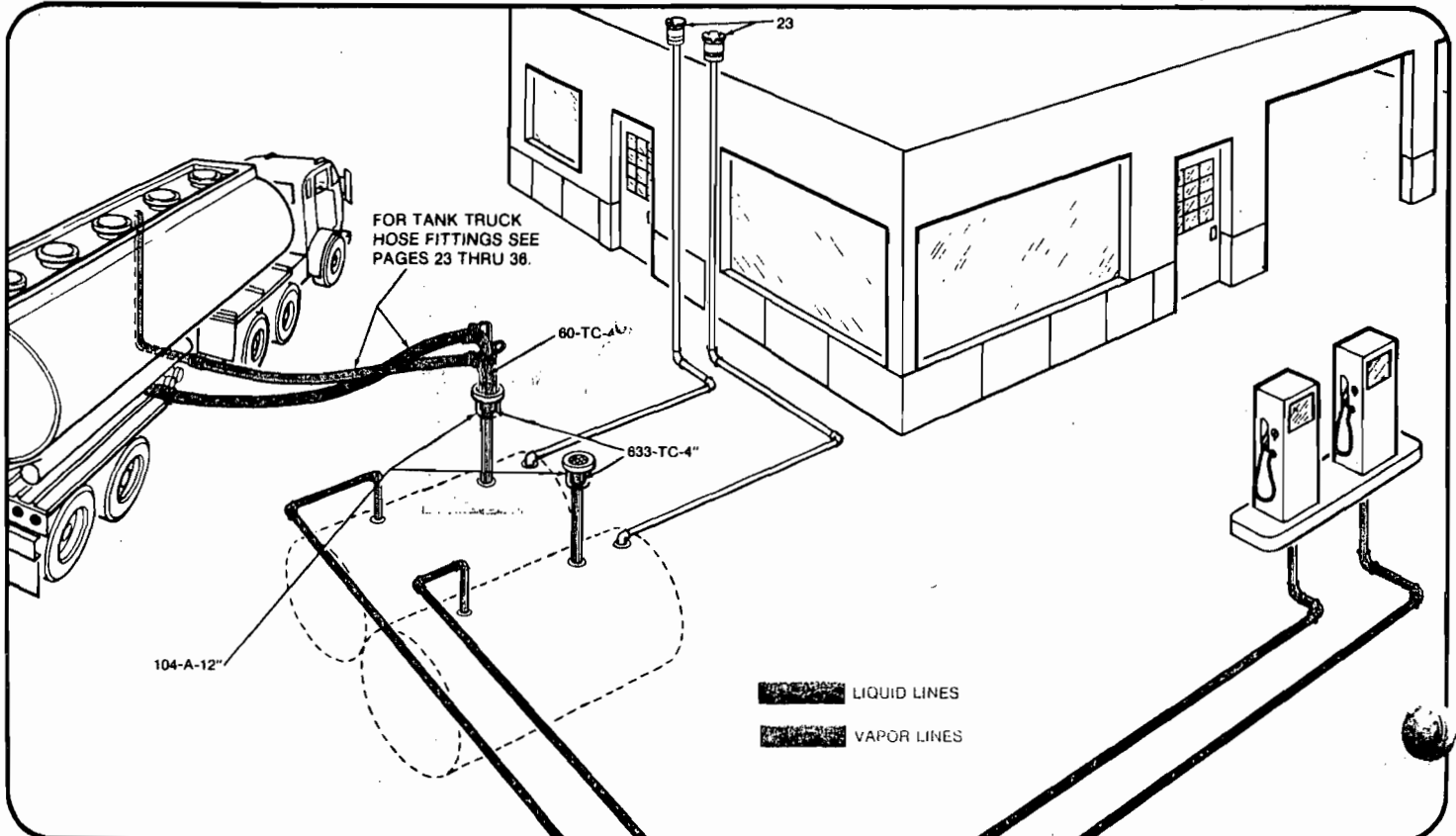
An easy, inexpensive means to convert existing conventional 3" & 4" size underground storage tank fill pipes to coaxial installations. Normally little to no digging or tearing up of concrete is necessary. Simply remove existing tight fill adaptor and fill pipe (if installation is so equipped), inserting Type 1 or Type 2 coaxial fill tube and adaptor as required.

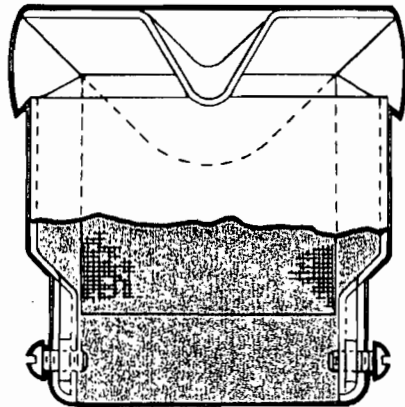
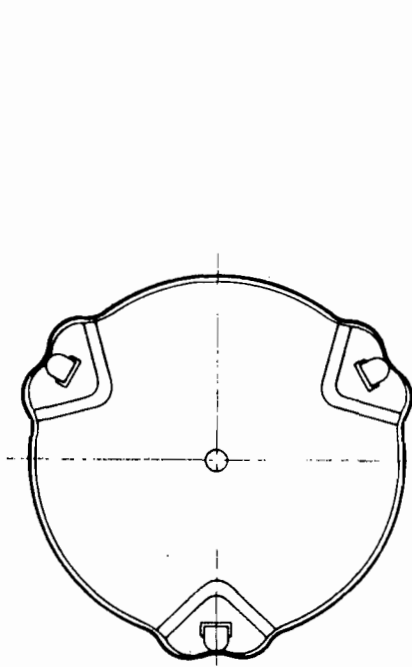
NOTE: Type 2 poppeted fittings are certified by California Air Resources Board (CARB)

SHOWN BELOW IS A TYPE 1 - 68TC-4" TYPICAL INSTALLATION

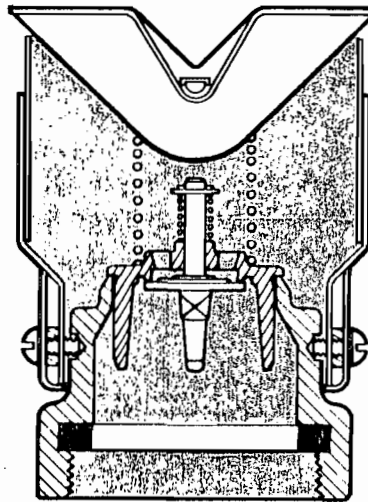
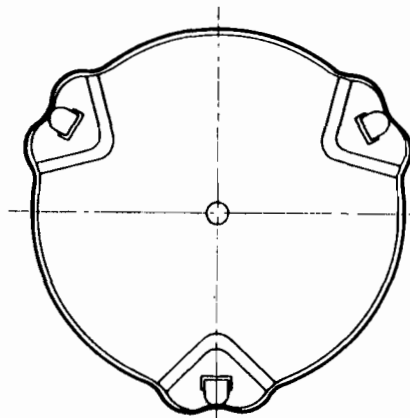


**TYPE 1 68-TC - 4" SERIES
(NON-POPPETED) FITTINGS**





OPW 23



OPW 523



OPW 523

OPW 23, 523, 523-S VENTS

The OPW 23 Vent has an aluminum body and a 40-mesh brass screen. The OPW 23 is an open vent and directs vapors upward in accordance with NFPA Code 30. Set screws make installation easy. Available in 1½", 2" and 3" sizes.

The OPW 523 Pressure Vacuum Vent is an upward vapor discharge vent and is available with either an 8 oz. or 12 oz. pressure setting and a ½ oz. vacuum setting.

The 523 is available in 2" size, and is attached to the 2" threaded vent pipe.

The OPW 523-S Pressure Vacuum Vent is similar to the 523 above, except it is attached to the vent line with set screws rather than pipe threads. A gasket is provided.

Available in 2" size.

The rated maximum flow pressure drop for all OPW 523 Vents is 28 oz. per square inch at 7000 SCFH.

<u>COMPOUND</u>	<u># SAMPLES ANALYZED</u>	<u>AVG LBS. TOLUENE/100 LBS.</u>	<u>AVG LBS RUBBER/ 100 LBS.</u>
7002	48	58.9	41.1
5503	22	63.7	36.3
1101	48	47.8	52.2
6607	40	63.4	36.6
1100	23	49.3	50.7
1492	20	61.2	38.8
6609	31	66.5	33.5
106E	1	63.3	36.7
7001	2	62.1	37.9
108	12	32.7	67.3
1280	12	57.1	42.9
106	9	65.6	34.4
105	13	33.8	66.2
3303	2	64.9	35.1
7007	7	62.0	38

Note: Encompasses all but 3 compounds currently used in production and would be a suitable cross section of those compounds used most frequently.

8. How many pounds of volatile organic compounds will be in a gallon of applied coating?

The chart below delineates lbs. solvent/gallon of applied coating for the most frequently used compounds.

<u>COMPOUND</u>	<u>HI</u>	<u>LO</u>	<u>AVERAGE</u>	<u>NUMBER TESTED</u>
7007	5.6	4.8	5.3	16
5503	6.1	5.4	5.7	8
6607	5.8	4.6	5.2	21
1492	5.4	4.6	5.2	4
3303	5.3	5.6	5.4	2
1280	5.1	4.9	5.0	2
1100	5.6	4.6	4.9	9
1101	5.0	4.3	4.7	16
105	4.1	3.8	3.9	9

* The above examples are typical of those compounds current in use, however, they do not include all compounds currently in use.

9. a) How will Talc be received, stored and conveyed to the process?

Talc is received, stored and conveyed to the process in 55 pound bags.

- b) What air pollution control equipment or operation practices will be used to minimize Talc emissions?

Appendix III contains a drawing of the bag house and ancillary piping and collectors to be used in the talcing operation for the purpose of dust collection.

- c) How much Talc will be emitted to the atmosphere?

Talc emissions will be fugitive since the collector will be located in-doors. The dust collector supplier is being contacted in order to ascertain estimated collection and removal efficiencies for the purpose of a fugitive emissions estimate. We will transmit that information to you as soon as it becomes available.

10. Will solvent laden air discharged from the plant contain any particulates or oil mist?

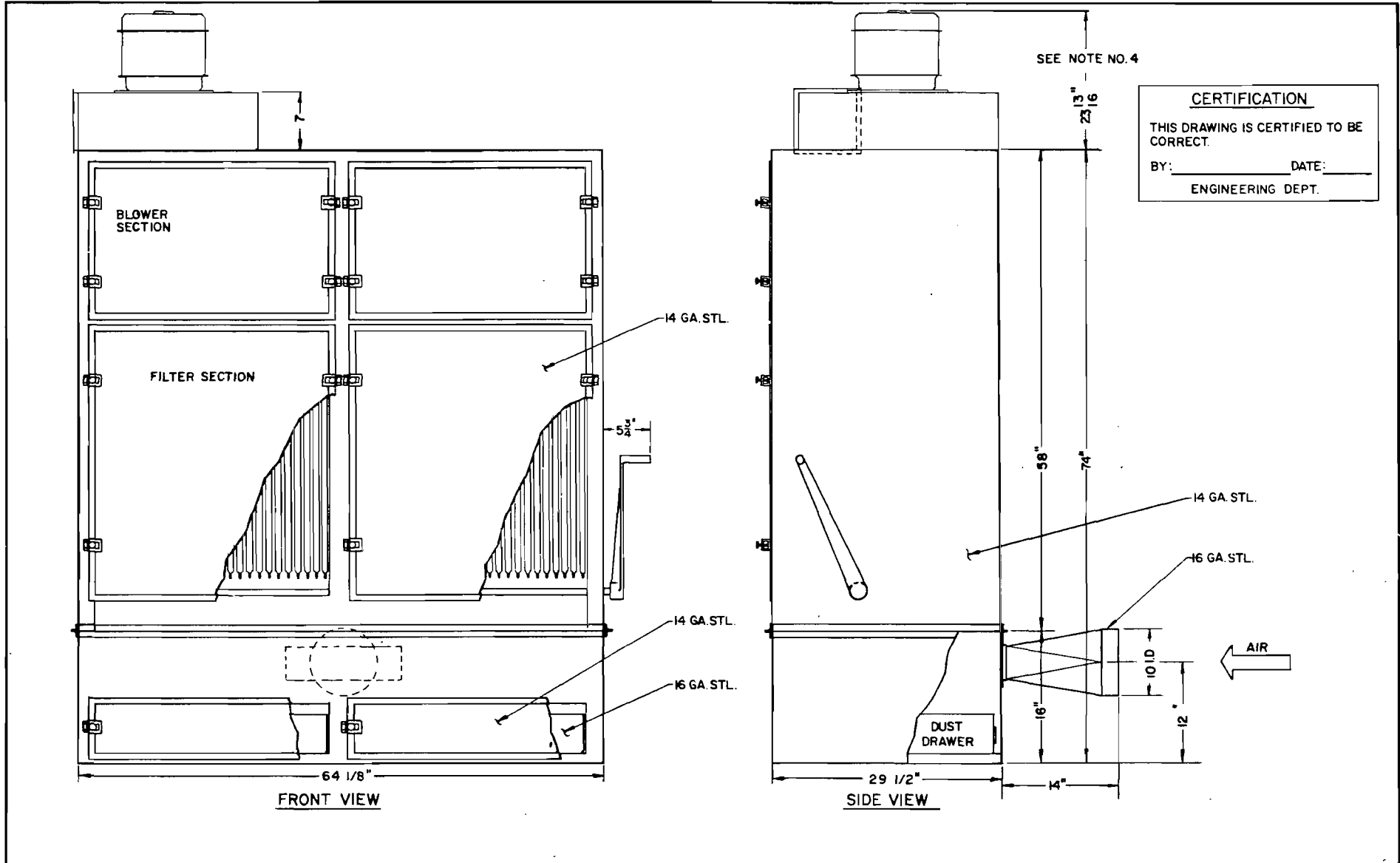
The solvent laden air will contain only fugitive particulates and no oil mist. Should solvent recovery be the selected control device, air discharged will be filtered at an efficient removal level of 99.0%+ of particles \geq .3 microns in diameter.

11. a) What will be the particulate and VOC emissions from emission point No. 2 mills?

Particulate and VOC emissions combined from the milling operation would approximate 1.7 - 8.0 lbs/day. These numbers were generated by weighing all the individual components to be added to a batch before milling and then weighing the total batch post milling and noting the lost material. The plasticizers used in the milling process are not appreciably volatile. Appendix IV contains several example data sheets on plasticizers commonly used in the milling operation and Appendix V contains volatility data on many plasticizers used in the rubber industry.

- b) What control equipment will be used to limit particulate emissions from the mills?

Since nearly all rubber compounds are compounded from master-batches, particulate emissions are quite low and as such no control device is currently proposed for the operation, see (11a) above.



CERTIFICATION
 THIS DRAWING IS CERTIFIED TO BE CORRECT.
 BY: _____ DATE: _____
 ENGINEERING DEPT.

....MULTIPLE RATING TABLES....

C.F.M.	F.P.M.	EXT.S.P.	C.F.M.	F.P.M.	EXT.S.P.
3625	6654	4.65	1912	3351	12.15
3365	6179	5.70*			
2995	5491	7.38			
2390	4386	9.95			

- SPECIFICATIONS**
1. MOTOR SHALL BE 10 HP ON NO. 124-20-5-10 3600 RPM, 230-460/60/3.
 2. DUST STORAGE CAPACITY SHALL BE 4.4 CU. FT.
 3. FILTER AREA SHALL BE 400 SQ. FT. (40 FILTERS 30" x 24" EACH.)
 4. NET WT. 1100 SHIPPING WT. 1475

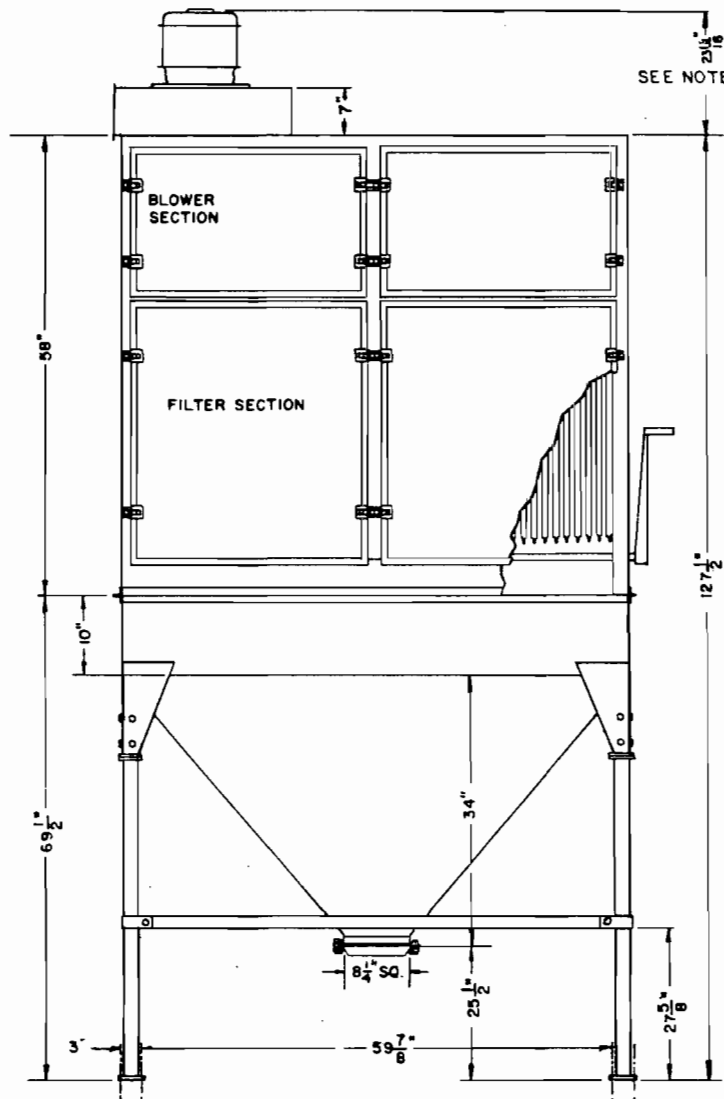
- NOTES**
1. THIS DRAWING IS FOR A 124-20-5-10 DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST.
 2. THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
 3. ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR & WHEEL ASS'Y
 4. MOTOR HEIGHT BASED ON STOCK, CAN VARY WITH MFRS.

TORIT DIVISION, DONALDSON COMPANY, INC.
 P.O. BOX 1299 - MINNEAPOLIS, MINNESOTA 55440

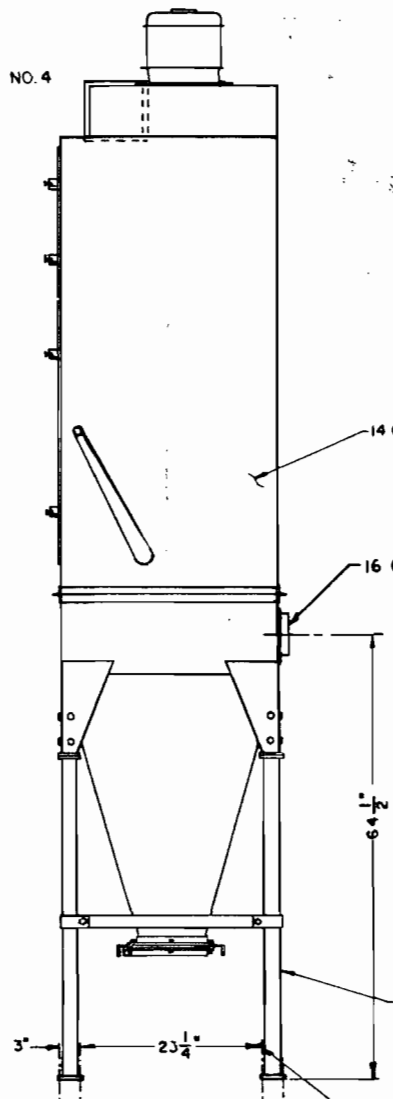
SPECIFICATION CONTROL DRAWING
NO. 124-20-5-10 DUST COLLECTOR

DRAWN: T. H. [Signature]
 APP'D: [Signature]

DRWG. NO: **124-020**
 REV. E 2-25-72

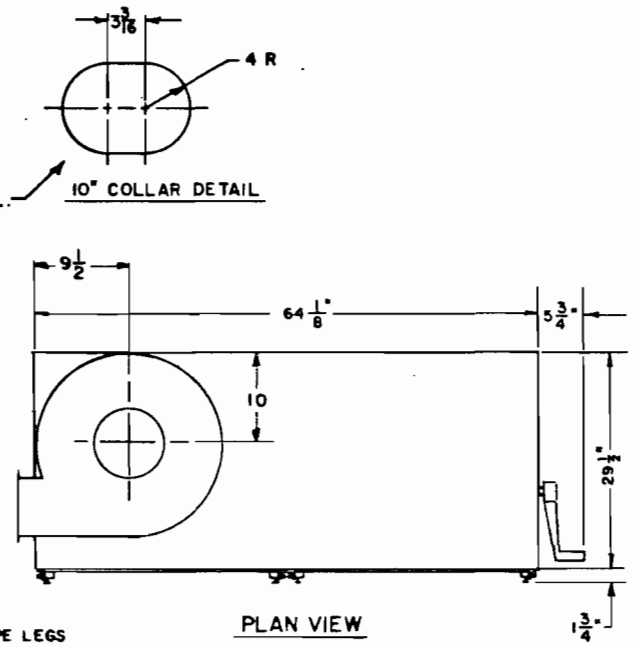


FRONT VIEW



SIDE VIEW

CERTIFICATION
 THIS DRAWING IS CERTIFIED TO BE CORRECT.
 BY: _____ DATE: _____
 ENGINEERING DEPT.



PLAN VIEW

LOCATING DIMS. FOR 9/16" DIA. MTG. HOLES.

MULTIPLE RATING TABLES

C.F.M.	F.P.M.	EXT. S.P.	C.F.M.	F.P.M.	EXT. S.P.
3625	6654	4.65	1912	3351	12.15
3365	6179	5.70 *			
2995	5491	7.38			
2390	4386	9.95			

SPECIFICATIONS

- MOTOR SHALL BE 10 HP
3500 RPM, 230-460/60/3.
- HOPPER CAPACITY SHALL BE 160 CU. FT.
- FILTER AREA SHALL BE 400 SQ. FT.
(40 FILTERS 30" X 24" EACH)
- INLET LOCATION: CENTERED ON BACK OR
CENTERED ON EITHER END
- NET WT. 1070 SHIPPING WT. 1365
1100 1405

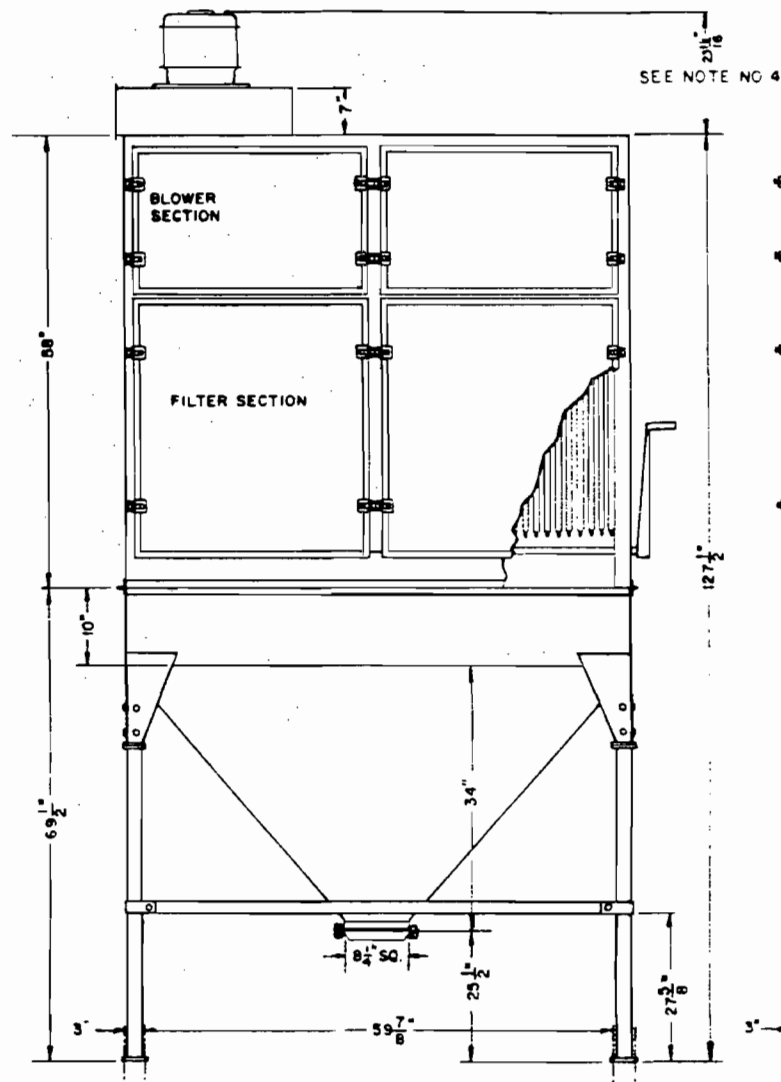
NOTES

- THIS DRAWING IS FOR A 124-20-5-10H DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST
- THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
- ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR AND WHEEL ASY
- MOTOR HEIGHT BASED ON STOCK CAN VARY WITH MFRS.

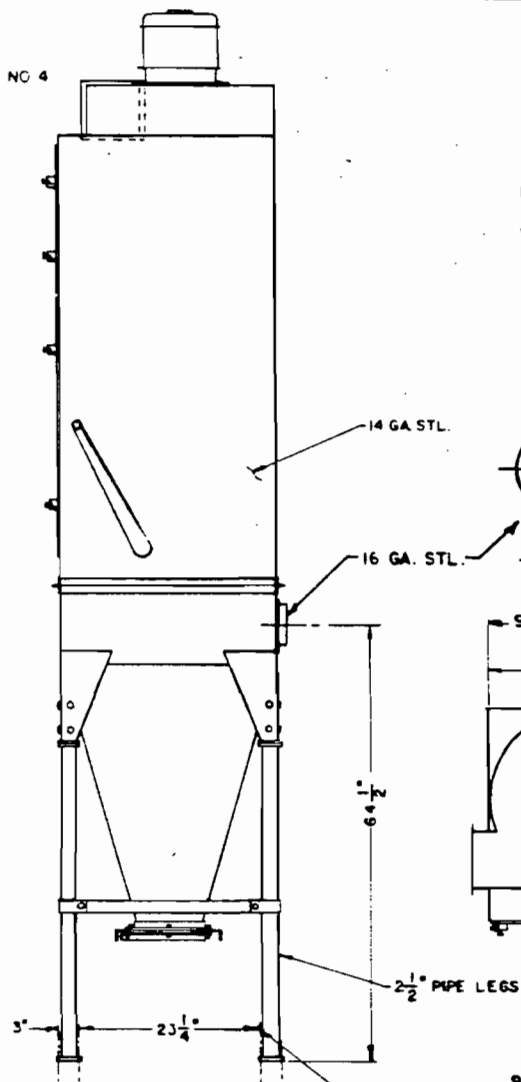
TORIT DIVISION, DONALDSON COMPANY, INC.
 P.O. BOX 1299 - MINNEAPOLIS, MINNESOTA 55440

SPECIFICATION CONTROL DRAWING
NO. 124-20-5-10H DUST COLLECTOR

DRAWN: T. Johnson DRWG. NO. 124-021
 APP'D: [Signature] REV. C 1-82

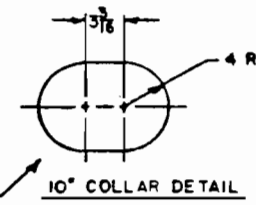


FRONT VIEW

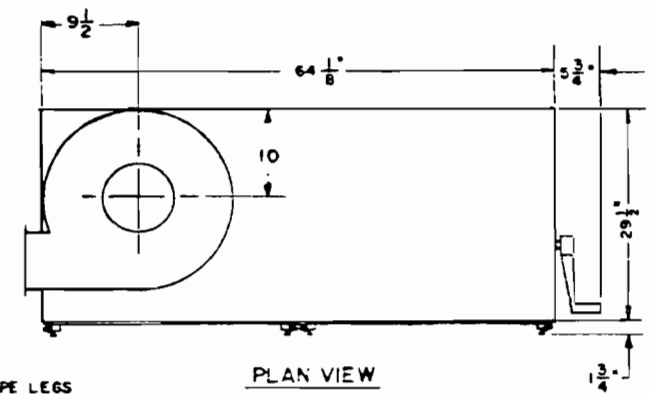


SIDE VIEW

CERTIFICATION
 THIS DRAWING IS CERTIFIED TO BE CORRECT.
 BY: _____ DATE: _____
 ENGINEERING DEPT.



10" COLLAR DETAIL



PLAN VIEW

MULTIPLE RATING TABLES					
C.F.M.	F.P.M.	EXT. S.P.	C.F.M.	F.P.M.	EXT. S.P.
3625	6654	4.65	1912	3351	12.15
3365	6179	5.70			
2995	5491	7.38			
2390	4386	9.95			

SPECIFICATIONS	
1. MOTOR SHALL BE 10 HP 3500 RPM, 230-460/60/3.	
2. HOPPER CAPACITY SHALL BE 160 CU. FT.	
3. FILTER AREA SHALL BE 400 SQ. FT. (40 FILTERS 30" X 24" EACH)	
4. INLET LOCATION, CENTERED ON BACK OF CENTERED ON EITHER END	
5. NET WT. 1070 SHIPPING WT. 1365	
1100	1405

- NOTES
1. THIS DRAWING IS FOR A 124-20-5-10H DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST.
 2. THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
 3. ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR AND WHEEL ASY.
 4. MOTOR HEIGHT BASED ON STOCK CAN VARY WITH MFRS.

TORIT DIVISION, DONALDSON COMPANY, INC.
 P.O. BOX 1299 - MINNEAPOLIS, MINNESOTA 55440

SPECIFICATION CONTROL DRAWING
NO. 124-20-5-10H DUST COLLECTOR

DRAWN: T.W.J. DRWG. NO: 124-021
 APP'D: [Signature] REV. C 1-83

- phenylacetone. Use: Polymerization inhibitor.
- diphenylbromoarsine** (C_6H_5)₂AsBr.
Properties: White crystals. M.p. 54–56°C.
Derivation: (a) Hydrobromic acid and diphenylarsinous oxide are heated together for about 4 hours at 115–120°C; (b) by action of arsenic tribromide on triphenyl arsine at 300–350°C.
Hazard: Highly toxic; strong irritant.
Shipping regulations: (Rail, Air) Arsenical compounds, solids, n.o.s.. Poison label.
- 1,3-diphenyl-2-buten-1-one.** See dypnone.
- diphenylcarbazide** (C_6H_5NHNH)₂CO.
Properties: White crystals or flakes. Insoluble in water; soluble in alcohol and benzene. M.p. 173°C.
Decomposes in light.
Derivation: Phenylhydrazine and urea.
Use: Determination of copper and other metals.
- diphenylcarbinol.** See benzhydrol.
- diphenyl carbonate** (C_6H_5O)₂CO.
Properties: White, crystalline solid. Can be halogenated and nitrated in characteristic manner. Readily undergoes hydrolysis and ammonolysis. Soluble in acetone, hot alcohol, benzene, carbon tetrachloride, ether, glacial acetic acid and other organic solvents; insoluble in water. B.p. 302°C; m.p. 78°C; sp. gr. 1.1215 (87/4°C).
Grade: Technical.
Uses: Plasticizer and solvent; synthesis of polycarbonate resins.
- diphenylchloroarsine** (C_6H_5)₂AsCl.
Properties: Colorless crystals or dark-brown liquid, which slowly becomes semi-solid. Decomposed by water (slowly). Soluble in carbon tetrachloride, chloropicrin, phenyldichloroarsine; practically insoluble in water. Sp. gr. 1.363 (40°C) (solid), or 1.358 (45°C) (liquid); b.p. 333°C (in CO₂ atmosphere); m.p. 41°C.
Derivation: Benzene and arsenic trichloride are heated in presence of aluminum chloride.
Grade: Technical.
Hazard: Highly toxic by inhalation; strong irritant to tissue.
Use: Military poison gas.
Shipping regulations: (Rail) Tear Gas label. (Air) Poison label. Not accepted on passenger planes.
- diphenyldecyl phosphite** (C_6H_5O)₂POC₁₀H₂₁.
Properties: Nearly water-white liquid; sp. gr. 1.023 (25/15.5°C); m.p. 18°C; refractive index 1.5160 (n_D²⁵/D). Combustible.
Uses: Chemical intermediate; stabilizer for polyvinyl and polyolefin resins.
- diphenyldichlorosilane** (C_6H_5)₂SiCl₂.
Properties: Colorless liquid; b.p. 305°C; f.p. –22°C; sp. gr. 1.19 (20°C); refractive index (n_D²⁵/D) 1.5713; flash point (COC) 288°F. Readily hydrolyzed by moisture, with liberation of hydrochloric acid. Combustible.
Derivation: (a) Reaction of powdered silicon and chlorobenzene in the presence of copper powder as catalyst; (b) reaction of phenylmagnesium chloride with silicon tetrachloride.
Grade: Technical.
Hazard: Highly toxic; strong irritant to tissue.
Use: Intermediate for silicone lubricants.
- diphenyldimethoxysilane** (C_6H_5)₂Si(OCH₃)₂.
Properties: Liquid; sp. gr. 1.080 (25°C); b.p. 191°C (53 mm); refractive index 1.5404 (25°C). Soluble in acetone, benzene, methyl alcohol; combustible.
Hazard: Probably toxic.
Uses: Treatment of powders, glass, paper, and fabrics.
- diphenyldimide.** See azobenzene.
- diphenyleneimine.** See carbazole.
- alpha-diphenylenemethane.** See fluorene.
- diphenylene oxide** (dibenzofuran) C₁₂H₈O (tricyclic).
Properties: Crystalline solid; m.p. 87°C; b.p. 288°C; insoluble in water; slightly soluble in alcohol, ether and benzene.
Derived from coal-tar.
Hazard: Probably toxic.
Use: Insecticide.
- 1,1-diphenylethane.** See uns-diphenylethane.
- 1,2-diphenylethane.** See sym-diphenylethane.
- uns-diphenylethane** (1,1-diphenylethane) (C_6H_5)₂CHCH₃.
Properties: Colorless liquid. Soluble in chloroform, ether, carbon disulfide. B.p. 286°C; sp. gr. 1.004 (20°C); f.p. –21.5°C. Flash point 264°F; combustible.
Derivation: Action of acetaldehyde on benzene in presence of concentrated sulfuric acid.
Uses: Solvent for nitrocellulose; organic synthesis.
- sym-diphenylethane** (bibenzyl; dibenzyl; 1,2-diphenylethane) C₆H₅CH₂CH₂C₆H₅.
Properties: White, crystalline needles or small plates. Soluble in alcohol, chloroform, ether, carbon disulfide; insoluble in water. Sp. gr. 0.9782; b.p. 284°C; m.p. 52°C.
Derivation: (a) By treating benzyl chloride with metallic sodium. (b) Action of benzyl chloride on benzylmagnesium chloride.
Use: Organic synthesis.
- diphenyl ether.** See diphenyl oxide.
- diphenylethylene.** See stilbene.
- N,N-diphenylethylenediamine** (ethyl diphenyldiamine) C₆H₅NHCH₂CH₂NHC₆H₅.
Properties: Cream-colored solid; sp. gr. 1.14; softening point 54°C; insoluble in water; soluble in acetone, ethylene dichloride, benzene, and gasoline.
Use: Antioxidant in rubber compounding.
- diphenylglycolic acid.** See benzilic acid.
- diphenylguanidine** (DPG; melaniline) HN:C(NHC₆H₅)₂.
Properties: White powder; bitter taste; slight odor; sp. gr. 1.13; m.p. 147°C; decomposes above 170°C; soluble in ethyl alcohol, carbon tetrachloride, chloroform, hot benzene and toluene; slightly soluble in water.
Derivation: Treatment of aniline with cyanogen chloride.
- diphenylmethane-4,4'-diisocyanate** (MDI; methylene-diparaphenylene isocyanate; methylenebis(phenyl isocyanate)) CH₂(C₆H₄NCO)₂.
Properties: Light-yellow, fused solid; solidification point 37°C; sp. gr. (70°C) 1.197; soluble in acetone, benzene, kerosine and nitrobenzene; combustible.
Derivation: para, para'-Diaminodiphenylmethane and phosgene.
Hazard: Highly toxic by inhalation of fumes. Strong irritant. Tolerance, 0.02 ppm in air.
Uses: Preparation of polyurethane resin and spandex fibers; bonding rubber to rayon and nylon.
- diphenylmethanol.** See benzhydrol.
- diphenylmethyl bromide** (benzhydryl bromide) BrCH(C₆H₅)₂.
Properties: Solid; m.p. 45°C; b.p. 193°C (26 mm). Decomposes in hot water; soluble in alcohol; very soluble in benzene.
Hazard: Strong irritant to eyes and skin.
Use: Organic synthesis.
Shipping regulations: (Air) Corrosive label. (Rail) Corrosive material; n.o.s.. White label.
- diphenylmethylchlorosilane** (C_6H_5)₂(CH₃)SiCl.
Properties: Colorless liquid; sp. gr. 1.107 (25°C); b.p. 295°C; flash point 135°F. Combustible.
Derivation: Grignard reactoin of diphenyldichlorosilane with methylmagnesium chloride.
Hazard: Moderate fire risk. Probably toxic.
Use: Intermediate; end stopper for silicone oils.
- diphenylnaphthylenediamine** C₁₀H₆(NHC₆H₅)₂.
Properties: Silvery, crystalline plates. Slightly soluble in alcohol; insoluble in water. M.p. 164°C.
Derivation: By heating 2,7-dihydroxynaphthalene with aniline and aniline hydrochloride.
Use: Organic synthesis.
- diphenylnitrosamine.** See N-nitrosodiphenylamine.
- 2,5-diphenyloxazole** (DPO) OOC₆H₄CC₆H₄.
Properties: White, fluffy solid; m.p. 70–72°C.
Grade: Scintillation.
- phenylacetone** C₆H₅CH₂COCH₃.
Use: Wavelength shifter in solution scintillation counting.
- diphenyl isophthalate** (DPIP) C₆H₄OOCC₆H₄COOC₆H₅.
Properties: White solid; m.p. 138–139°C. Combustible.
Use: Manufacture of polybenzimidazoles, high temperature-resistant polymers.
- diphenylketone.** See benzophenone.
- diphenylmethane** (benzylbenzene) (C_6H_5)₂CH₂.
Properties: Long colorless needles. Soluble in alcohol and ether; insoluble in water. Sp. gr. 1.0056; m.p. 26.5°C; b.p. 264.7°C. Flash point 266°F; combustible.
Derivation: Condensation of benzyl chloride and benzene in presence of aluminum chloride.
Hazard: Probably irritant and narcotic in high concentrations.
Uses: Organic synthesis; dyes; perfumery.
- diphenylmethane-4,4'-diisocyanate** (MDI; methylene-diparaphenylene isocyanate; methylenebis(phenyl isocyanate)) CH₂(C₆H₄NCO)₂.
Properties: Light-yellow, fused solid; solidification point 37°C; sp. gr. (70°C) 1.197; soluble in acetone, benzene, kerosine and nitrobenzene; combustible.
Derivation: para, para'-Diaminodiphenylmethane and phosgene.
Hazard: Highly toxic by inhalation of fumes. Strong irritant. Tolerance, 0.02 ppm in air.
Uses: Preparation of polyurethane resin and spandex fibers; bonding rubber to rayon and nylon.
- diphenylmethanol.** See benzhydrol.
- diphenylmethyl bromide** (benzhydryl bromide) BrCH(C₆H₅)₂.
Properties: Solid; m.p. 45°C; b.p. 193°C (26 mm). Decomposes in hot water; soluble in alcohol; very soluble in benzene.
Hazard: Strong irritant to eyes and skin.
Use: Organic synthesis.
Shipping regulations: (Air) Corrosive label. (Rail) Corrosive material; n.o.s.. White label.
- diphenylmethylchlorosilane** (C_6H_5)₂(CH₃)SiCl.
Properties: Colorless liquid; sp. gr. 1.107 (25°C); b.p. 295°C; flash point 135°F. Combustible.
Derivation: Grignard reactoin of diphenyldichlorosilane with methylmagnesium chloride.
Hazard: Moderate fire risk. Probably toxic.
Use: Intermediate; end stopper for silicone oils.
- diphenylnaphthylenediamine** C₁₀H₆(NHC₆H₅)₂.
Properties: Silvery, crystalline plates. Slightly soluble in alcohol; insoluble in water. M.p. 164°C.
Derivation: By heating 2,7-dihydroxynaphthalene with aniline and aniline hydrochloride.
Use: Organic synthesis.
- diphenylnitrosamine.** See N-nitrosodiphenylamine.
- 2,5-diphenyloxazole** (DPO) OOC₆H₄CC₆H₄.
Properties: White, fluffy solid; m.p. 70–72°C.
Grade: Scintillation.

types; curing and hardening of polymers (e.g., core-binding resins); corrosion inhibitor; propellant. Shipping regulations: (Rail) Flammable liquids, n.o.s., Red label. (Air) Flammable Liquid label.

triethylborane (triethylborine; boron triethyl). (C₂H₅)₃B.

Properties: Colorless liquid. Sp. gr. (25°C); flash point -32°F; f.p. -93°C; b.p. 95°C; refractive index 1.3971; heat of combustion 20,000 Btu/lb. Miscible with most organic solvents; immiscible with water.

Derivation: Reaction of triethylaluminum and boron halide, or diborane and ethylene.

Hazard: Flammable, dangerous fire risk. Ignites spontaneously in air. Highly toxic by inhalation; strong irritant. Reacts violently with water and oxidizing materials.

Uses: Igniter or fuel for jet and rocket engines; fuel additive; olefin polymerization catalyst; intermediate.

Shipping regulations: (Rail) Pyrophoric liquid, n.o.s., Red label. (Air) Not acceptable.

triethyl borate (ethyl borate) (C₂H₅)₃BO₃.

Properties: Colorless liquid; mild odor. Hydrolyzes rapidly, depositing boric acid in finely divided crystalline form. B.p. 120°C; sp. gr. 0.863-0.864 (20/20°C); flash point 51.8°F (C.C.); wt/gal 7.20 lb. (20°C); refractive index 1.37311 (20°C).

Uses: Antiseptics; disinfectants; antiknock agent.

Hazard: Flammable, dangerous fire risk. Moderately toxic.

Shipping regulations: (Rail) Flammable liquid, n.o.s., Red label. (Air) Flammable Liquid label. (Air) Legal label name, ethyl borate.

triethylborine. See triethylborane.

triethyl citrate (ethyl citrate) C₃H₅(COOC₂H₅)₃.

Properties: Colorless, mobile liquid. Bitter taste; b.p. 294°C; b.p. (1 mm) 126-127°C; sp. gr. 1.136 (25°C); refractive index 1.4405 (24.5°C); pour point -50°F; solubility in water 6.5 g/100 cc; solubility in oil 8 g/100 cc. Flash point 303°F (COC); combustible. Low toxicity.

Derivation: Esterification of citric acid.

Grades: Technical; refined; F.C.C.

Containers: Metal drums and cans; tank cars.

Uses: Solvent and plasticizer for nitrocellulose and natural resins; softener; paint removers; agglutinant; perfume base; food additive (not over 0.25%).

triethylenediamine N(CH₂CH₂)₃N. Catalyst used in the production of polyurethanes. Combustible.

triethylene glycol (TEG) HO(C₂H₄O)₃H.

Properties: Colorless, hygroscopic, practically odorless liquid. Sp. gr. 1.1254 (20/20°C); b.p. 287.4°C; vapor pressure less than 0.01 mm (20°C); flash point 350°F (C.C.); wt/gal 9.4 lb (20°C); freezing point -7.2°C; viscosity 0.478 poise (20°C). Autoignition temp. 700°F. Soluble in water; immiscible with benzene, toluene and gasoline. Combustible; low toxicity. Derivation: From ethylene and oxygen, as a by-product of ethylene glycol manufacture.

Grades: Technical; C.P.

Containers: 1-, 5-gal cans; 55-gal drums; tank cars.

Uses: Solvent and plasticizer in vinyl, polyester and polyurethane resins; dehydration of natural gas; humectant in printing inks; extraction solvent ("Udex" process).

triethylene glycol diacetate

CH₃COOCH₂CH₂OCH₂CH₂OCH₂CH₂COOCH₃. Properties: Colorless liquid. Sp. gr. (25°C) 1.112; re-

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fractive index n (25°C) 1.437; b.p. 300°C; f.p. less than -60°C. Combustible; low toxicity. Use: Plasticizer.

triethylene glycol dibenzoate

C₆H₅CO(OCH₂CH₂)₃OOCC₆H₅. Properties: Crystals; b.p. 210-223°C; m.p. 46°C; flash point 457°F (TOC); sp. gr. 1.168. Combustible. Low toxicity. Uses: Plasticizer for vinyl resins; adhesives.

triethylene glycol dicaprylate (triethylene glycol dioctoate) C₇H₁₅COO(CH₂CH₂O)₃OCC₇H₁₅.

Properties: Clear liquid; sp. gr. 0.973 (20°C); acidity 0.3% max. (caprylic); moisture 0.05% max; f.p. -3°C; b.p. 243°C (5 mm). Soluble in most organic solvents. Combustible; low toxicity. Uses: Low-temperature plasticizer for elastomers.

triethylene glycol dichloride. See triglycol dichloride.

triethylene glycol didecanoate

C₉H₁₉COO(C₂H₄O)₃OCC₉H₁₉. Properties: Colorless liquid. B.p. 237°C (2.0 mm); sp. gr. 0.9584 (20/20°C); viscosity 28.6 cp (20°C). Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol di(2-ethylbutyrate)

C₅H₁₁OCOCCH₂(CH₂OCH₂)₃CH₂OCOC₅H₁₁. Properties: Light-colored liquid; sp. gr. 0.9946 (20/20°C); 8.3 lb/gal (20°C); b.p. 196°C (5 mm); vapor pressure 5.8 mm Hg (200°C); solubility in water 0.02% by wt (20°C); viscosity 10.3 cp (20°C). Flash point 385°F. Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol di(2-ethylhexoate)

C₇H₁₅OCOCCH₂(CH₂OCH₂)₃CH₂OCOC₇H₁₅. Properties: Light-colored liquid; sp. gr. 0.9679 (20/20°C); 8.1 lb/gal (20°C); b.p. 219°C (5 mm); vapor pressure 1.8 mm Hg (200°C); insoluble in water; viscosity 15.8 cp (20°C). Flash point 405°F. Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol dihydroabietate

C₁₉H₃₁COO(C₂H₄O)₃OCC₁₉H₃₁. Properties: Liquid. Sp. gr. (25°C) 1.080-1.090; refractive index (20°C) 1.5180; vapor pressure (225°C) 2.5; flash point 226°C; insoluble in water. Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol dimethyl ether

CH₃(OCH₂CH₂)₃OCH₃. Properties: Water-white liquid; mild ether odor; sp. gr. (20/20°C) 0.9862; refractive index 1.4233 (n_D 20/D); flash point 232°F; b.p. (760 mm) 216.0°C; (100 mm) 153.6°C; f.p. -46°C. Autoignition temp. 1166°F. Completely soluble in water and hydrocarbons at 20°C. May contain peroxides. Combustible. Low toxicity. Containers: Glass bottles; cans; 55-gal drums. Uses: Solvent for gases; coupling immiscible liquids.

triethylene glycol dioctoate. See triethylene glycol dicaprylate.

triethylene glycol dipelargonate

C₈H₁₇COO(C₂H₄O)₃OCC₈H₁₇. Properties: Clear liquid; sp. gr. 0.964 (20/20°C); b.p. 251°C (5 mm); f.p. +1 to -4°C; refractive index 1.4470 (23°C); flash point 410°F. Almost insoluble in water; soluble in most organic solvents. Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol dipropionate

C₂H₅CO(OCH₂CH₂)₃OOCC₂H₅. Properties: Colorless liquid. Sp. gr. (25°C) 1.066; refractive index (25°C) 1.436; b.p. (2 mm) 138-142°C; f.p. less than -60°C; solubility in water, 6.70% by weight. Combustible. Low toxicity. Use: Plasticizer.

triethylene glycol monobutyl ether. See butoxytriethylene glycol.

triethylenemelamine (tretamine; TEM; 2,4,6-tris(1-aziridinyl)-s-triazine)

N[C]N(CH2)2N[C]N(CH2)2N[C]N(CH2)2. Properties: White, crystalline, odorless powder; m.p. 160°C (polymerizes); polymerizes readily with heat or moisture; soluble in alcohol, water, methanol, chloroform, and acetone. Grade: N.F.

Hazard: Highly toxic. Uses: Medicine (see nitrogen mustards); insecticide; chemosterilant.

triethylenephosphoramidate (tepa; tris-(1-aziridinyl)phosphine oxide; APO) (NCH₂CH₂)₃PO.

Properties: Colorless crystals; m.p. 41°C; soluble in water, alcohol and ether. Combustible. Derivation: From ethyleneimine.

Hazard: Highly toxic. Strong irritant to skin and tissue.

Uses: Medicine (see nitrogen mustards); insect sterilant. Also used with tetrakis(hydroxymethyl)phosphonium chloride (THPC) to form a condensation polymer suitable for flameproofing cotton. See also tris[1-(2-methylaziridinyl)phosphine oxide]. Shipping regulations: (Rail) White label. (Air) Corrosive label. Legal label name: tris(1-aziridinyl)phosphine oxide. See also tepa.

triethylenetetramine NH₂(C₂H₄NH)₂C₂H₄NH₂.

Properties: Moderately viscous yellowish liquid. Less volatile than diethylenetriamine but resembles it in many other properties. Soluble in water. B.p. 277.5°C; sp. gr. 0.9818 (20/20°C); m.p. 12°C; flash point 275°F (C.C.); wt 8.2 lb/gal (20°C). Combustible. Autoignition temp. 640°F.

Grades: Technical; anhydrous.

Containers: Cans; drums; tank cars.

Hazard: Strong irritant to tissue. Causes skin burns and eye damage.

Uses: Detergents and softening agents; synthesis of dyestuffs, pharmaceuticals and rubber accelerators.

Shipping regulations: (Air) Corrosive label.

tri(2-ethylhexyl)phosphate [C₈H₁₇CH(C₂H₅)CH₂]₃PO₄.

Properties: Light-colored liquid; sp. gr. 0.9260 (20/20°C); 7.70 lb/gal (20°C); b.p. 220°C (5 mm); vapor pressure 1.9 mm Hg (200°C); insoluble in water; viscosity 14.1 cp (20°C); pour point -74°C; flash point 405°F. Combustible; low toxicity. Use: Plasticizer.

tri(2-ethylhexyl) phosphite (C₈H₁₇O)₃P.

Properties: Straw-colored liquid; sp. gr. 0.897 (25/15°C); m.p., glass at low temperature; refractive index 1.451 (n_D 25/D); flash point 340°F (COC). Combustible; low toxicity. Uses: Plasticizer; intermediate.

tri(2-ethylhexyl) trimellitate C₆H₃(COOC₈H₁₇)₃.

Properties: Clear liquid, mild odor; sp. gr. 0.992 (20/

fractive index n (25°C) 1.437; b.p. 300°C; f.p. less than -60°C. Combustible; low toxicity.
Use: Plasticizer.

triethylene glycol dibenzoate

$C_{26}H_{30}O_6$; $CO(OCH_2CH_2)_3OOC_6H_5$.
Properties: Crystals; b.p. 210–223°C; m.p. 46°C; flash point 457°F (TOC); sp. gr. 1.168. Combustible. Low toxicity.
Uses: Plasticizer for vinyl resins; adhesives.

triethylene glycol dicaprylate (triethylene glycol dioctoate)

$C_{52}H_{100}O_{10}$; $COO(CH_2CH_2O)_3COOC_8H_{17}$.
Properties: Clear liquid; sp. gr. 0.973 (20°C); acidity 0.3% max. (caprylic); moisture 0.05% max; f.p. -3°C; b.p. 243°C (5 mm). Soluble in most organic solvents. Combustible; low toxicity.
Uses: Low-temperature plasticizer for elastomers.

triethylene glycol dichloride. See triglycol dichloride.

triethylene glycol didecanoate

$C_{46}H_{86}O_{10}$; $COO(C_2H_4O)_3OOC_9H_{19}$.
Properties: Colorless liquid. B.p. 237°C (2.0 mm); sp. gr. 0.9584 (20/20°C); viscosity 28.6 cp (20°C). Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol di(2-ethylbutyrate)

$C_{26}H_{48}O_8$; $OCOC_2H_5(CH_2OCH_2)_3CH_2OCOC_2H_5$.
Properties: Light-colored liquid; sp. gr. 0.9946 (20/20°C); 8.3 lb/gal (20°C); b.p. 196°C (5 mm); vapor pressure 5.8 mm Hg (200°C); solubility in water 0.02% by wt (20°C); viscosity 10.3 cp (20°C). Flash point 385°F. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol di(2-ethylhexoate)

$C_{30}H_{54}O_8$; $OCOC_2H_5(CH_2OCH_2)_3CH_2OCOC_2H_5$.
Properties: Light-colored liquid; sp. gr. 0.9679 (20/20°C); 8.1 lb/gal (20°C); b.p. 219°C (5 mm); vapor pressure 1.8 mm Hg (200°C); insoluble in water; viscosity 15.8 cp (20°C). Flash point 405°F. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol dihydroabietate

$C_{19}H_{31}O_6$; $COO(C_2H_4O)_3OOC_{19}H_{31}$.
Properties: Liquid. Sp. gr. (25°C) 1.080–1.090; refractive index (20°C) 1.5180; vapor pressure (225°C) 2.5; flash point 226°C; insoluble in water. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol dimethyl ether

$CH_3(OCH_2CH_2)_3OCH_3$.
Properties: Water-white liquid; mild ether odor; sp. gr. (20/20°C) 0.9862; refractive index 1.4233 (n 20/D); flash point 232°F; b.p. (760 mm) 216.0°C; (100 mm) 153.6°C; f.p. -46°C. Autoignition temp. 1166°F. Completely soluble in water and hydrocarbons at 20°C. May contain peroxides. Combustible. Low toxicity.
Containers: Glass bottles; cans; 55-gal drums.
Uses: Solvent for gases; coupling immiscible liquids.

triethylene glycol dioctoate. See triethylene glycol dicaprylate.

triethylene glycol dipelargonate

$C_{48}H_{86}O_{10}$; $COO(C_2H_4O)_3OOC_{17}H_{33}$.
Properties: Clear liquid; sp. gr. 0.964 (20/20°C); b.p. 251°C (5 mm); f.p. +1 to -4°C; refractive index 1.4470 (23°C); flash point 410°F. Almost insoluble in water; soluble in most organic solvents. Combustible. Low toxicity.

triethylene glycol dipropionate

$C_8H_{16}O_6$; $CO(OCH_2CH_2)_3OOC_3H_7$.
Properties: Colorless liquid. Sp. gr. (25°C) 1.066; refractive index (25°C) 1.436; b.p. (2 mm) 138–142°C; f.p. less than -60°C; solubility in water, 6.70% by weight. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol monobutyl ether. See butoxytriethylene glycol.

triethylenemelamine (tretamine; TEM; 2,4,6-tris(1-aziridinyl)-s-triazine)

$NC[N(CH_2)_2]NC[N(CH_2)_2]NC[N(CH_2)_2]$.
Properties: White, crystalline, odorless powder; m.p. 160°C (polymerizes); polymerizes readily with heat or moisture; soluble in alcohol, water, methanol, chloroform, and acetone.
Grade: N.F.
Hazard: Highly toxic.
Uses: Medicine (see nitrogen mustards); insecticide; chemosterilant.

triethylenephosphoramidate (tepa; tris-(1-aziridinyl)phosphine oxide; APO) $(NCH_2CH_2)_3PO$.

Properties: Colorless crystals; m.p. 41°C; soluble in water, alcohol and ether. Combustible.
Derivation: From ethylenimine.
Hazard: Highly toxic. Strong irritant to skin and tissue.
Uses: Medicine (see nitrogen mustards); insect sterilant. Also used with tetrakis(hydroxymethyl)phosphonium chloride (THPC) to form a condensation polymer suitable for flameproofing cotton. See also tris[1-(2-methyl)aziridinyl]phosphine oxide.
Shipping regulations: (Rail) White label. (Air) Corrosive label. Legal label name: tris(1-aziridinyl)phosphine oxide.
See also tepa.

triethylenetetramine $NH_2(C_2H_4NH)_3C_2H_4NH_2$.

Properties: Moderately viscous yellowish liquid. Less volatile than diethylenetriamine but resembles it in many other properties. Soluble in water. B.p. 277.5°C; sp. gr. 0.9818 (20/20°C); m.p. 12°C; flash point 275°F (C.C.); wt 8.2 lb/gal (20°C). Combustible. Autoignition temp. 640°F.
Grades: Technical; anhydrous.
Containers: Cans; drums; tank cars.
Hazard: Strong irritant to tissue. Causes skin burns and eye damage.
Uses: Detergents and softening agents; synthesis of dyestuffs, pharmaceuticals and rubber accelerators.
Shipping regulations: (Air) Corrosive label.

tri(2-ethylhexyl)phosphate $[C_8H_{17}CH(C_2H_5)CH_2]_3PO_4$.

Properties: Light-colored liquid; sp. gr. 0.9260 (20/20°C); 7.70 lb/gal (20°C); b.p. 220°C (5 mm); vapor pressure 1.9 mm Hg (200°C); insoluble in water; viscosity 14.1 cp (20°C); pour point -74°C; flash point 405°F. Combustible; low toxicity.
Use: Plasticizer.

tri(2-ethylhexyl) phosphite $(C_8H_{17}O)_3P$.

Properties: Straw-colored liquid; sp. gr. 0.897 (25/15°C); m.p., glass at low temperature; refractive index 1.451 (n 25/D); flash point 340°F (COC). Combustible; low toxicity.
Uses: Plasticizer; intermediate.

tri(2-ethylhexyl) trimellitate $C_6H_3(COOC_8H_{17})_3$.

Properties: Clear liquid, mild odor; sp. gr. 0.992 (20/

20°C); distillation range at 3 mm, 278–284°C (5–95%); f.p., a gel at -35°C; refractive index 1.4846 (23°C); wt/gal 8.26 lb (20°C). Combustible; low toxicity.
Use: Plasticizer.

triethylmethane. See 3-ethylpentane.

triethylmethyl malonaldehyde diacetal. See 1,1,3-triethoxy-3-methoxypropane.

triethyl orthoformate (triethoxymethane) $CH(OC_2H_5)_3$.

Properties: Colorless liquid; pungent odor; b.p. 145.9°C; refractive index 1.39218 (18.8°C); sp. gr. 0.895 (20/20°C). Soluble in alcohol, ether; decomp. in water. Flash point 86°F (C.C.). Low toxicity.
Derivation: Reaction of sodium ethylate with chloroform or reaction of hydrochloric acid with hydrogen cyanide in ethyl alcohol solution.
Containers: 55-gal steel drums.
Hazard: Flammable, moderate fire risk.
Uses: Organic synthesis; pharmaceuticals.

triethyl phosphate (TEP) $(C_2H_5)_3PO_4$.

Properties: Colorless, high-boiling liquid. Mild odor; very stable at ordinary temperatures. Compatible with many gums and resins. Soluble in most organic solvents; completely miscible in water. When mixed with water is quite stable at room temperature, but at elevated temperatures it hydrolyzes slowly. F.p. -56.4°C; b.p. 216°C. Flash point 240°F; refractive index 1.4055 (20°C); wt/gal 8.90 lb (68°F). Combustible.

Grades: Technical; 97%.

Containers: Drums; tank cars; tank trucks.

Hazard: May cause nerve damage but to less extent than other cholinesterase-inhibiting compounds.

Uses: Solvent; plasticizer for resins, plastics, gums; manufacture of pesticides; catalyst; lacquer remover.
Shipping regulations: (Rail, Air) Organic phosphate, liquid, n.o.s., Poison label. Not acceptable on passenger planes.

triethyl phosphite $(C_2H_5)_3PO_3$.

Properties: Colorless liquid; sp. gr. 0.9687 (20°C); b.p. 156.6°C; refractive index 1.413 (n 25/D); flash point 130°F; insoluble in water; soluble in alcohol and ether. Combustible.

Containers: Glass bottles; 5-, 55-gal drums.

Hazard: Moderate fire risk. May be toxic.

Uses: Synthesis; plasticizers; stabilizers; lube and grease additives.

O,O,O-triethyl phosphorothioate (triethyl thiophosphate) $(C_2H_5O)_3PS$.

Properties: Colorless liquid with characteristic odor. B.p. (10 mm) 93.5–94°C; sp. gr. 1.074; flash point 225°F (COC); combustible.

Containers: Bottles; drums.

Hazard: Toxic by ingestion; cholinesterase inhibitor.
Uses: Plasticizer; lubricant additive; antifoam agent; hydraulic fluid; intermediate.

Shipping regulations: (Rail, Air) Organic phosphate, liquid, n.o.s., Poison label. Not acceptable on passenger planes.

triethyl tricarballylate $(C_2H_5OCOCH_2)_3CHCOOC_2H_5$.

Properties: Colorless liquid. Sp. gr. (20°C) 1.087; refractive index (26°C) 1.4234; b.p. (5 mm) 158–160°C; solubility in water (20°C) 0.62% by weight. Combustible.

Use: Plasticizer.

ner intermediate; sol- parts and precision

ethyl)trimethylammo-
(OH)₃; C₆H₅O₇.
ion); drums.

chloroethylene (q.v.).

cobaltic oxide.

tricyclamol chloride

soluble in alcohol; in-
9 (48°C); b.p. 234°C
ible.

ners.

OOH. A saturated
in natural fats or oils.
crystalline solid; m.p.
ed in medical research
as chromatography.

C₆H₄)₃BO₃.
sp. gr. 1.065 (25°C);
x 1.5480 (24°C); flash
in all proportions in
hydrolyzes on contact

sis.

phate; TCP)
of isomers.
, odorless liquid. Sta-
refractive index 1.556
); wt/gal 9.7 lb; crys-
Miscible with all the
s, also with vegetable
point 437°F; autoigni-

osphorus oxychloride.
, tank cars.
ngestion and skin ab-
highly toxic; its toler-
of air.

chloride, polystyrene,
for plastics; air filter
waterproofing; additive
s; hydraulic fluid and

p.
slight phenolic odor.
(20/4°C) 1.115; flash
ible in water; miscible
, ether, and kerosine.

for plastics and resins.

H₂N · CH₃Cl. 1-Cyclo-

der; faint characteristic odor; soluble in alcohol and
in water.
Use: Medicine.

tricyclic. An organic compound comprised of three
(only) ring structures, which may be the same or dif-
ferent, e.g., anthracene.

sym-tricyclodecane. See adamantane.

tricyclohexyl borate. See boric acid ester.

n-tridecane CH₃(CH₂)₁₁CH₃.

Properties: Colorless liquid. Soluble in alcohol; insol-
uble in water. Sp. gr. 0.755 (20/4°C); b.p. 225.5°C;
f.p. -5.45°C; refractive index 1.4250 (20/D); flash
point 175°F. Combustible; low toxicity.

Grades: 95%; 99%; research.

Containers: Glass bottles; 1-, 5-gal drums.

Uses: Organic synthesis; distillation chaser.

n-tridecanoic acid (tridecyclic acid; tridecoic acid)
CH₃(CH₂)₁₁COOH. A saturated fatty acid usually
prepared synthetically.

Properties: Colorless crystals; m.p. 44.5°C; sp. gr.
0.8458 (80/4°C); b.p. 312.4°C, 192.2°C (16 mm); re-
fractive index 1.4328 (50°C). Slightly soluble in wa-
ter; soluble in alcohol and ether. Combustible; low
toxicity.

Grade: 99% pure.

Uses: Organic synthesis; medical research.

tridecanol. See tridecyl alcohol.

tridecoic acid. See n-tridecanoic acid.

tridecyl alcohol (tridecanol). A commercial mixture of
isomers of the formula C₁₂H₂₅CH₂OH.

Properties: Low-melting white solid with pleasant
odor; b.p. 274°C; m.p. 31°C; sp. gr. (20/20°C)
0.845; wt/gal 7.0 lb; flash point (TCC) 180°F. Com-
bustible; low toxicity.

Derivation: Oxo process (q.v.) from C₁₅ hydrocarbons.

Grade: Technical.

Containers: 55-gal drums; tank cars.

Uses: Esters for synthetic lubricants; detergents; anti-
foam agent; other tridecyl compounds; perfumery.

tridecylbenzene (1-phenyltridecane) C₆H₅(CH₂)₁₁CH₃.

Properties: Colorless liquid; sp. gr. 0.85-0.86 (60/
60°F); refractive index 1.4815-1.4830. Combustible.
Use: Detergent intermediate.

tridecyllic acid. See n-tridecanoic acid.

tri(decyl) orthoformate CH(OC₁₀H₂₁)₃.

Properties: Liquid; b.p. 194°C; f.p. -15 to -20°C;
refractive index 1.448; insoluble in water; soluble in
benzene, naphtha, ether, and alcohol.

Use: To remove small quantities of water from ethers
or other solvents where acid catalysts can be em-
ployed.

tri(decyl) phosphite (C₁₀H₂₁O)₃P.

Properties: Water-white liquid; decyl alcohol odor;
sp. gr. 0.892 (25/15.5°C); m.p. less than 0°C; re-
fractive index 1.4565 (25°C). Flash point 455°F.
Combustible.

Containers: 55-gal drums.

Uses: Chemical intermediate; stabilizer for polyvinyl
and polyolefin resins.

tridihexethyl chloride

C₆H₁₁(C₆H₅)(OH)CH₂CH₂N(C₂H₅)₂ · C₂H₅Cl. (3-
Diethylamino-1-phenyl-1-cyclohexylpropanol etho-

form, and in alcohol. Practically insoluble in ether
and in acetone. Melting range 198-202°C.

Grade: N.F.

Use: Medicine.

2,4,6-tri(dimethylaminomethyl)phenol

[(CH₃)₂NCH₂]₃C₆H₂OH.

Properties: Liquid; refractive index 1.5181. Combust-
ible.

Hazard: May be toxic.

Uses: Antioxidants, acid neutralizers, stabilizers, and
catalysts for epoxy and polyurethane resins.

tri(dimethylphenyl)phosphite (trixylenyl phosphate)
[(CH₃)₂C₆H₄O]₃PO.

Properties: Liquid. Sp. gr. 1.155; refractive index
1.5535; b.p. (10 mm), 243-265°C; flash point 450°F;
solubility in water (85°C), 0.002% by weight. Com-
bustible.

Use: Plasticizer.

"**Tridione.**"³ Trademark for trimethadione, U.S.P.

tridodecyl amine. See trilauryl amine.

tridodecyl borate. See boric acid ester.

tridymite SiO₂. A vitreous, colorless or white, native
form of pure silica. Found variously but not so com-
monly as quartz (q.v.). Quartz will change into tridy-
mite with a 16.2% increase in volume at 870°C.
Unlike quartz, it is soluble in boiling sodium car-
bonate solution. Sp. gr. 2.28-2.3; Mohs hardness 7.

trietazine. Generic name for 2-chloro-4-diethylamino-
6-ethylamino-s-triazine ClC₂N₃[N(C₂H₅)₂]NHC₂H₅.

Properties: Solid; practically insoluble in water.

Uses: Herbicide; plant growth regulator.

"**Tri-Ethane.**"¹⁷⁷ Trademark for 1,1,1-trichloroethane
(q.v.).

triethanolamine (TEA; tri(2-hydroxyethyl)amine)
(HOCH₂CH₂)₃N.

Properties: Colorless, viscous, hygroscopic liquid
with slight ammoniacal odor; m.p. 21.2°C; b.p.
335°C (dec); vapor pressure < 0.01 mm (20°C); sp.
gr. 1.126; flash point (open cup) 375°F; wt/gal 9.4
lb; miscible with water, alcohol; soluble in chloro-
form; slightly soluble in benzene and ether; slightly
less alkaline than ammonia. Commercial product
contains up to 25% diethanolamine and up to 5%
monoethanolamine. Combustible; low toxicity.

Derivation: Reaction of ethylene oxide and ammonia.

Grades: Technical; regular; 98%; U.S.P.

Containers: Drums; tank cars.

Uses: Fatty acid soaps used in drycleaning, cosmet-
ics, household detergents, and emulsions; wool
scouring; textile antifume agent and water-repellent;
dispersion agent; corrosion inhibitor; softening agent,
humectant, and plasticizer; insecticide; chelating
agent; rubber accelerator.

triethanolamine lauryl sulfate

(HOC₂H₄)₃NOS(O)₂OC₁₂H₂₅. A liquid or paste.

Containers: Drums, tank cars; tank trucks.

Uses: Detergent; wetting, foaming and dispersing
agent for industrial, cosmetic and pharmaceutical
applications, especially shampoos.

triethanolamine methanearsonate

CH₃As(O)[ONH(C₂H₅OH)]₂.

Hazard: Highly toxic by ingestion.

Use: Herbicide.

Shipping regulations: (Rail, Air) Arsenical compounds,
n.o.s. Poison label.

triethanolamine oleate. See trihydroxyethylamine ole-
ate.

triethanolamine stearate. See trihydroxyethylamine
stearate.

triethanolamine titanate. See titanium chelate.

1,1,3-triethoxyhexane

CH(OC₂H₅)₂CH₂CH(OC₂H₅)C₂H₅.

Properties: Colorless liquid; sp. gr. 0.8746 (20/20°C);
b.p. 133°C (50 mm); f.p. -100°C; wt/gal 7.3 lb;
flash point 210°F. Insoluble in water. Combustible.
Low toxicity.

Use: Synthesis of aldehydes, acids, esters, chloride,
amines, etc.

trithoxymethane. See triethyl orthoformate.

1,1,3-triethoxy-3-methoxypropane (triethylmethyl
malonaldehyde diacetal)

(CH₃O)(C₂H₅O)CHCH₂CH(OC₂H₅)₂.

Properties: Colorless liquid. Sp. gr. (25/4°C), 0.9300;
b.p. (6 mm) 86°C. Combustible.

Grade: 99%.

Uses: Intermediate; crosslinking and insolubilizing
agent.

triethylaconitate

C₂H₅OOCCCH(COOC₂H₅)CH₂COOC₂H₅.

Properties: Liquid. Sp. gr. (25°C) 1.096; refractive
index (26°C) 1.4517; b.p. (5 mm) 154-156°C. Com-
bustible.

Use: Plasticizer.

triethylaluminum (ATE; TEA; aluminum triethyl).

(C₂H₅)₃Al.

Properties: Colorless liquid; sp. gr. 0.837; f.p.
-52.5°C; b.p. 194°C; specific heat 0.527 (91.4°F).
Miscible with saturated hydrocarbons. Flash point
-63°F.

Derivation: By introduction of ethylene and hydrogen
into an autoclave containing aluminum. The reaction
proceeds under moderate temperature and
varying pressures.

Grades: 88-94%.

Containers: Cylinders.

Hazard: Highly toxic; destructive to tissue. Flam-
mable, dangerous fire risk; ignites spontaneously in
air; reacts violently with water, acids, alcohols, halo-
gens, and amines.

Uses: Catalyst intermediate for polymerization of ole-
fins, especially ethylene; pyrophoric fuels; production
of alpha-olefins and long chain alcohols; gas plating
of aluminum.

Shipping regulations: (Rail) Red label. (Air) Not ac-
ceptable. (Rail) Legal label name; aluminum triethyl.

triethylamine (C₂H₅)₃N.

Properties: Colorless liquid; strong ammoniacal odor.
B.p. 89.7°C; f.p. -115.3°C; sp. gr. (20/20°C) 0.7293;
wt/gal (20°C) 6.1 lb; flash point (open cup) 20°F.
Soluble in water and alcohol.

Derivation: From ethyl chloride and ammonia under
heat and pressure.

Containers: 1-, 5-gal cans; 55-gal drums; tank cars.

Hazard: Flammable, dangerous fire risk. Explosive
limits in air 1.2 to 8.0%. Tolerance, 25 ppm in air.
Toxic by ingestion and inhalation; strong irritant to
tissue.

Uses: Catalytic solvent in chemical synthesis; accel-
erator activators for rubber; wetting, penetrating and
waterproofing agents of quaternary ammonium

P-AH TP-90B - SPGR, 0.97. High boiling, straw colored liquid; little or no odor; non-toxic in ordinary handling. Visc (27°C) 8 cps.
Bp 660-760°F.

APPENDIX V

TABLE 15.3

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers</i>												
1	Acetin	C ₆ H ₁₀ O ₄	134	1.190 at 25° C.	1.445 at 25° C.	68.8 at 25° C.						
2	<i>o</i> - and <i>p</i> -Toluenesulfonamides (mixture)	C ₇ H ₉ O ₂ NS	171	1.353 solid		f.p. 105						
3	Diacetin	C ₇ H ₁₂ O ₅	176	1.178	1.438 at 25° C.	35.7 at 25° C.						
4	<i>N</i> -Ethyl- <i>p</i> -toluenesulfonamide	C ₉ H ₁₃ O ₂ NS	199	1.171 at 85° C.	1.522 at 65° C.	59.0 at 25° C.						
5	<i>N</i> -Ethyl- <i>o</i> - and <i>p</i> -toluenesulfonamides (mixture)	C ₉ H ₁₃ O ₂ NS	199	1.190 at 25° C.	1.540 at 25° C.	426 at 25° C.				0.55	101	
6	Triacetin	C ₉ H ₁₄ O ₆	218	1.157 (av.)	1.429 at 25° C.	15.1 at 25° C.						
7	Dimethyl phthalate	C ₁₀ H ₁₀ O ₄	194	1.191	1.513 at 25° C.	16.3 at 25° C.			0.5	0.34	>4.0	>1000
8	<i>o</i> -Nitrobiphenyl	C ₁₂ H ₉ O ₂ N	199	1.203 at 25° C.	1.613 at 25° C.	38 at 25° C.						
9	Diethyl phthalate	C ₁₂ H ₁₄ O ₄	222	1.118	1.501 at 20° C.	12.6 at 20° C.	0.15	0.7	2.8	1.1	3.40	>500
10	Triethyl citrate	C ₁₂ H ₂₀ O ₇	276	1.130 at 25° C.	1.441 at 24.5° C.							
11	Dimethyl sebacate	C ₁₂ H ₂₂ O ₄	230	0.986 at 30° C.	1.436 at 30° C.	3.5 at 38° C.						

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^m	
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		
									10 days, 60° C.	24 hours, 70° C.	Water		Oil
<i>Simple Plasticizers</i>													
1	Acetin												Kee
2	<i>o</i> - and <i>p</i> -Toluenesulfonamides (mixture)												Mon
3	Diacetin												Kee
4	<i>N</i> -Ethyl- <i>p</i> -toluenesulfonamide												Mon
5	<i>N</i> -Ethyl- <i>o</i> - and <i>p</i> -toluenesulfonamides (mixture)												Mon
6	Triacetin												Kee, TE
7	Dimethyl phthalate												Mon, TE
8	<i>o</i> -Nitrobiphenyl												Mon
9	Diethyl phthalate												Mon, TE, UBI
10	Triethyl citrate												Pf
11	Dimethyl sebacate												Har

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.
^m For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
12	Di-n-butyl tartrate	C ₁₂ H ₂₂ O ₆	262	1.094	1.444 at 25° C.	67.7 at 25° C.	0.5	0.7	0.3	2.4	1.50	143
18	Tri-n-butyl phosphate	C ₁₂ H ₂₇ O ₄ P	266	0.978	1.424 at 20° C.	3.4 at 25° C.	< 0.07	6.0	M	M	> 4	502
14	Monomethyl phthalate ethyl glycolate ester	C ₁₃ H ₁₄ O ₆	266	1.227	1.504 at 25° C.	185	0.1	1.2	0.5	0.1	0.65	61
15	N-Cyclohexyl-p-toluenesulfonamide	C ₁₃ H ₁₉ O ₂ NS	253	1.125	1.498 solid at 25° C.	87° C.	< 0.08	1.38	1.60	0.18	0.50	93
16	Monoethyl phthalate ethyl glycolate ester	C ₁₄ H ₁₈ O ₆	280	1.186	1.501 at 25° C.	53	0.85	3.2	0.50	0.10	0.16	61
17	Di(methyl CELLLOSOLVM) phthalate	C ₁₄ H ₁₈ O ₆	282	1.169	1.439 at 25° C.	70						> 525
18	Triethyl acetylacrylate	C ₁₄ H ₂₂ O ₈	318	1.13	1.428 at 25° C.	70						
19	Diisobutyl adipate	C ₁₄ H ₂₆ O ₄	258	0.957	1.433 at 25° C.	8.9						
20	Tributyrin	C ₁₅ H ₂₆ O ₆	302	1.036	1.440 at 25° C.	7.5						
21	Methyl CELLLOSOLVM laurate	C ₁₅ H ₃₀ O ₈	258	0.894 at 25° C.	1.440 at 25° C.	7.5						

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ²		
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac- tion, 10 days, 23° C., 4-mil Film		Spew, 2 days, 60° C., 100% Humidity ^b	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
<i>Simple Plasticizers (Continued)</i>														
12	Di-n-butyl tartrate												Kes	
13	Tri-n-butyl phosphate												CS	
14	Monomethyl phthalate ethyl glycolate ester	55.4°						17.5		6.1	5.9		Mon	
15	N-Cyclohexyl-p-toluenesulfonamide												Mon	
16	Monoethyl phthalate ethyl glycolate ester												Mon	
17	Di(methyl CELLLOSOLVM) phthalate	46°						6.6		4.6	3.7	None	OA, TE	
18	Triethyl acetylacrylate												PT	
19	Diisobutyl adipate												Cab	
20	Tributyrin												TE	
21	Methyl CELLLOSOLVM laurate												Kes	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²²²² For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	217° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
22	Di-n-butyl phthalate	C ₁₆ H ₂₂ O ₄	278	1.048	1.493 at 20° C.	20.3	Insol.	0.46	34	30.0	0.98	148
23	Di(CELLOSOLVE) phthalate	C ₁₆ H ₂₂ O ₆	310	1.120	1.491 at 25° C.	f.p. 31° C.	0.2	2.0	2	0.64	0.15	30
24	n-Butyl laurate	C ₁₈ H ₃₄ O ₂	256	0.857 at 25° C.								
25	Methyl CELLOSOLVE myristate	C ₁₇ H ₃₄ O ₃	286	0.895 at 25° C.								
26	Di(butyl CARBITOL) formal	C ₁₇ H ₃₆ O ₆	336	0.97								
27	Triphenyl phosphate	C ₁₈ H ₁₅ O ₄ P	326	1.268 at 60° C.	1.552 at 60° C.	f.p. 49.9° C.					0.02	22
28	Bis(dimethylbenzyl) ether	C ₁₈ H ₂₂ O	286	1.008	1.553 at 25° C.	29.0						
29	n-Butyl cyclohexyl phthalate	C ₁₈ H ₂₄ O ₄	304	1.078	1.508 at 20° C.	117.0						75
30	Mono-n-butyl phthalate n-butyl glycolate ester	C ₁₈ H ₂₄ O ₆	336	1.103	1.490 at 25° C.	64.9	Insol.	0.37	4.65	2.70	0.08	31
31	Tri-n-butyl citrate	C ₁₈ H ₃₂ O ₇	360	1.045	1.443 at 25° C.		Insol.	<0.2	10.0	14.5	0.16	44
32	Triethylene glycol di(2-ethylbutyrate)	C ₁₈ H ₃₄ O ₆	346	0.995	1.440 at 20° C.	11.5	0.02	1.0	0.2	24.0	0.86	

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a											Supplier ^m	
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film			Spew, 2 days, 60° C., 100% Humidity ^b
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Simple Plasticizers (Continued)</i>														
22	Di-n-butyl phthalate	41.4 ^c							17.5		2.7	6.8	None	Bar, CS, Har, Mon, Nev, Pitt, TE, USI, DA
23	Di(CELLOSOLVE) phthalate												—	OA
24	n-Butyl laurate												—	AH
25	Methyl CELLOSOLVE myristate												—	AH
26	Di(butyl CARBITOL) formal	55.3 ^c							10.1		15.6	14.5	None	Thi
27	Triphenyl phosphate												—	Dow, Mon, Pitt
28	Bis(dimethylbenzyl) ether												—	OA
29	n-Butyl cyclohexyl phthalate	58.8				+3	-20	6.5		0.6	8.8	None	Bar	
30	Mono-n-butyl phthalate n-butyl glycolate ester	53.2 ^c	3000	300				3.3		17	4.3	None	Mon	
31	Tri-n-butyl citrate	55.1	2800	300		-4	-32	3.1		8.1	14.6	None	Pf	
32	Triethylene glycol di(2-ethylbutyrate)												—	CCCC

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.⁵⁶^m For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
33	Butyl CELLOSOLVE laurate	C ₁₈ H ₃₆ O ₂	300	0.882 at 25° C.	1.439 at 25° C.	6.5 at 25° C.						
34	Di-n-hexyl adipate	C ₁₈ H ₃₄ O ₄	314	0.933 (av.) at 25° C.	1.439 at 25° C.	7.8 at 25° C.						
35	Di(1,3-dimethylbutyl) adipate	C ₁₈ H ₃₄ O ₄	314	0.926 at 25° C.	1.433 at 25° C.	5.6 ca. at 38° C.						
36	Di-n-butyl sebacate	C ₁₈ H ₃₄ O ₄	314	0.936 at 25° C.	1.440 at 25° C.	9.3 at 25° C.	Insol.	1.6	M	M		63
37	Di(butyl CELLOSOLVE) adipate	C ₁₈ H ₃₄ O ₆	346	0.997 at 25° C.	1.442 at 25° C.	12.5						58
38	n-Butyl myristate	C ₁₈ H ₃₆ O ₂	284	0.861 at 25° C.	1.022 at 25° C.	12.2	0.08	6.3	Almost miscible			
39	Tri(butyl CELLOSOLVE) phosphate	C ₁₈ H ₃₉ O ₇ P	398	1.208 at 25° C.	1.563 at 20° C.	39.8						
40	Cresyl diphenyl phosphate	C ₁₉ H ₁₇ O ₄ P	340	1.116 at 25° C.	1.538 at 20° C.	39.3 at 25° C.						29
41	n-Butyl benzyl phthalate	C ₁₉ H ₂₀ O ₄	312	1.083 at 25° C.	1.547 at 20° C.	155						32
42	Bis(dimethylbenzyl) carbonate	C ₁₉ H ₂₂ O ₃	298									

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalences ^a										Supplier ^m		
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extraction, 10 days, 23° C., 4-mil Film		Spec. 2 days, 60° C., 100% Humidity ^b	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
<i>Simple Plasticizers (Continued)</i>														
33	Butyl CELLOSOLVE laurate												Kes	
34	Di-n-hexyl adipate												Kes	
35	Di(1,3-dimethylbutyl) adipate												Har	
36	Di-n-butyl sebacate	37.8 ^c						7.2		0.4	10.5	None	Har, Pitt, RH	
37	Di(butyl CELLOSOLVE) adipate	41.4 ^c									18.0	None	Kes, OA	
38	n-Butyl myristate												AH	
39	Tri(butyl CELLOSOLVE) phosphate	43.7 ^c						1.1		14.2	13.5	None	OA	
40	Cresyl diphenyl phosphate	72	3000	292	64	475	4	-15		6.1	0.9	14.8	None	Mon
41	n-Butyl benzyl phthalate	59	2930	281	66	730	6	-11		9.5	0.6	7.8	None	Mou
42	Bis(dimethylbenzyl) carbonate	71	3000	284	63	450	2	-9		11.8	1.5	9.2		OA

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁰^m For key to supplier abbreviations see page 903

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
43	Methyl pentachlorostearate	C ₁₉ H ₂₂ O ₂ Cl ₅	471	1.204	1.497 at 20° C. 1.482 at 25° C.	975.0						7
44	Methyl ricinoleate	C ₁₉ H ₃₆ O ₃	312	0.925 at 25° C.	1.462 at 25° C.							
45	Methyl CELLOSOLV ₈ palmitate	C ₁₉ H ₃₈ O ₃	314	0.891 at 25° C.								
46	Methyl hydroxystearate	C ₁₉ H ₃₈ O ₃	314			f.p. 48° C.						
47	Diphenyl phthalate	C ₂₀ H ₁₄ O ₄	318	1.28 solid	1.572 at 74° C.						0.006	10
48	Dicyclohexyl phthalate	C ₂₀ H ₂₆ O ₄	330	1.23		f.p. 63.5° C.						
49	2-Ethylhexyl diphenyl phosphate	C ₂₀ H ₃₇ O ₄ P	362	1.092	1.511 at 20° C. 1.487 at 30° C.	29.5	Insol.	0.24				44
50	Di-n-hexyl phthalate	C ₂₀ H ₃₀ O ₄	334	1.007	1.488 at 25° C.	56	Insol.	<0.05	M	M	0.20	43
51	Di(2-ethylbutyl) phthalate	C ₂₀ H ₃₀ O ₄	334	1.016	1.488 at 25° C.	28.8 cs. at 38° C.	Insol.	<0.05				65
52	Di(1,3-dimethylbutyl) phthalate	C ₂₀ H ₃₀ O ₄	334	0.995	1.480 at 25° C.							150

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^b		
		Plasticizer p.p.h. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extraction, 10 days, 23° C., 4-mil Film		Specs, 2 days, 60° C., 100% Humidity, %	
		10 days, 60° C.	24 hours, 70° C.	Water	Oil									
<i>Simple Plasticizers (Continued)</i>														
43	Methyl pentachlorostearate	69 ^d	2900	338	64	600	-6	-19	0.2		0.2	17.1	None	Hook
44	Methyl ricinoleate													Bak
45	Methyl CELLOSOLV ₈ palmitate													AH
46	Methyl hydroxystearate													Bak
47	Diphenyl phthalate													Mon
48	Dicyclohexyl phthalate	64 ^e	3000	290			+5	-5	1.0		0	4.5	None	Bar, OA
49	2-Ethylhexyl diphenyl phosphate	60	2580	260	64	540	-2	-28	1.6		0.2	14.8	None	Mon
50	Di-n-hexyl phthalate	57	2720	293	63	570	-8	-81	8.8	11.6	0	18.5	None	CCCC, TE
51	Di(2-ethylbutyl) phthalate	45.3 ^o							4.4		0.2	8.4	None	TE
52	Di(1,3-dimethylbutyl) phthalate													Har

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Ref. 23^d Mixture of 1 part plasticizer to 2 parts FLAXOL plasticizer DOP.^e Mixture of 0.9 part plasticizer to 1 part FLAXOL plasticizer DOP.^f For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
53	Di(butyl CELLOSOLVE) phthalate	C ₂₀ H ₃₀ O ₆	360	1.063	1.483 at 25° C.	42.0	Insol.	0.63	2.80	6.50	0.06	24
54	Di(CARBITROL) phthalate	C ₂₀ H ₃₀ O ₆	398	1.150	1.492 at 25° C.	82.0	Insol.	2.80	0.09	0.20	0.08	18
55	Tri-n-butyl acetylacrylate	C ₂₀ H ₃₄ O ₆	402	1.046 at 25° C.	1.441 at 25° C.							61
56	n-Butyl palmitate	C ₂₀ H ₄₀ O ₂	313	0.866 at 25° C.								
57	Tricresyl phosphate	C ₂₁ H ₃₁ O ₄ P	368	1.165	1.557 at 20° C.	120.0	Insol.	0.37	5.5	2.0	0.01	7
58	n-Butyl benzyl sebacate	C ₂₁ H ₃₂ O ₄	348	1.004	1.479 at 25° C.	8.6 ca. at 38° C.						
59	Methyl acetylricinoleate	C ₂₁ H ₃₈ O ₄	355	0.936	1.456 at 20° C.	23.1						
60	Methyl CELLOSOLVE oleate	C ₂₁ H ₄₀ O ₃	341	0.902	1.453 at 25° C.	10.2				0.12		29
61	Di(2-ethylbutyl) aselate	C ₂₁ H ₄₀ O ₄	357	0.930	1.442 at 25° C.	11.4						43
62	Propylene glycol monoricinoleate	C ₂₁ H ₄₀ O ₄	357	0.950 at 25° C.								
63	Methyl CELLOSOLVE stearate	C ₂₁ H ₄₂ O ₃	343	0.877 at 25° C.	1.443 at 25° C.	8.9 at 25° C.						30

PLASTICIZERS AND PLASTICIZATION

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^m		
		Plasticizer p.h.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Specs, 2 days, 60° C., 100% Humidity	
		10 days, 60° C.	24 hours, 70° C.	Water	Oil									
<i>Simple Plasticizers (Continued)</i>														
53	Di(butyl CELLOSOLVE) phthalate	47.8°						2.1		4.8	11.4	None	OA	
54	Di(CARBITROL) phthalate	54.4°						2.0		10.1	9.9	None	OA	
55	Tri-n-butyl acetylacrylate											—	PI	
56	n-Butyl palmitate											—	AH	
57	Tricresyl phosphate	75	2880	274	58	580	+ 5	- 3	0.1	0.7	6.1	None	Cal, Mon, OA, Pitt	
58	n-Butyl benzyl sebacate											—	Har	
59	Methyl acetylricinoleate	47.8°							8.8	9.6	11.0	—	Bak	
60	Methyl CELLOSOLVE oleate	50.5° ^f							3.6	3.0	10.3	—	AH, Kes, OA	
61	Di(2-ethylbutyl) aselate	44.5	2800	260					-40	5.2	1.5	18.0	None	Em
62	Propylene glycol monoricinoleate											—	Bak	
63	Methyl CELLOSOLVE stearate											—	AH, Kes	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁵^f Mixture of 0.63 part plasticizer to 1 part tricresyl phosphate.^m For key to supplier abbreviations see page 903.

DESCRIPTION OF TABLES AND FIGURES

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TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (131° C.)
Simple Plasticizers (Continued)												
64	n-Butyl ricinoleate	C ₃₂ H ₄₂ O ₃	355									
65	Di(1,3-dimethylbutyl) sebacate	C ₂₂ H ₄₂ O ₄	371	0.911	1.439 at 25° C. 1.447 at 20° C.	9.2 cs. at 38° C. 18.7 at 20° C.	Insol.	0.12				41
66	Di(2-ethylhexyl) adipate	C ₂₂ H ₄₂ O ₄	371	0.927	1.448 at 25° C. 1.448 at 20° C.	17.7 at 25° C. 17.7 at 20° C.						38
67	Diisooctyl adipate	C ₂₂ H ₄₂ O ₄	371	0.928	1.440 at 25° C. 1.440 at 20° C.	8.3 cs. at 38° C. 19 at 20° C.						
68	Dicapryl adipate	C ₂₂ H ₄₂ O ₄	371	0.915	1.440 at 25° C. 1.447 at 20° C.	19 at 25° C. 19 at 20° C.						
69	Diethylene glycol dipelargonate	C ₂₂ H ₄₂ O ₅	387	0.963	1.447 at 20° C.							
70	Di(butyl CELLOSOLVE) sebacate	C ₂₂ H ₄₂ O ₆	403	0.970								0.04
71	Triethylene glycol di(2-ethyl hexanoate)	C ₂₂ H ₄₂ O ₆	403	0.968	1.444 at 20° C.	16.1	Insol.	0.4	M	M		0.18
72	Triethylene glycol dicaprylate	C ₂₂ H ₄₂ O ₆	406 (av.)	0.973								
73	Di(butyl CARBITROL) adipate	C ₂₂ H ₄₂ O ₆	435	1.02 at 16° C.		15 at 22° C. 7.9 at 25° C.						
74	n-Butyl stearate	C ₃₂ H ₆₄ O ₂	341	0.858	1.444 at 20° C.							59

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^m		
		Plasticizer p.p.h. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extraction, 10 days, 23° C., 4-mil Film		Spew, 2 days, 60° C., 100% Humidity ^b	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
Simple Plasticizers (Continued)														
64	n-Butyl ricinoleate												Dee	
65	Di(1,3-dimethylbutyl) sebacate												Har	
66	Di(2-ethylhexyl) adipate	51	2730	318	67	600	-15	-53	3.5	12.2	2.2	21.0	None	CCCC, Gdr, MWI, Kes, OA, Har
67	Diisooctyl adipate													Kes, MWI, OA, Pitt, RCA, Har, Cab
68	Dicapryl adipate													Har
69	Diethylene glycol dipelargonate	51.3	2800	330			-15	-48	6.9		6.0	22.0	Yes	Em
70	Di(butyl CELLOSOLVE) sebacate													Dee
71	Triethylene glycol di(2-ethylhexanoate)	59	2680	300	69	800	-17	-50	4.7	12.4	2.0	22.8	None	CCCC
72	Triethylene glycol dicaprylate													RCA, Drew
73	Di(butyl CARBITROL) adipate													Thi
74	n-Butyl stearate													CS, AH, Har, Kes, OA

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.

^b Dash (—) indicates no data available.

^m For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.		
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)	
Simple Plasticizers (Continued)													
75	n-Butyl oleate	C ₂₂ H ₄₂ O ₂	339	0.864 at 25° C.	1.449 at 25° C.	7.7 at 25° C.							
76	Tetrahydrofurfuryl oleate	C ₂₃ H ₄₂ O ₃	366	0.925	1.466 at 20° C.	17.8							
77	Methyl CALLOSOLVE acetylricinoleate	C ₂₃ H ₄₂ O ₅	399	0.955	1.456 at 20° C.	42.1	Insol.	0.6	M	M		12	
78	o-Xenyl diphenyl phosphate	C ₂₄ H ₁₉ O ₄ P	402	1.233	ca. 1.550 at 60° C.	30-55 at 60° C.	Insol.	0.35	2.22	0.3	0.005	20	
79	Dibenzyl sebacate	C ₂₄ H ₃₈ O ₄	382	1.055 at 30° C.	1.482 at 30° C.	13.0 at 35° C.							
80	Di(2-ethylhexyl) phthalate	C ₂₄ H ₃₈ O ₄	390	0.986	1.486 at 20° C.	81.0	Insol.	0.20	M	M	0.02	24	
81	Diisooctyl phthalate	C ₂₄ H ₃₈ O ₄	390	0.987	1.488 at 20° C.	83.0						19	
82	Dicaprlyl phthalate	C ₂₄ H ₃₈ O ₄	390	0.966	1.478 at 20° C.	66.7						22	
83	Di(2-ethylhexyl) tetrahydrophthalate	C ₂₄ H ₄₂ O ₄	395	0.969	1.486 at 20° C.	42.0	Insol.	0.26				80	
84	Tetra-n-butyl thiodisuccinate	C ₂₄ H ₄₂ O ₄ S	491	1.054	1.463 at 20° C.	63.6	Insol.	0.22				6	
85	Di(2-ethylhexyl) hexahydrophthalate	C ₂₄ H ₄₄ O ₄	397	0.956	1.481 at 20° C.	35.9						37	
86	n-Butyl acetylricinoleate	C ₂₄ H ₄₄ O ₄	397	0.929	1.456 at 25° C.	35.3						30	

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^m			
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Swow, 2 days, 60° C., 100% Humidity ^b		
									10 days, 60° C.	24 hours, 70° C.	Water			Oil	
Simple Plasticizers (Continued)															
75	n-Butyl oleate												AH, Kes		
76	Tetrahydrofurfuryl oleate	59 ^d	2590	297	67	650	-8	-34	0.8		0.3	17.2	None	AH, Em, Pitt	
77	Methyl CALLOSOLVE acetylricinoleate	50.8 ^c							2.4		1.3	14.5	None	Bak, Dee	
78	o-Xenyl diphenyl phosphate													Dow	
79	Dibenzyl sebacate	45.3 ^c							0.7		0.3	11.8	None	Har	
80	Di(2-ethylhexyl) phthalate	61	2360	253	67	700	-3	-31	0.7	5.1	0	15.3	None	Bar, CCCC, Har, Mon, MWI, OA, Pitt, TE, DA, Gdr	
81	Diisooctyl phthalate	56	2700	300			-1	-27					None	Bar, Cab, Har, MWI, OA, Pitt, RCA	
82	Dicaprlyl phthalate	64	2500	270			-5	-35	0.6		0	33.8	None	RH, Har	
83	Di(2-ethylhexyl) tetrahydrophthalate	66	2590	320	88	560	-13	-37	4.1	11.5	0.3	21.8	None	CCCC	
84	Tetra-n-butyl thiodisuccinate	64	3200	318	62	760	-2	-22	0	2.5	0.8	15.2	None	CCCC	
85	Di(2-ethylhexyl) hexahydrophthalate	64	2580	300	68	750	-10	-34			9.7	0	25.4	None	CCCC
86	n-Butyl acetylricinoleate	55.4 ^c											Yes	Bak	

Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence^a

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.
^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁶
^d Mixture of 1 part plasticizer to 2 parts Flexol plasticizer DOP.
^m For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, sp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
87	Butyl CELLOSOLVE oleate	C ₂₄ H ₄₆ O ₂	383	0.885 at 25° C.	1.454 at 25° C.	5.5 at 25° C.						
88	Dinonyl adipate	C ₂₄ H ₄₆ O ₄	398	0.914 at 25° C.	1.445 at 25° C.	18.6 at 25° C.						
89	n-Octyl n-decyl adipate (mixture)	C ₂₄ H ₄₆ O ₄ (est.)	398 (av.)	0.920	1.450 at 20° C.	15.4 at 25° C.						
90	Polyethylene glycol di(2-ethylhexanoate)	C ₂₄ H ₄₆ O ₇	447	0.989	1.447 at 20° C.	25.1 at 25° C.	Insol.	1.4				44
91	Butyl CELLOSOLVE stearate	C ₂₄ H ₄₆ O ₂	385	0.882	1.446 at 25° C.	13.3 at 25° C.						26
92	Tri(2-ethylhexyl) phosphate	C ₂₄ H ₅₀ O ₄ P	435	0.926	1.443 at 20° C.	14.1 at 25° C.	Insol.	1.4	M	M		89
93	Di(2-ethylhexyl) azelate	C ₂₆ H ₄₈ O ₄	413	0.918	1.450 at 20° C.	20 at 25° C.						20
94	n-Octyl n-decyl phthalate (mixture)	C ₂₈ H ₄₈ O ₄ (est.)	418 (av.)	0.975	1.484 at 20° C.	72 at 25° C.						8
95	Isooctyl n-octyl n-decyl phthalate (mixture)	C ₂₈ H ₄₈ O ₄ (est.)	418 (av.)	0.978	1.485 at 20° C.	42.9 at 25° C.						
96	Di(2-ethylhexyl) sebacate	C ₂₈ H ₅₀ O ₄	427	0.912	1.451 at 20° C.	19.9 at 25° C.						12
97	Diisooctyl sebacate	C ₂₈ H ₅₀ O ₄	427	0.917	1.447 at 25° C.	20.4 at 25° C.						
98	Dicapryl sebacate	C ₂₈ H ₅₀ O ₄	427	0.907	1.444 at 25° C.	13.0 ca. at 38° C.						

M = Miscible.

PLASTICIZERS AND PLASTICATION

No.	Name	Comparison of Properties of Vynlite Resin VYNW-5 Specimens Plasticized to Equivalence ^a											Supplier ^m	
		Plasticizer p.p.h. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film			Spew, 2 days, 60° C., 100% Humidity ^b
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Simple Plasticizers (Continued)</i>														
87	Butyl CELLOSOLVE oleate													Kee
88	Dinonyl adipate													Pitt
89	n-Octyl n-decyl adipate (mixture)	53	2550	295	78	1050	-7	-54	5.5	0.3	23.5	None	None	Hero, OA
90	Polyethylene glycol di(2-ethylhexanoate)	62	2690	308	70	750	-12	-44	2.8	8.8	3.0	19.4	None	CCCC
91	Butyl CELLOSOLVE stearate													AH, Kee, OA
92	Tri(2-ethylhexyl) phosphate													CCCC
93	Di(2-ethylhexyl) azelate	59	2640	300	66	980	-16	-57	3.2	6.8	0.6	24.8	None	Eu
94	n-Octyl n-decyl phthalate (mixture)	57	2590	334	66	750	-17	-56	0.5		0.4	22.5	None	Hero
95	Isooctyl n-octyl n-decyl phthalate (mixture)	62	2640	318	64	640	-8	-38	0.2		0.3	21.8	None	Hero
96	Di(2-ethylhexyl) sebacate	67	2720	334	66	470	-12	-39		3.9	0	22.4	None	Hero
97	Diisooctyl sebacate	60	2420	293	67	1700	-19	-56	0.08	4.8	0.3	30.2	None	Dee, Har, MWI, RH
98	Dicapryl sebacate													Har, Pitt, MWI Har

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.
^m For key to supplier abbreviations see page 903.

DESCRIPTION OF TABLES AND FIGURES

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

922

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
99	2-Ethylhexanoic acid diester of <i>N,N</i> -bis(2-hydroxyethyl)-2-ethylhexanamide	C ₂₈ H ₅₃ O ₅ N	484	0.956	1.458 at 20° C.	139.2	Insol.	0.5				12
100	Tri(<i>p</i> - <i>tert</i> -butylphenyl) phosphate	C ₃₀ H ₃₉ O ₄ P	495			f.p. 95-99.5° C.						0.9
101	Glyceryl triricinoleate	C ₆₇ H ₁₀₄ O ₉	933	0.959 at 25° C.	1.477 at 25° C.							
102	Glyceryl tri(acetylricinoleate)	C ₆₃ H ₁₁₀ O ₁₂	1080	0.964	1.469 at 25° C.	271						0.7
<i>Resinous Plasticizers</i>												
103	AROCLOR 1242, chlorinated biphenyl				1.378-1.388 at 25° C.		Insol.	<0.1	<i>M</i>	<i>M</i>	2.50	146
104	FLEXOL plasticizer R-2H, polyester				1.055	ca. 20,000	Insol.	1.0				0.5
105	G.E. 2557, polyester				1.028	306						3
106	G.E. 2559, polyester				1.060	4278						1
107	PARACRIL C, nitrile rubber				0.984							
108	PARAPLEX G-25, polyester		ca. 8000		1.06	300,000						0.2

PLASTICIZERS AND PLASTICIZATION

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a												Supplier ^m
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.a.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Stiffness at 23° C., p.a.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C. 4-mil Film		Spec. 2 days, 60° C., 100% Humidity	
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Simple Plasticizers (Continued)</i>														
99	2-Ethylhexanoic acid diester of <i>N,N</i> -bis(2-hydroxyethyl)-2-ethylhexanamide	04	2730	327	63	650	- 3	-23	0.4	1.7	1.1	11.0	None	CCCC
100	Tri(<i>p</i> - <i>tert</i> -butylphenyl) phosphate												—	Dow
101	Glyceryl triricinoleate												—	Bak
102	Glyceryl tri(acetylricinoleate)	68.0 ^c							1.4		2.5	9.5	None	Bak
<i>Resinous Plasticizers</i>														
103	AROCLOR 1242, chlorinated biphenyl	56.2 ^b							4.5		0.6	0.6	None	Mon
104	FLEXOL plasticizer R-2H, polyester	90	3080	360	57	440	+2	-10	0	0	0	2.4	None	CCCC
105	G.E. 2557, polyester	72	2730	310	63	580	- 2	-18	0.9	3.5	3.1	17.0	Very slight	GE
106	G.E. 2559, polyester	90 ^d	2110	215	72	740	6	- 6	0.6	2.3	2.7	7.1	Very slight	GE
107	PARACRIL C, nitrile rubber	106	2500	410	60	480	+ 7	-54	0.5		0.2	0.6	None	Nau
108	PARAPLEX G-25, polyester	82	2600	310	60	480	- 1	-25	0.2		1.0	0.6	None	RH

DESCRIPTION OF TABLES AND FIGURES

^a 1000 p.a.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.a.i. per minute.
^b Dash (—) indicates no data available.
^c 100% elongation, 1000 p.a.i., constant rate of loading of 810 p.a.i. per minute. See Reed.²⁶
^d Mixture of 0.9 part plasticizer to 1 part tricresyl phosphate.
^e Only composition tested, does not yield standard elongation.
^f Highest plasticizer content tested. Elongation at 1000 p.a.i. = 64%.
^m For key to supplier abbreviations see page 903.

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TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

924

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Resinous Plasticizers (Continued)</i>												
109	PANAFLEX G-40, polyester		ca. 6000	1.15	1.472 at 20° C.	290,000						0.8
110	PANAFLEX G-50, polyester		ca. 2200	1.084	1.470 at 20° C.	3000						1
111	PLASTOLEIN 9720, polyester		ca. 850	1.043	1.462 at 20° C.	431						2
<i>Hydrocarbon-Type Plasticizers</i>												
112	Dow 276-V2, polymolecular product derived from α -methylstyrene			1.021	1.582 at 20° C.							115
113	HB 40, partially hydrogenated mixture of isomeric terphenyls			1.007	1.570 at 25° C.							
114	KENFLEX L, hydrocarbon resin made by reaction of formaldehyde + dimethylnaphthalenes			1.01 at 15.6° C.		f.p. -1° C.						
115	NEVILLAC 10° (PHO), a hydrindyl phenol derivative	240 (av.)		1.075 to 1.10 at 15.6° C.	1.597 at 25° C.	23,500 at 25° C.						
116	NEVINOL			1.03 to 1.08 at 15.6° C.	1.506 (av.) at 25° C.	65 to 110 at 25° C.						
117	PANAFLEX BN-1, alkylated aromatic hydrocarbon mixture			0.955	1.561 at 20° C.	151.6						
118	SOVALOID C, alkylated aromatic hydrocarbon mixture			1.038		105						

PLASTICIZERS AND PLASTICIZATION

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a											Supplier ^m	
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film			Specs, 2 days, 60° C., 100% Humidity
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Resinous Plasticizers (Continued)</i>														
109	PANAFLEX G-40, polyester	67 ^j	2780	310	69	760	1	-15	0.3		1.0	5.2	Slight	RH
110	PANAFLEX G-50, polyester	75	2630	320	62	680	0	-16	0.2		1.3	6.5	None	RH
111	PLASTOLEIN 9720, polyester	69	2850	312	66	900	-7	-23	0.3	1.8	1.0	14	None	Em
<i>Hydrocarbon-Type Plasticizers</i>														
112	Dow 276-V2, polymolecular product derived from α -methylstyrene	74 ^k	2420	315	62	650	-5	-24	1.4		0.6	22.6	None	Dow
113	HB 40, partially hydrogenated mixture of isomeric terphenyls	62 ^d	2890	298	66	940	-1	-23	8.4		0.5	15.5	None	Mon
114	KENFLEX L, hydrocarbon resin made by reaction of formaldehyde + dimethylnaphthalenes													Ken
115	NEVILLAC 10° (PHO), a hydrindyl phenol derivative	56.3 ^{e, h}							13.3		0.1	0.1	None	Nev
116	NEVINOL												None	Nev
117	PANAFLEX BN-1, alkylated aromatic hydrocarbon mixture	63 ^d	2800	310	66	920	2	-21	6.1		1.8	16.8	None	Pan
118	SOVALOID C, alkylated aromatic hydrocarbon mixture	61.4 ^l	3100	280			+4	-16	11.5		1.7	9.0	None	SV

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁵^d Mixture of 1 part plasticizer to 2 parts FLEXOL plasticizer DOP.^e Only composition tested, does not yield standard elongation.^f Mixture of 1 part plasticizer to 1 part FLEXOL plasticizer DOP.^k Mixture of 1 part plasticizer to 3 parts FLEXOL plasticizer DOP.^l Mixture of 0.85 part plasticizer to 1 part FLEXOL plasticizer DOP.^m For key to supplier abbreviations see page 903.

DESCRIPTION OF TABLES AND FIGURES

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TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

926

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Miscellaneous Plasticizers</i>												
119	ARNEEL TOD, mixture of oleic, linoleic, and cyclic nitriles		ca. 278	0.910		30.2						
120	n-Butyl acetyl polyricinoleate, PG16		362 (av.)	0.913 at 25° C. 1.164 at 23° C.	1.460 at 25° C. 1.505 at 23° C.	3100 at 25° C. 153 at 20° C.						
121	Chlorinated paraffin 40			1.164	1.505							
122	FLEXOL plasticizer B-400, polypropylene glycol mono-n-butyl ether			0.995	1.449 at 20° C.		0.1	3.3				ca. 60
123	Glyceryl polyricinoleate, No. 15 oil			1.031 at 25° C.	1.483 at 25° C.							
124	HARFLEX 500, monomeric ester type			0.932	1.488 at 20° C.	17.0						
125	HERCOFLEX 600, a pentaerythritol ester		538 (av.)	1.006	1.453 at 20° C.	48.3						
126	OHOPLEX Q-10, octyl fatty-phthalic acid esters			0.952	1.475 at 25° C.	41						
127	PARAFLEX G-60, high-molecular-weight ester		ca. 1000	0.994	1.474 at 20° C. 1.478 at 25° C.	240 to 300 at 54° C.						0.9
128	Plasticizer DP-520, chlorinated aliphatic product			1.126 at 15.8° C. 0.97 at 25° C.	1.478 at 25° C.	66.4 at 54° C.						
129	Polyethylene glycol dilaurate											
130	SANTICIZER 180			1.071	1.506 at 20° C.	234						
131	STAFLEX KA, monomeric ester type											

PLASTICIZERS AND PLASTICIZATION

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^m		
		Plasticizer p.h.r. at 23° C. 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spew, 2 days, 60° C., 100% Humidity ^b	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
<i>Miscellaneous Plasticizers</i>														
119	ARNEEL TOD, mixture of oleic, linoleic, and cyclic nitriles	52.8	2000	290			-44	11.7		0.7	14.8	None	Arm	
120	n-Butyl acetyl polyricinoleate, PG16	Incompatible at 55.3 p.h.r.										Bak		
121	Chlorinated paraffin 40	68 ^c	2820	335	67	900	-5	-28	1.0	0	20	None	Halo, Dia, Hero, Hook CCCC	
122	FLEXOL plasticizer B-400, polypropylene glycol mono-n-butyl ether												Bak	
123	Glyceryl polyricinoleate, No. 15 oil												Bak	
124	HARFLEX 500, monomeric ester type	63	2400	319	67	980	-10	-61	1.1	0.4	23.0	None	Har	
125	HERCOFLEX 600, a pentaerythritol ester	58	2910	327	66	900	-2	-28	0	1.1	12.8	None	Hero	
126	OHOPLEX Q-10, octyl fatty-phthalic acid esters												OA	
127	PARAFLEX G-60, high-molecular-weight ester	69	2800	317	67	880	-2	-21	0.2	0.4	11.1	None	RH	
128	Plasticizer DP-520, chlorinated aliphatic product												Drew	
129	Polyethylene glycol dilaurate												Gly	
130	SANTICIZER 180	73	2670	308	67	430	-2	-17		2.7	1.3	12.2	None	Mon
131	STAFLEX KA, monomeric ester type												Des	

DESCRIPTION OF TABLES AND FIGURES

927

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c Mixture of 0.9 part plasticizer to 1 part FLEXOL plasticizer DOP.^m For key to supplier abbreviations see page 903.

DUST COLLECTOR

DUCT WORK -

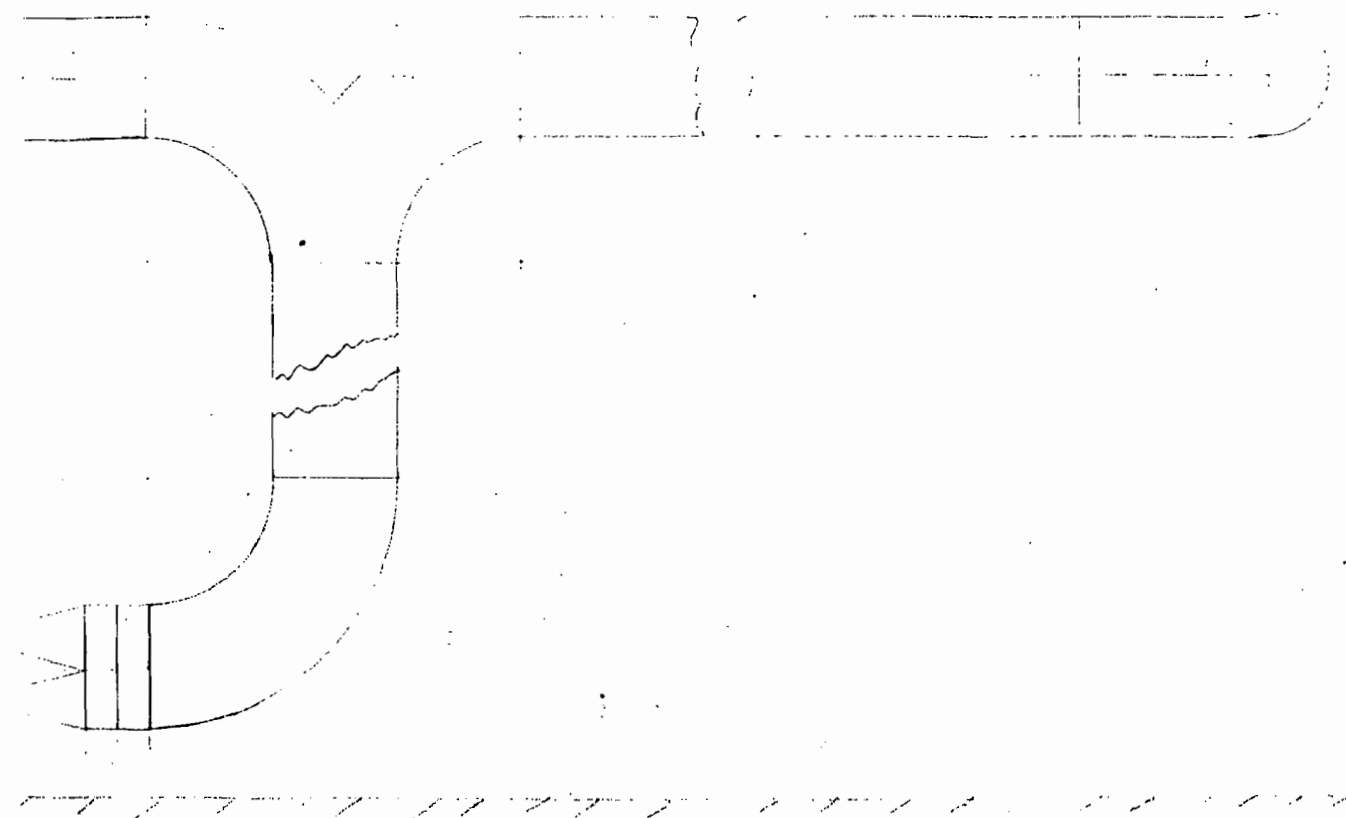
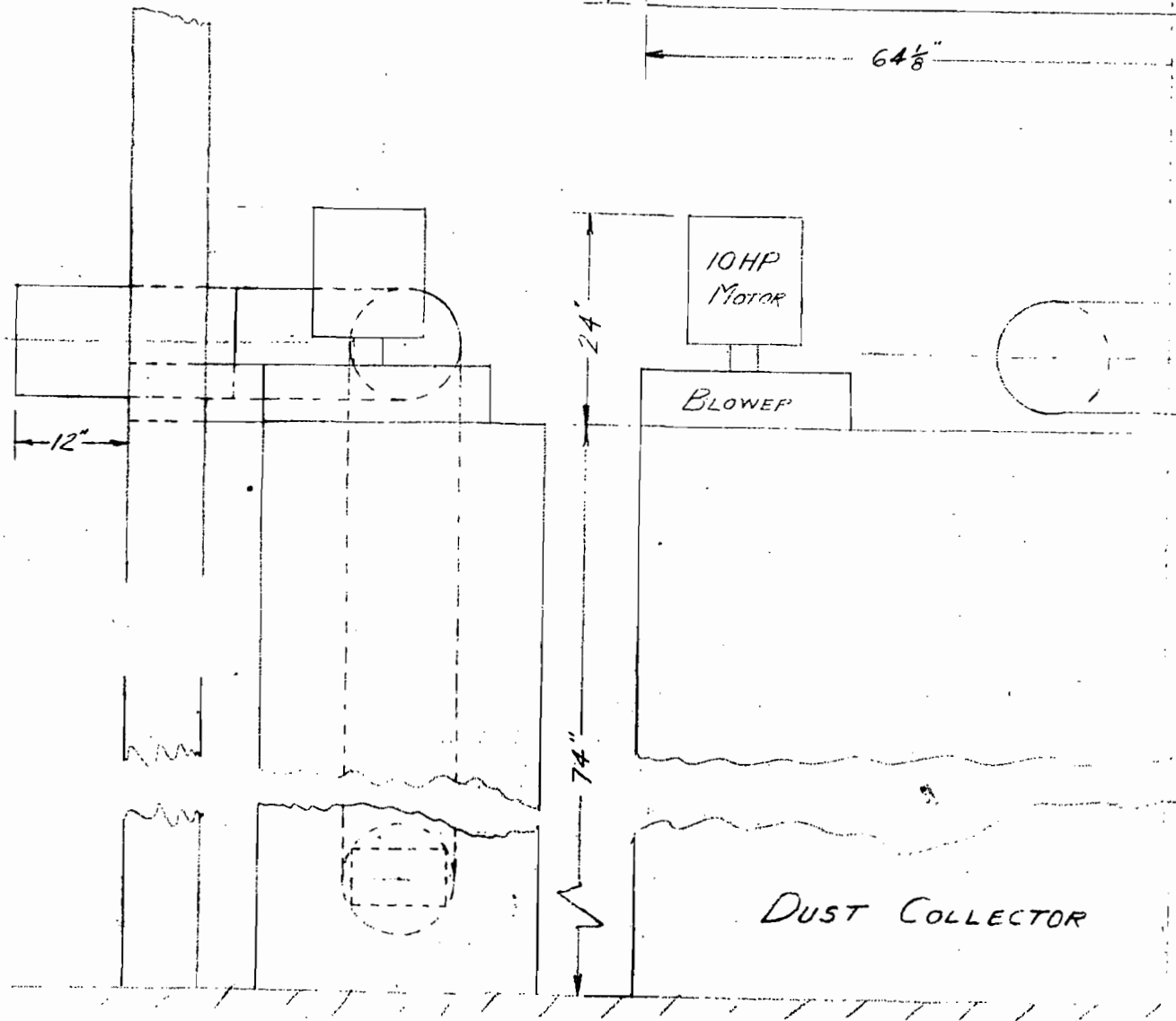
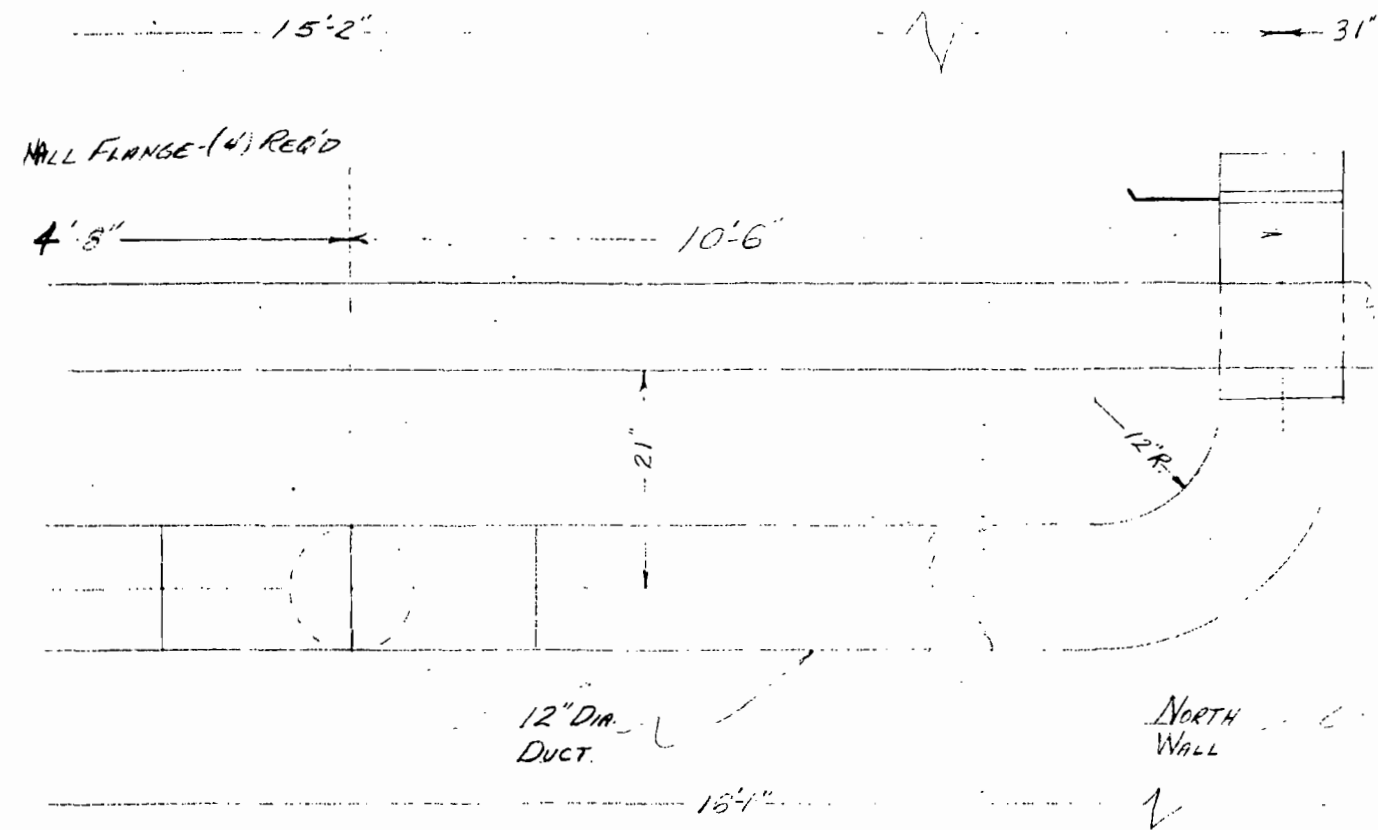
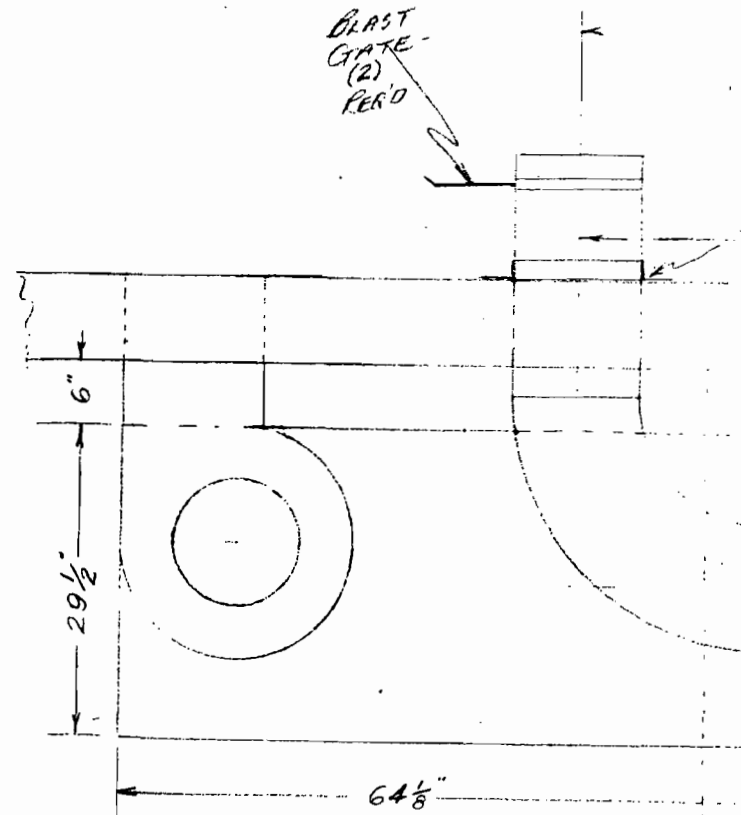
TALC APPLICATOR

DAVID M
ORLANDO

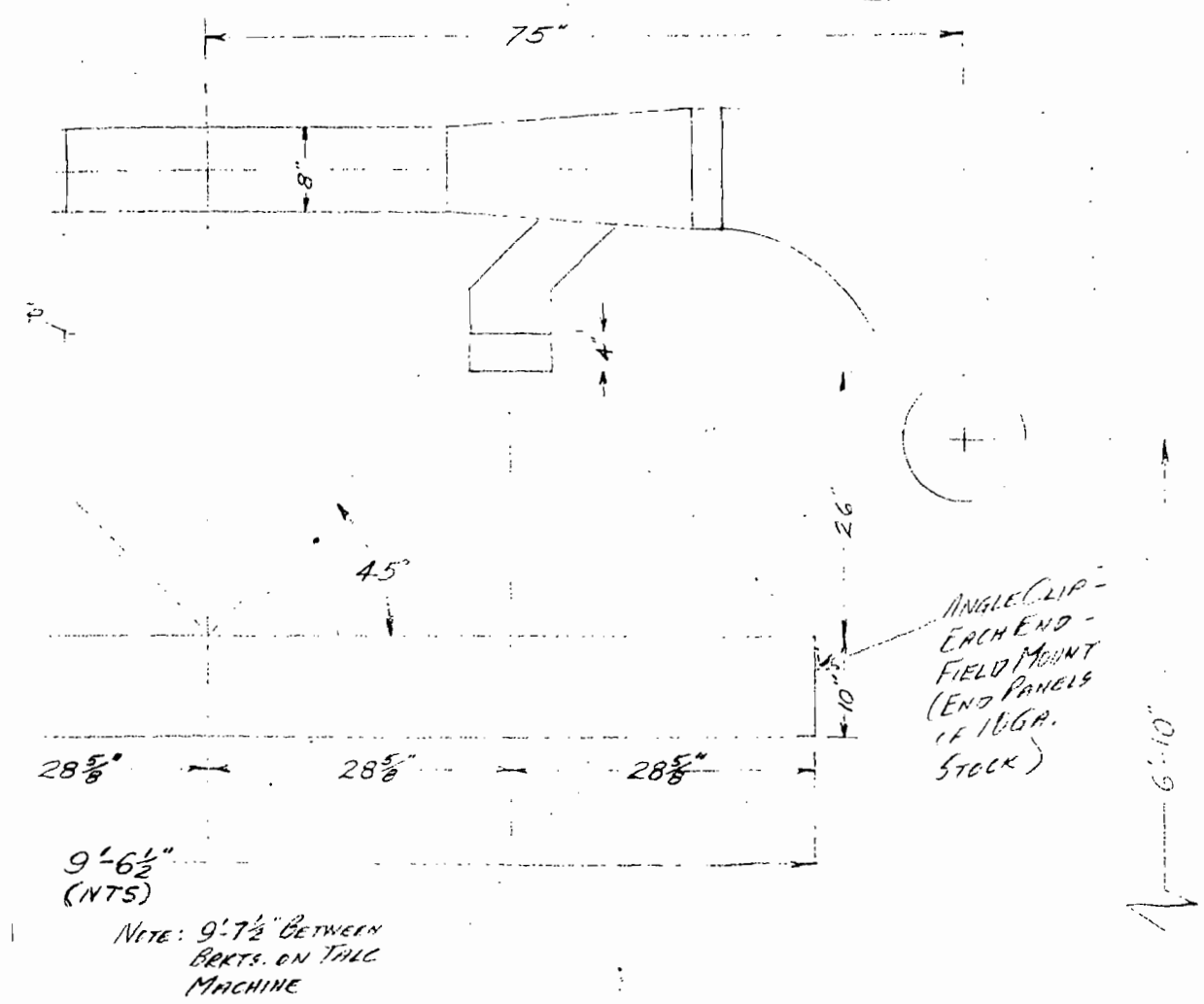
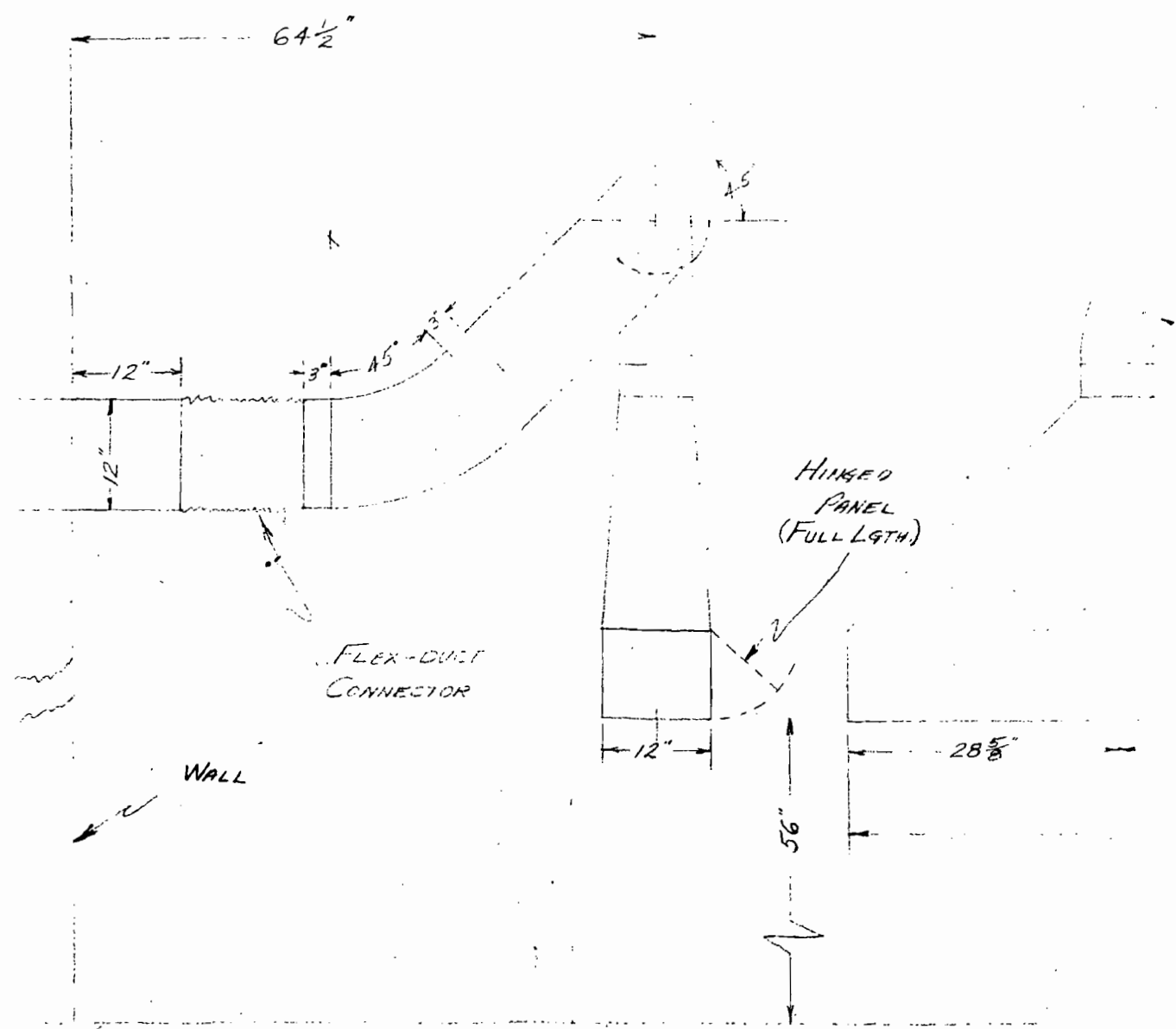
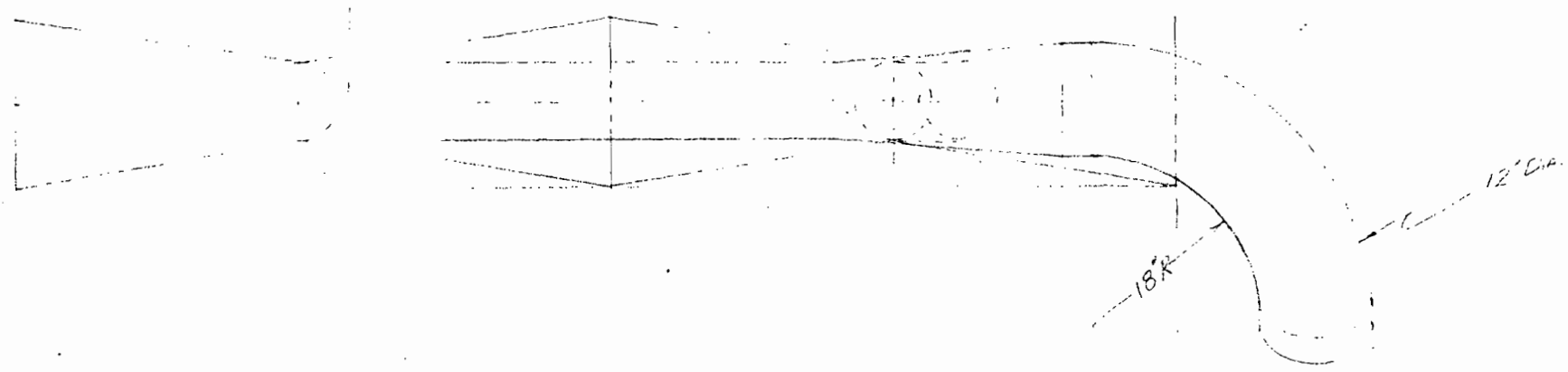
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DWG #2 OF 3 REV 9/22/83

REV 10/1/83



DUST COLLECTOR
HOOD & DUCT -
TALC APPLICATOR -
DAVID M
ORLANDO
SCALE: 1"=1'-0"
DWG. # 1 OF 3
9/1/83 RHB
REV 3/22/83
REV. 10/1/83



11. c) Will product scrap be recycled to the mills? If so, will VOC be emitted by the scrap?

No, and as such no VOC will be emitted by the scrap.

12. Are the flow rates listed in the application from points 4 and 5 correct?

Yes, each emission point has two (2) fans and the duct velocity in each is 800 fpm. Given the area of each duct (1.76 ft.²), the flow would equate to 1415 ACFM. In the original application the velocities in the ducts were incorrectly shown in feet per second. Those numbers were actually in feet per minute. Please excuse the typographical error.

13. What will be the percent of the lower explosive limit (LEL) at peak solvent evaporation rates for each exhaust stream?

The new ventilation system will be designed around peak solvent evaporation rates of 40% of the lower explosive limit and will incorporate the use of a lower explosive limit detection and control system designed to enhance safe operation.

14. What will be the percent of the threshold limit values (TLV) near the discharge points \approx 20 feet outside the plant during peak solvent evaporation rates?

The control device, which will provide at least a 90% removal efficiency can be expected to reduce emission levels 20 feet from the exhaust discharge points to well below the current (OSHA) TLV's for Toluene 200 ppm and MEK 200 ppm, however, without extensive modeling an estimated percent of the TLV at the property line cannot be established. Please advise of the necessity of performing said modeling in view of the proposed control program.

15. a) How many employees will work at this plant?

61 full-time employees.

- b) What impact will the emissions from the plant have on the ambient air quality, vegetation and visibility near the plant?

With the incorporation of a control device capable to continuous operation at or above 90% removal efficiency, adverse effects on air quality, visibility and vegetation are not anticipated since the control program reflects state of the art air pollution control technology.

16. If the solvents were recovered, could they be used or sold to a reclamation plant?

Yes, either avenue is a possibility.

17. What is the current delivered cost of the solvents used at the plant?

Toluene \$1.30/gallon

MEK \$.36/pound

18. Has the company made any studies on recovering and reusing the solvents at the plant and, if so, what conclusions were reached?

Yes, preliminary studies suggest that recovery and on-site re-use are well within the realm of feasibility, however, the necessary equipment requires a very substantial capital investment.

19. Please explain price estimate.

Enclosed you will find a copy of a budget quotation for a 25,000 SCFM solvent recovery plant (Appendix VI) which includes equipment and installation. Our original \$700,000.00 figure also included ventilation rework estimates and process control system installation. At this time it is not possible to guarantee that a solvent recovery or thermal incineration device will be of this size or cost since efforts are now underway to reduce the size of the system, however, if a system of such size is necessary due to the nature of the process (non-continuous peak evaporation type), the original \$700,000.00 figure is a very real number.

20. How many gallons of VOC per year will be shipped to Emille for disposal?

Approximately 220 (55 gallon drums) which would contain some 5,000 gallons of VOC. (By estimate.)

DISCUSSION:

In accordance with our recent conversation and the historical data provided in item one, David 'M' Company may be considered a major source of air pollution in an uncontrolled state. As such a B.A.C.T. type standard would be applicable and will likely approximate the 2.9 lbs. VOC/gallon of coating applied or 90% removal of VOC from the exhaust stream. David 'M' Company cannot meet the afore-mentioned low solvent technology standard and will be obligated to propose the control device intended for use in meeting the 90% removal standard referenced above. While I cannot guaranty at this time the exact type of control device which David 'M' Company will install, I can state that both recovery and thermal destruction techniques capable of delivering 90% removal of VOC's from exhaust streams are being evaluated.

The attached schedule estimates the time periods associated with specifying the type of control device and ultimately installing same. Would you please advise me of the proposed schedule's acceptability. We believe this schedule to be a fair appraisal of the time necessary to complete the installation of a project of this magnitude. Should you require further information and/or assistance, please do not hesitate to contact me directly.

Sincerely,



Michael A. Ware
Graphics Div. Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

MAW:bjw
Attach.

Mike Ware

October 19, 1983

ESTIMATED
OVERALL COMPLIANCE SCHEDULE
DAVID M COMPANY VOC CONTROL PROJECT

PHASE

Data Accumulation	January 1, 1984
Spreader Enclosure - (1) For Temperature Evaluation	January 15, 1984
Process Modifications and Development	April 1, 1984
Methods Comparison/Analysis - Thermal Incineration Vs. Recovery	May 15, 1984
Final Specification Development	July 1, 1984
Final Specification/Quotations and Analysis	September 1, 1984
Appropriations Request Development and Approval	December 1, 1984
Oven Enclosure and Exhaust System Installation	March 1, 1985
Solvent Recovery Delivery and Installation	August 1, 1985
Operational Shakedown, Debugging and Compliance Testing	December 1, 1985

APPENDIX VI



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~999020~~ 240189

COPY

YOUR REF:

OUR REF: E 0379
S 7212

DATE: 2 August 1983

PROPOSAL

FOR

SOLVENT RECOVERY SYSTEM

TO BE LOCATED AT:

DAVID M. COMPANY

201 VALENTINE WAY
LONGWOOD, FLORIDA



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~900020~~ 240189

YOUR REF:

OUR REF: S 7212

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

SOLVENT RECOVERY PLANT

ACTIVATED CARBON ADSORPTION UNIT

<u>AIRFLOW</u>	25,000 scfm.
<u>AIR TEMPERATURE</u>	140° F.
<u>SOLVENT CAPACITY</u>	50 gph (360 lb/hr)
<u>SOLVENT</u>	Toluene

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL

David M. Company

Proposal No. S 7212

PLANT DESCRIPTION

The solvent laden air would be filtered to remove dust, cooled to approximately 95° F by the air cooler and then delivered by the fan to the three adsorbers.

Two of the three adsorbers would normally be handling the solvent laden air whilst the other would be regenerating with steam to extract the solvent from the carbon or would be held on "stand-by" following "steaming" and a short dry/cool period.

The adsorbers would cycle consecutively and automatically with override of the cycle time and steam input by gas analyser control. The proposed adsorbers are of annular bed design, the solvent air after filtering and cooling passes through an automatic inlet valve into the outer annulus of the adsorber, through the carbon bed and then exhausts to atmosphere through the inner annulus and the automatic exhaust valve.

At the end of an adsorbing period the air inlet and outlet valves close and an automatic steam valve allows steam to enter the inner annulus of the adsorber and pass through the carbon bed to the outer annulus from which point the steam together with solvent vapour extracted from the carbon passes to the condenser. During steaming some condensation occurs on the outer shell of the adsorber and is drained through filters to join the main condensate flow from the condenser to the decanter.

At the end of the steaming period the adsorber automatic steam valve shuts and the air inlet and outlet valves open to allow solvent laden air to again pass through the adsorber for a short period to dry and cool the bed (during this dry/cool period the solvent laden air is shut off from the other adsorber). The inlet valve of the dried/cooled adsorber would then close and the adsorber would be held on "stand-by" until the gas analyser, sampling the exhaust of the other adsorber, or the override timer, gives the signal for that adsorber to change to the steaming cycle, the stand-by adsorber being first brought back into service, so that there is no interruption to the adsorber cycle.

Any incondensibles from the condenser return through the condenser vent pipe to the inlet duct for recycling through the adsorbers. The condenser vent is fitted with high temperature alarm to indicate overloading from excessive steam flow or shortage or failure of the cooling water supply.

Safety features include adsorber pressure and vacuum relief valves and liquid seals.

David M. Company

Proposal No. S 7212

We would supply the following:

1. AIR FILTER - 25,000 scfm capacity

a) Pre-Filter Section

With removable and disposable fiberglass panels to protect and extend the life of the main filter.

b) Main Filter Section

With disposable filter elements of the HEPA type of fiberglass material, with an efficiency of better than 99% for particles down to 0.3 microns per standard tests.

The whole of the above would be incorporated in a galvanized steel case with access doors to facilitate servicing.

A differential resistance gauge would be fitted.

2. AIR COOLER - 25,000 scfm capacity.

Would be of finned tube design with copper tubes and aluminum fins to cool the inlet airstream from 140° F to approximately 95° F when supplied with cooling water at a temperature of maximum 86° F.

The air cooler coils would be housed in a galvanized steel case and would be drainable.

3. FAN AND MOTOR

The fan would be of the high efficiency type with airfoil impeller capable of handling 25,000 scfm of solvent laden air, when at an actual temperature of approximately 95° F, at a total pressure of approximately 18" w.g., allowing for 2" w.g. suction before the air filters.

The casing would be of substantial carbon steel construction and the unit would have spark-proof features to AMCA standards.

The impeller would be driven through a flexible coupling by a 125 HP explosion proof motor to Class 1, Group D, Division 1, 460 volt, 3 phase, 60 cycles.

A manually operated radial leaf damper would be fitted at the fan inlet for regulation of the airflow.

David M. Company

Proposal No. S 7212

4. THREE ADSORBERS - 6'8" diameter x approximately 10'0" high shell.

Each capable of handling 12,500 scfm of solvent laden air calculated at 70° F, when at an actual temperature of approximately 95° F.

The adsorbers would be of the annular carbon bed type with the carbon contained between stainless steel panels formed up into cylinders and reinforced by carbon steel stiffening bands. The activated carbon bed and screens would be supported upon a carbon steel tray to which would be connected the solvent laden air inlet valve, vapour piping to condenser and liquor drains. The air exhaust valve and the steam valve would be connected to the outlet branch of the adsorber.

The carbon bed and screens would be contained within a removable carbon steel shell, fitted with access and inspection holes with bolted cover plates. Flanged connections would be fitted to the tray so that the carbon could be easily and quickly run off into drums.

Vacuum and pressure relief valves would be provided for each adsorber.

Handrails would be fitted round the tops of the adsorbers, with a walkway over and between the adsorbers, and with an access ladder from ground level, in accordance with the requirements of OSHA.

5. ACTIVATED CARBON

We would provide the initial charge of approximately 6400 lbs of high quality carbon for each of the three adsorbers. The carbon would be manufactured by Sutcliffe Speakman and would be of the grade most suited to the required duty.

6. INLET AND OUTLET AIR VALVES - 26" diameter

We would provide an inlet and outlet valve for each adsorber 26" diameter of the mushroom type of substantial construction arranged for pneumatic operation through the automatic control mechanism.

Renewable seats would be fitted in the valve bodies and access holes with bolted cover plates provided. An indicator would be fitted to each valve so that the attitude of the valve could be easily ascertained.

David M. Company

Proposal No. S 7212

7. CONDENSER

To condense the steam and solvent vapours leaving an adsorber during the steaming period.

The unit would be of the tubular surface type with the shell in 304 stainless steel and the tubes in 316 stainless steel. The water boxes and tubeplates would be in carbon steel.

The condenser vent pipe (fitted with thermometer with high temperature alarm contacts) would be connected to the inlet duct so that any incondensibles displaced from the adsorbers would recycle.

8. ONE DECANTER - 3'0" diameter x 4'6" high

It would be of the self-adjusting decanting type to separate automatically the condenser water and solvent. The tank would be of carbon steel welded construction fitted with a bolted cover plate and all the necessary connections. A sight glass would be fitted in the solvent outlet.

9. TRUNKING

We would provide the interconnecting trunking between the air filter, air cooler, fan and adsorbers, but do not allow for any trunking prior to the inlet manifold of the air filter. The trunking would be of carbon steel of welded construction flanged at suitable intervals and be adequately supported.

10. THREE EXHAUSTS

An exhaust stack would be provided for each adsorber, of carbon steel welded construction, galvanized and would terminate about 6' above the adsorber platform.

11. PIPING AND VALVES

We would provide pipework to interconnect the above and manifold the service pipes to terminal flanges at the battery limits.

Vapour and liquor piping would be in 304 stainless steel, utility piping in carbon steel.

We include for all isolating valves, steam traps, pipe supports, etc.

David M. company

Proposal No. S 7212

12. PLANT CONTROL UNIT

Would consist of the following:

a) Gas Analyser (Infra-Red Type)

To measure the solvent content in the exhaust airstream of the adsorber next due for desorbing and to delay steaming until the set point was achieved. However, an adjustable override timer would be incorporated into the program to limit the period of such a delay.

A strip recorder for the exhaust gas concentration would be included and the unit would have a "high level" alarm.

b) Programmable Logic Controller

Of the solid state type which on receipt of the appropriate signals would cause the adsorber valves to sequence as appropriate. The program logic would be such that the next step would not be commenced until the previous one had been proven completed.

In the unlikely event of failure of the P.L.C. the solvent recovery plant could be operated manually by the use of pilot air solenoid valves inbuilt into the panel.

c) Annunciation Section

Of the 'Panalarm' type or similar to alarm:
"high level" - gas analyser output
"high temperature" - condenser vent

d) Control Panel

The above items would be mounted in a free-standing enclosure, suitable for indoor location in a non-hazardous area, assumed within 50' of the adsorbers.

The panel would be pre-wired and tubed to terminal points on the frame. We exclude the impulse tubing from the panel to the plant items.

David M. Company

Proposal No. S 7212

13. INSTRUMENTS

We would provide the following indicating type instruments which would be locally mounted on the plant:

- 4 Dial type thermometers
- 1 Steam pressure gauge
- 1 Differential pressure gauge (resistance through air filter)
- 1 4-Point pressure gauge (air pressure and suctions within the plant)

14. STRUCTURES

We would provide structures to support items of the plant.

15. PAINTING

Where applicable, plant items would be primer coated before shipment.

16. COMPRESSED AIR

You to supply compressed air, clean, dry and oil free at a pressure of approximately 80 psig.

17. DRAWINGS AND INSTRUCTIONS

We would provide three (3) copies of each of the following:

- Foundation drawing
- Flowsheet
- Plant arrangement drawing
- Major assemblies

Plus two (2) copies of:

- Erection drawings and instructions
- Operating and Maintenance Manual

A list of recommended spares would also be provided.



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: 908020 240189

YOUR REF:

OUR REF:

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

TERMS, PRICE AND CONDITIONS

PRICE OF PROPOSAL

PROPOSAL NO. 5 7212

For the supply of the whole of the items as specified in this proposal, delivered only to your works in Longwood, Florida (offloading from the transport and installation by others) would be the sum of approximately.....\$ 370,000.00

(THREE HUNDRED SEVENTY THOUSAND DOLLARS)

Any Federal, State or local taxes that may be applicable are excluded.

TERMS OF PAYMENT

25% of quoted price with order.

45% of quoted price in three (3) equal payments pursuant to invoices on the following basis:

- a) upon submission of preliminary drawings comprising plant arrangement and foundation drawings and flowsheet.
- b) upon submission of confirmation that the manufacture is at least 40% complete.
- c) upon submission of confirmation that the manufacture is at least 70% complete.

25% of quoted price upon despatch or when ready for despatch if despatch date is delayed by you.

5% of quoted price on start-up and successful operation, at latest three (3) months after despatch, or when ready for despatch, whichever is sooner.

Continued.....

David M. Company

Proposal No. S 7212

TERMS, PRICE AND CONDITIONS...continued

DESPATCH

Approximately six (6) months from receipt of order assuming that there are no delays due to changes in specifications or approval of drawings.

EXCLUSIONS

We do not include for any motor controls, electrical wiring, or any special cold weather protection or insulation, or any parts, materials, work for services of any kind whatsoever except those specifically mentioned in the body of the proposal.

STANDARD OF MANUFACTURE

The goods described in this proposal would be manufactured to normal commercial standards and be suitable for the duty specified. Should you desire manufacture to any specific standards, codes or inspection schedules other than those allowed for then we will be pleased to comply with your requests, but any extra charges so incurred would be in addition to the prices quoted.

VALIDITY OF PROPOSAL

This proposal is open for acceptance for a period of thirty (30) days from the date of this quotation and after this period confirmation must be obtained that the terms and conditions contained herein are still valid.

SUTCLIFFE SPEAKMAN INCORPORATED



William C. Moses
Vice President

E. & O. E.

WCM/pmd

David M. Company

Proposal No. S 7212

UTILITY REQUIREMENTS

The following are the approximate utility requirements of the plant when operating at design load conditions. The figures are subject to confirmation by our final design study.

STEAM.....1190 lb/hr, dry saturated quality
at 40 psig.

COOLING WATER.....170 gpm, inlet 86^o, maximum,
return 115^o.

POWER.....100 KwH.

COMPRESSED AIR.....20 scfh.

SPACE

The equipment, we estimate, would require a floor area of approximately 45' x 20'.

The height of the equipment would be approximately 18', but if it is to be located inside, then provision should be made for lifting of the adsorber shells.

APPENDIX 'A'

OUR PRELIMINARY PROPOSAL FOR THE TURNKEY INSTALLATION

OF THE PLANT QUOTED IN MAIN PROPOSAL NO. S 7212

1. FOUNDATIONS

We would provide concrete foundations for the appropriate items of our equipment.

Our price is conditional on the understanding that the ground is of suitable load bearing capacity, that no rock blasting or piling would be required and that all work would be above the water table.

2. MOTOR CONTROL CENTER

We include the starters for the fan motor, the water cooling tower fan and the recirculating water pumps, and which would be suitable for location in a non-hazardous area which we assume will be by the control panel.

3. WATER COOLING TOWER

Unosion Lubricator S? Chemicals

We would supply and install a water cooling tower designed to cool the required amount of water from 115° F to approximately 86° F. The tower would be of induced draft type and complete with fan and motor. The system would include two pumps with motors (one normally operating, one stand-by) and the water circulating piping and valves between the water cooling tower and the air cooler and condenser of the recovery plant, the water cooling tower to be at ground level and located adjacent to the recovery plant.

4. INSTALLATION

We include for the necessary labour, tools, moving and lifting gear including cranes, for the installation of the items of equipment specified in this proposal onto the foundation.

4. INSTALLATION...Continued

It is understood that the work would be undertaken during our normal working hours and that we and our sub-contractors or agents would be allowed free and unrestricted access to the site.

Any overtime worked at your request would be chargeable extra at the rate in force at the time.

You would be responsible for offloading from the transport any items delivered prior to the start of the installation work and provide suitable storage and protection for all parts close to the battery limits.

All temporary site support facilities including but not limited to water, electricity, steam, drain, office and telephone would be provided by you at your expense, adjacent to the site.

Note that the quoted price assumes we will have continuity of work on site.

5. SUPERVISION

We include for the services of our own suitably qualified supervisor to supervise and give instructions to the labour provided by our sub-contractors, to secure the installation of the equipment in a competent manner.

6. IMPULSE TUBING

The pneumatic tubing from the appropriate plant items to the control panel would be supplied and installed in a workmanlike manner. It is assumed that the tubing run is a maximum of fifty feet.

7. ELECTRICAL WIRING

We would supply and install all necessary electrical wiring, cable trays, etc. to interconnect the various items of plant to the motor control center and the control panel, assuming a maximum cable run of fifty feet from the adsorbers. Where appropriate, the wiring would be to explosion proof, classification.

David M. Company

Proposal No. S 7212

8. INSULATION/WINTERIZATION

Insulation and heat tracing of appropriate equipment within our battery limits effected, assuming a minimum ambient temperature of 25° F.

9. START-UP

We include for the services of a skilled engineer for a period of two (2) consecutive weeks on site to start-up the plant and provide instructions to your personnel on its operation.

Any extra time required by you or non-continulty in attendance would be at additional cost.

10. INSURANCE

We would provide and maintain the following insurances and would require any and all sub-contractors employed by us at your site to procure and maintain the same type and amount of insurance.

- a) Workman's Compensation and employer's Liability
 - \$100,000.00 each accident and aggregate disease.
- b) Comprehensive General Liability
 - Bodily injury and Property damage of up to \$1,000,000 combined single limit.
- c) Comprehensive Automobile Liability
 - To cover owned, non-owned, leased and hired cars, combined single limit \$1,000,000.

Certificates of the above shall be provided to you prior to the start of our on site work.

In addition to the above and in order to insure the equipment from its delivery to your site, through installation to your acceptance of the plant, we would take out "all risks" insurance plus "builders risk" and "installation floaters" at the appropriate time and at your expense.

David M. Company

Proposal No. S 7212

BUDGETARY PRICE

Our approximate additional price to that quoted in Main Proposal No. S 7212 for the turnkey installation of the plant as detailed in this Appendix 'A' would be the sum of\$ 160,000.00.

(ONE HUNDRED SIXTY THOUSAND DOLLARS)

COMPLETION

We would expect to supply and install the specified plant in approximately nine (9) months from receipt of order.

TERMS AND CONDITIONS

Otherwise as Main Proposal No. S 7212.

A handwritten signature in black ink, appearing to be 'D. M. Company', is located in the lower right quadrant of the page.

SOLVENT RECOVERY

WITH

ACTIVE CARBON

SUTCLIFFE SPEAKMAN, INC.

Sutcliffe Speakman have specialised for over 50 years in the design and manufacture of Solvent Recovery plant. Their unique position in the industry stems from a background of almost 80 years experience in the manufacture of active carbon for general adsorption and catalytic applications.

SOLVENT RECOVERY WITH ACTIVE CARBON

Our recovery plants are in world-wide use in a great variety of solvent-using industries, ranging from coating, impregnation and printing of materials to the manufacture of man made fibres.

Our active carbon plants for the recovery of industrial solvents, from air or gas mixtures, can give a recovery efficiency of at least 99% by volume depending on the conditions. In addition to vapour phase recovery, active carbon plants can efficiently extract and recover solvents and organic compounds from water and other liquid mixtures.

MONEY SAVING

Recovery plants save industry millions a year and the dramatic savings on solvent costs ensure more economic production.

CLEAN AIR

The adsorption efficiency of Sutcliffe Speakman plant is extremely high, effecting an important contribution to clean air.

LIQUID EFFLUENTS

Can also be purified with an active carbon plant.

WORKING CONDITIONS

A general improvement in working and safety conditions often results from the effective collection system required to achieve high overall efficiency.

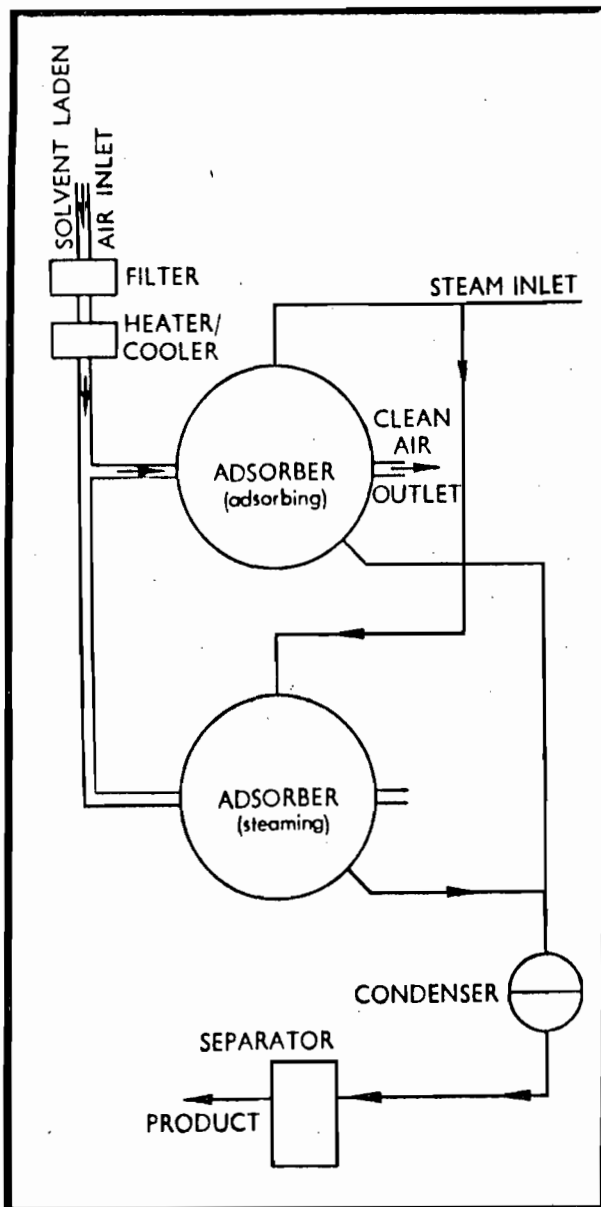
WHEN IS IT WORTHWHILE ?

If you are using more than 200 gallons a week of alcohols, esters, ethers, ketones, hydrocarbons, chlorinated compounds and other organic solvents, it is certainly worth giving preliminary consideration to the economic potential of installing recovery plant. The actual economics depend upon the cost of new solvent, the concentration of solvent in air entering the plant, the number of hours per week which the process will operate and the possible ancillary equipment which may be necessary to prepare the recovered solvent for re-use. This may be required if the solvent is water miscible or where a mixture of different solvents may be recovered together and necessitate separation.

Sometimes solvent quantities as low as 1 gallon per hour can be economically recovered. WRITE TO US, WE WILL ADVISE WHETHER A DETAILED STUDY IS JUSTIFIED.

GASEOUS PHASE ADSORPTION

BASIC FLOW DIAGRAM



HOW IT WORKS

The Sutcliffe Speakman system draws air laden with solvent vapour through a ducting into the plant when, if necessary, it is first filtered to remove dust or other contaminants and then heated or cooled to a suitable temperature before passing into an adsorber which contains a bed of active carbon. The solvent vapour is adsorbed on the carbon and the clean air is discharged to atmosphere. When the carbon bed has adsorbed the designed charge of solvent, the vapour/air stream is switched to a second adsorber and steam at low pressure is introduced to the charged adsorber which causes the carbon to release, in vapour form, the solvent which it adsorbed. The steam/solvent vapour mixture is then condensed and the solvent recovered.

If the solvent is insoluble in water, a separator is provided to divide the condensate into a solvent layer and water layer which automatically leave the separator by gravity flow. If the solvent is water soluble, the condensate is taken to a distillation unit or liquid phase adsorption system for final separation.

CARBONS

The wide range of Sutcliffe Speakman carbons developed specially for solvent recovery give a high degree of adsorption efficiency at relatively high velocities. These carbons also have low retention characteristics, so that steam usage for the regeneration of beds is minimised. The most suitable carbon is selected to meet the particular conditions under which the plant will operate.

PLANT

Sutcliffe Speakman design their adsorbers to give maximum surface area of carbon, thereby ensuring the minimum resistance to air-flow, and a relatively lower power consumption. Adsorbers are designed with static or rotary beds.

Automatic operation of the adsorber valves with variable time cycle, is effected by instrumentation or by the robust Sutcliffe Speakman standard control unit.

In normal applications the adsorbers are not subjected to pressures of more than 0.5 lbs/square inch. Where necessary, plants can be designed for operating at higher pressures.

If required, plant can be designed for further extension to suit clients needs.

The rotary bed plants and a range of static units can be supplied as fully "packaged" plants. Wherever practicable, all plants are pre-assembled in our works before despatch. Full erection and maintenance services are available including contract visits by Technical Service Engineers.

SOLVENT CONCENTRATION IN AIR

The adjacent diagram shows the quantities of solvent in relation to air flow at given volume/volume concentrations, corrected to the normal temperature and pressure conditions at the inlet to the recovery plant, with indication of solvent value per annum.

It is important that the concentration of solvent in the air stream from a process is, where applicable, well below the lower explosive limit of that particular solvent vapour in air, to ensure a satisfactory operational safety margin. It is normally considered that the quantity of air flow from a process should be sufficient to ensure that the concentration of solvent vapour in air is no more than a maximum of 50% of the lower explosive limit.

Conversely it is usually necessary to ensure that the concentration is not too low, or the size of plant required to handle air flow may be larger than can be justified economically in relation to the quantity of solvent to be recovered. The ideal concentration range is usually between 20%-40% of the lower explosive limit.

Sutcliffe Speakman technical staff are available to advise on the collection of solvent from your process to ensure that maximum overall recovery efficiency is achieved, and that the desirable air flow from an economic point of view is sufficient to collect the solvent vapours effectively.

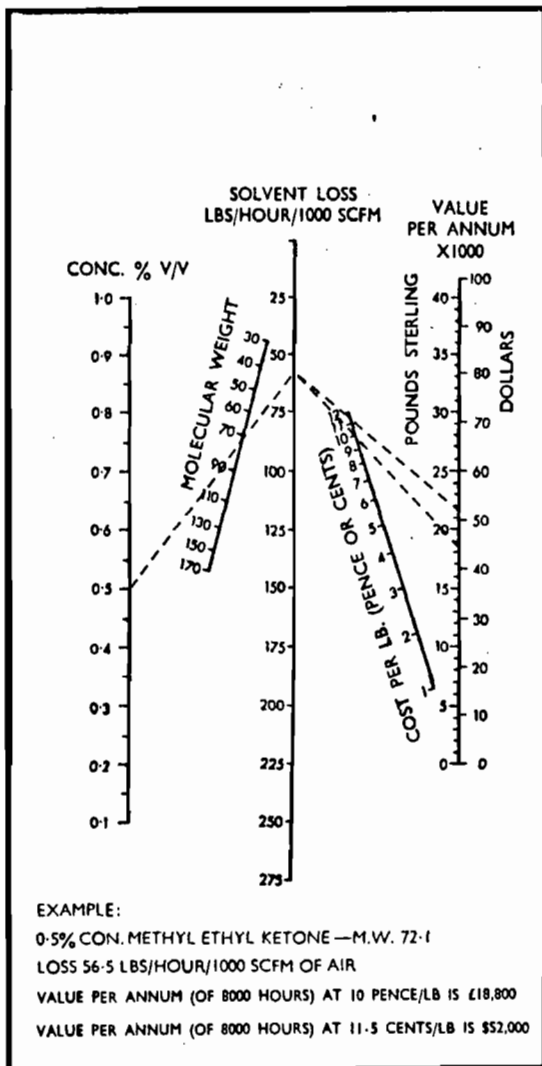


TABLE OF MOLECULAR WEIGHTS AND LOWER EXPLOSIVE LIMITS FOR SOME TYPICAL INDUSTRIAL SOLVENTS

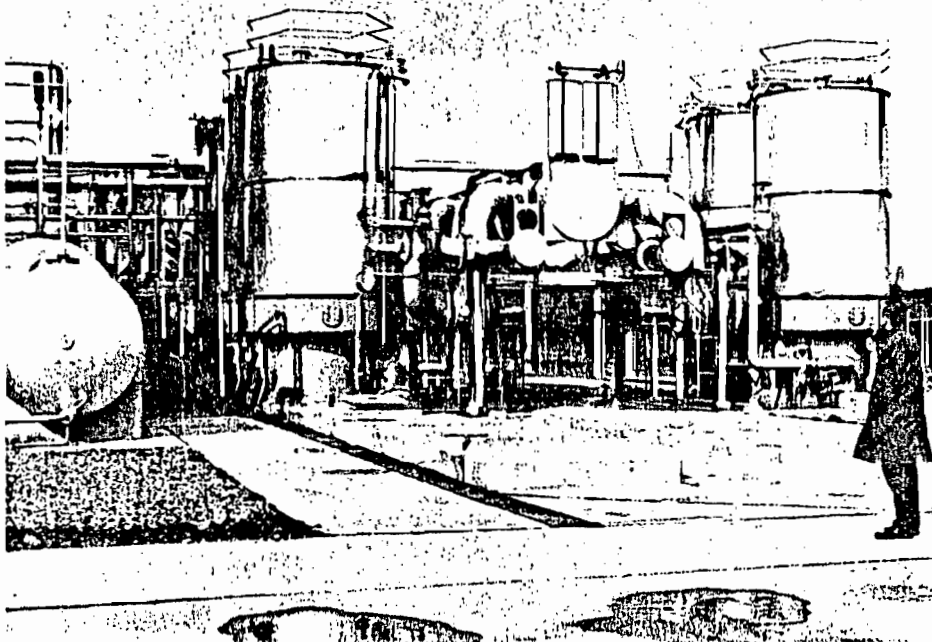
	A	B		A	B
Acetone	58.08	2.15	Methyl Acetate	74.08	4.1
Benzene	78.11	1.4	Methyl Alcohol	32.04	6.72
Butyl Acetate (Normal)	116.16	1.7	Methyl Ethyl Ketone	72.10	1.81
Butyl Acetate (Iso)	116.16	2.4	Methylene Chloride	84.93	Non-flam.
Butyl Alcohol (Normal)	74.12	1.45	Nonane	128.25	0.74
Butyl Alcohol (Iso)	74.12	1.68	Octane	114.23	0.95
Carbon Disulphide	76.13	1.0	Pentane (Normal)	72.15	1.3
Carbon Tetrachloride	153.84	Non-flam.	Pentane (Iso)	72.09	1.3
Chloroform	119.39	Non-flam.	Perchloroethylene	165.85	Non-flam.
Cyclohexane	84.16	1.35	Propyl Acetate (Normal)	102.3	2.0
Decane (Normal)	142.28	0.7	Propyl Acetate (Iso)	102.3	2.0
Dichloroethylene	97	9.7	Propyl Alcohol (Normal)	60.09	2.15
Ether (Diethyl)	74.12	1.85	Propyl Alcohol (Iso)	60.09	2.02
Ethyl Acetate	88.10	2.25	Tetrachlorethane	167.86	Non-flam.
Ethyl Alcohol	46.07	3.3	Tetrahydrofuran	72.10	1.84
Ethylene Dichloride	98.97	6.2	Toluene	92.13	1.3
Furfural	98.08	2.1	Trichloroethylene	131.4	Non-flam.
Heptane (Normal)	100.2	1.0	Xylene	108.16	1.0
Hexane (Normal)	86.17	1.25			

A Mol. Wt.
 B Lower Explosive Limit in Air % V/V at 20°C and approx. atmospheric Pressure

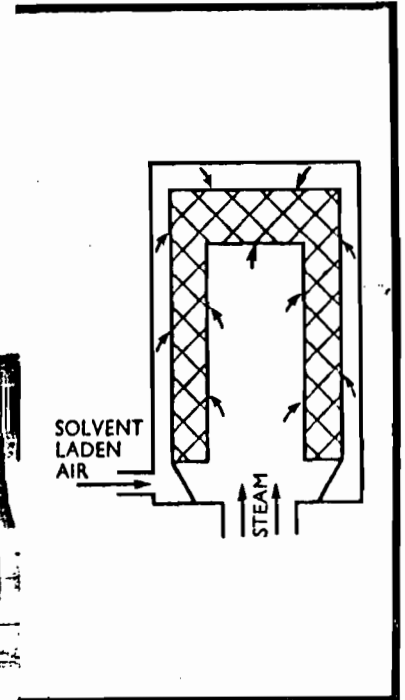
Sutcliffe Speakman manufacture two principal types of Gaseous Phase Recovery Plant— Both types employ active carbon.

STATIC BED PLANT

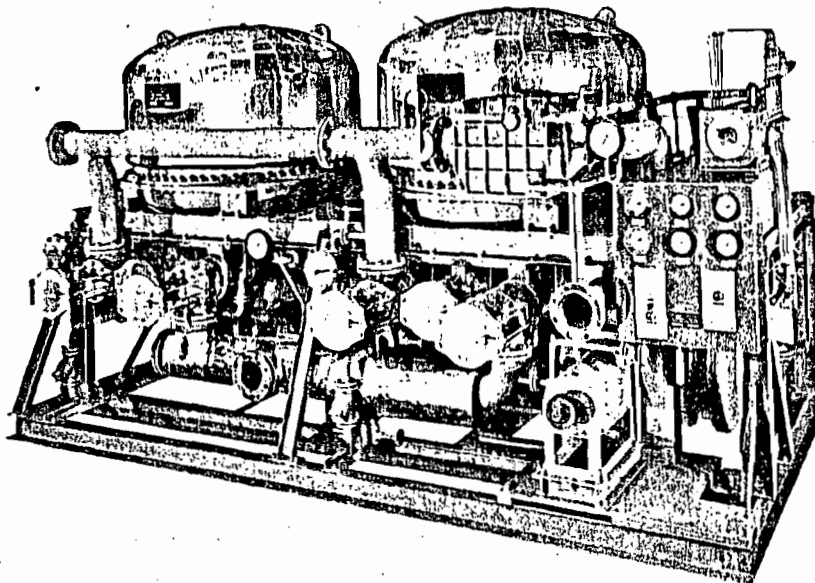
A static plant usually consists of two or more adsorbers each containing a vertical annular or flat bed of active carbon. One or more of the adsorbers receives the solvent laden air whilst one or more are being steamed out to recover the solvent. The adsorbers are suitably valved and cycle automatically so that the process is continuous. The smaller plants can be of packaged design, skid mounted. Plants are individually designed to meet customers specifications.



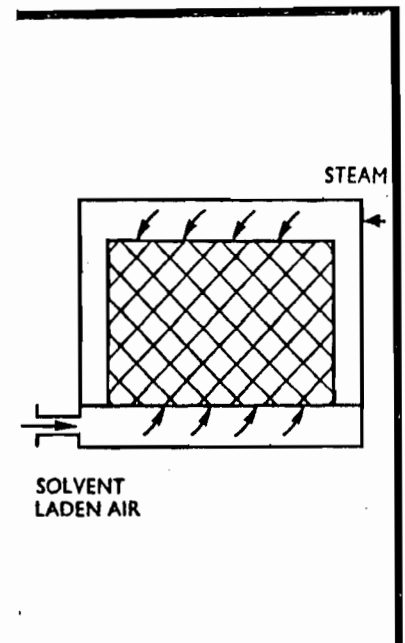
Adsorption section of a large recovery complex at an Acetate Fibre Factory in Belgium. The plant is arranged for future extension.



ANNULAR BED



Packaged type fully automatic plant of cylindrical flat bed design.



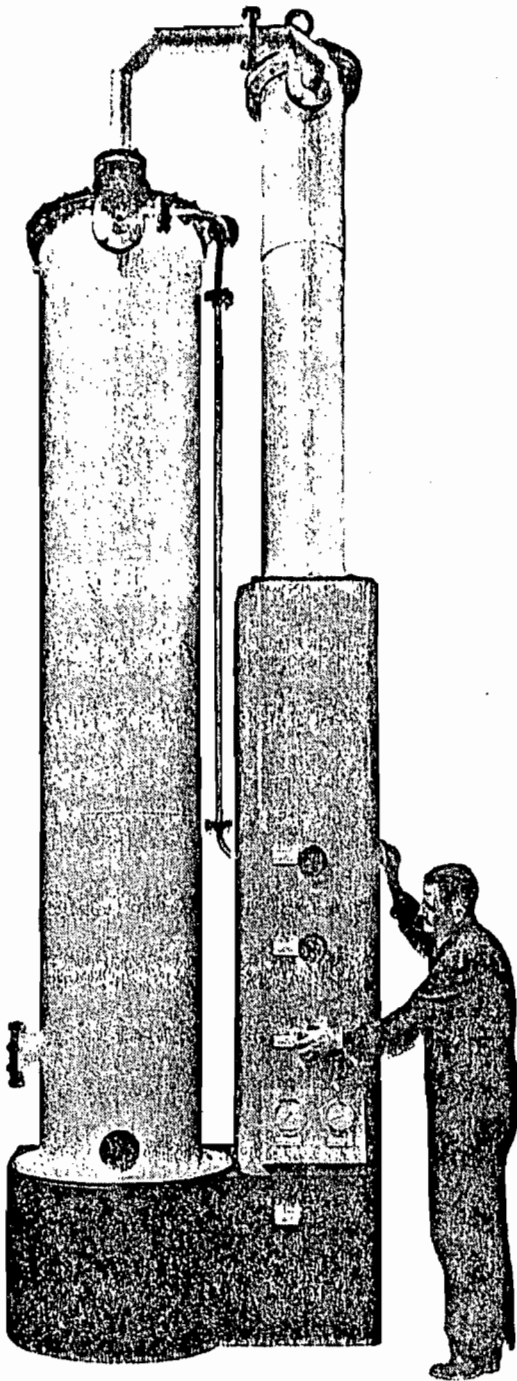
FLAT BED

LIQUID PHASE ADSORPTION

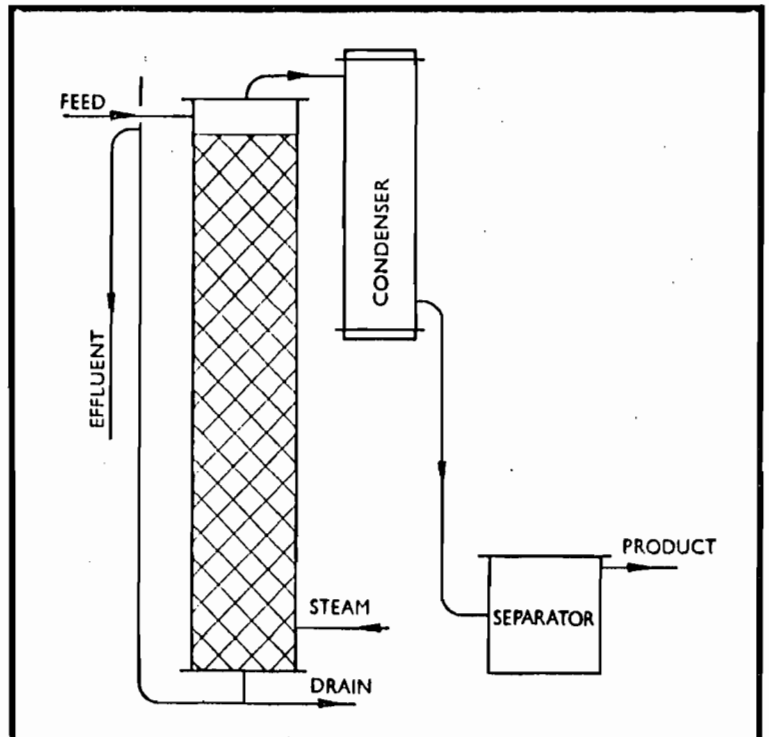
This system uses active carbon for the adsorption of small amounts of certain water-miscible solvents from solutions with water. The weak solvent/water liquor is passed through an active carbon bed and, when the carbon has adsorbed the designed charge of solvent, the solvent is distilled off by our patented system. This process effects substantial increase in concentration of a weak solvent/water solution, the water in the inlet feed passing through the bed and to effluent drain. During steaming of the carbon a small amount of water is introduced and the products of steaming taken to a separating tank from which the water layer recycles through the system.

In certain cases the carbon can be used to adsorb selectively a particular solvent component from a mixture of solvents with water.

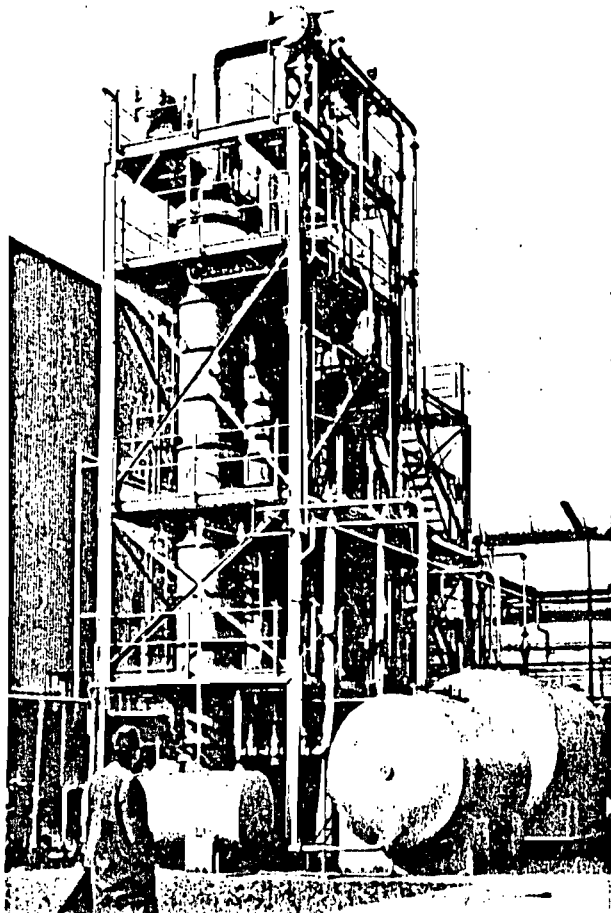
A particular feature of this system is the extreme simplicity of operation and control which can be manual or automatic.



Liquid phase adsorption unit.



DISTILLATION



Section of a large distillation complex.

Sutcliffe Speakman supply a complete range of distillation equipment, which may be required in the separation of water-miscible solvents from the products of recovery of the main adsorption plant, or for other purposes.

The range of distillation equipment includes both perforated plate and bubble cap designs, and the operations can be automatically controlled in relation to steam input and reflux ratio.

Stills can be supplied either for continuous or batch operation.

There is also a range of small units for the recovery of solvents from contaminated mixtures, i.e. by extraction of residues from waste lacquers or in cases where solvents are used for washing down printing and paint machines, etc., the solvent can be cleaned for further use.

ENQUIRIES

In submitting enquiries for evaluation, it is helpful if the following preliminary details can be submitted.

Gas or liquid flow rate

Operating pressures

Temperature and relative humidity of air or gas stream

Quantity, nature and value of compounds to be extracted

Other substances in air, gas or liquid stream of a contaminant or corrosive nature

Normal operating time of process

Details of steam, cooling water, power and compressed air services available.

TYPICAL SOLVENT RECOVERY APPLICATIONS

Acetate Fibre · Insulated Cables · Balata Belting · Leather Cloth · Hydrocarbon Extraction (Natural Gas) · Metallic Foil Printing and Lacquering · Compressed Asbestos Fibre Products
Oil Extraction · Degreasing · Paper Impregnating and Lacquering · De-sulphurising of Gas
Paper Printing · Dry Cleaning · Pressure-Sensitive Tapes · Explosives · Rotogravure Printing
Film Casting · Rubber-Dipped Goods · Rubber-Spread Goods · Surgical Bandages
Shoes, etc. · Solvent Manufacture · Tail Gas from Petroleum Wells · Viscose Fibre
Viscose and Acetate Film · Weatherproof Sheeting · Process Off-Gas



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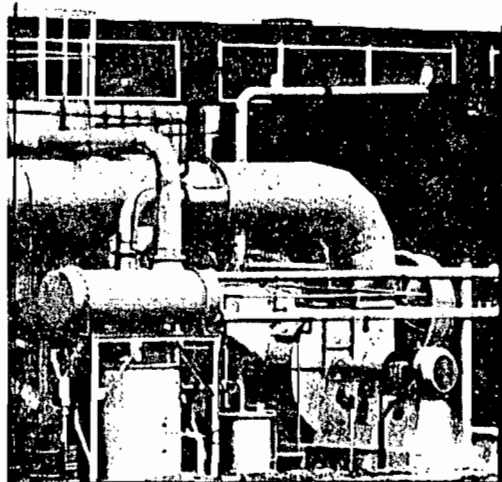
SUITE 200, HEAVER PLAZA,
1301 YORK RD. LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: 908020

The many facets of activated carbon

Reprinted from
Chemical Age 26 June 1981

activated carbon

and business advantages of solvent recovery



toluene and SBP2.

annular bed is the most efficient when it comes to steaming the solvent off the carbon bed (desorption). This is because the steam enters the inner annulus of the adsorber working its way outwards in contrast to other

designs where the adsorber vessels have to be efficiently lagged to prevent waste of heat energy. With the annular bed design by the time the steam has worked its way through the carbon bed to the outer annulus it has performed its useful work, and the cooler outer shell of the adsorber vessel serves as an air condenser and relieves the load on the main condenser with regard to the amount of cooling water required.

The basic components of a typical solvent recovery plant are:

- Air filter (to remove any airborne particulate matter in the solvent-laden air stream).
- Air cooler/heater (to temperature adjust air stream for safe and good adsorption on the carbon bed).
- Main fan(s) (to provide correct rate of airflow through the adsorbers).
- Activated carbon adsorbers (where the solvent(s) are adsorbed/desorbed).
- Condenser (to condense solvent vapour and steam from the desorption cycle).

- Separator (decanter) — (where non-miscible solvent/water mixture separates).

- Product tank (for collection of solvent from the separator).

In operation the solvent-laden air is passed to two of the adsorbers while the solvent is desorbed (by steaming) in the third adsorber.

The process operates automatically and continuously, usually with infra-red gas analyser control to ensure optimum efficiency, and each adsorber consecutively undergoes an adsorption and desorption cycle.

The simplest type of solvent recovery is where there is a single solvent which is non-miscible with water, eg, xylene. This solvent will collect as the upper layer in the decanter and, being virtually insoluble in water, can simply be decanted for re-use whilst the lower (water) layer is passed to drain.

If, however, a mixture of solvents is being recovered and some of these are miscible with water then additional plant is necessary to separate the solvents. This usually takes the form of distillation and dehydration equipment depending on the solvents involved and the final specification required by the customer for the recovered solvents.

Each solvent recovery system has its own particular technique for recovery, but the basic adsorption/desorption on activated carbon is common to all. When designing a solvent recovery plant for a particular set of conditions one must have detailed information on such parameters as airflow, air temperature, humidity, solvent concentrations, maximum/minimum process conditions, and details of any impurities in the solvent-laden air stream. The last factor can be particularly important since it can have an important bearing on materials of construction and expected carbon life.

There are several aspects which require careful attention to ensure that the best efficiency of solvent recovery is achieved and that at the same time minimum energy is consumed in the running of the plant. The first practical step which can be taken to maximise recovery efficiency is to present as high a solvent concentration as is safely possible with minimum airflow.

This will be largely governed by the type of process upstream of the

Table 1: Some typical results of solvent adsorption by carbons

Solvent	Concentration (%V/V)	*Initial adsorption (%W/W)	§Cyclic adsorption (%W/W)	**Steam ratio	Boiling point range (°C)
Methylene chloride	1.0	28.3	17.3	1.4	40.5
Arklone P	0.5	44.9	20.8	1.4	47.6
Acetone	1.0	20.3	12.5	2.3	56
Tetrahydrofuran	0.5	22.0	9.0	2.0	66
Hexane	0.48	21.3	8.2	3.5	68.7
Ethyl acetate	0.5	27.6	13.6	2.1	77.2
Trichlorethylene	0.5	44.6	19.9	1.8	86.7
n-Heptane	0.12	22.4	5.9	4.3	98
Toluene	0.4	23.3	9.6	3.5	110
Methyl isobutyl ketone	0.2	22.0	9.0	3.5	115.9
Mixed rubber solvent	0.3	25.7	10.3	3.8	120-160
Shellsol E	0.3	35.6	11.9	3.7	153-193
3, 3, 5-trimethyl cyclohexyl acetate	0.1	36.4	7.9	4.5	206

*Initial adsorption refers to the solvent adsorbed by new carbon to 'breakpoint', i.e. when solvent can be detected in the effluent air stream. The 'slip' level at this point is normally of the order of 20 ppm.

§Cyclic adsorption refers to the solvent which can be adsorbed and desorbed consistently at an economical steam consumption.

**Steam ratio refers to the total steam used to desorb the solvent from the adsorbent and is an overall figure inclusive of condensate and moisture condensed on the carbon bed.

SUTCLIFFE SPEAKMAN INC

S E N S O R C O N T R O L S

for SOLVENT RECOVERY

WITH PARTICULAR REFERENCE
TO THE USE OF INFRA-RED GAS ANALYSERS

Extract from paper prepared by
Dr. Robert D. Hill
Previously of Chemical Plant Division
Sutcliffe Speakman & Co. Ltd.

and

Presented at the 30th Annual Meeting of the
Gravure Technical Association
Chicago, April, 1979

There are many different types of sensors one can use on a solvent recovery plant but all function either to report information and/or to act on that information to control the solvent recovery plant's operation in some way. For example, proximity sensors can be used to sense whether a valve is open or shut, and flow metering and LEL meters can be used in conjunction with each other to measure the total solvent presented to an adsorber. My remarks will concentrate on using sensor control to optimize the conservation of energy, and maximize the efficiency of solvent recovery, and the role of the infra-red gas analyser in helping to achieve these aims.

The first practical step we can take in maximizing solvent recovery plant efficiency is to ensure that we are presenting, safely, as high a solvent load as possible in the minimum airflow passed to the solvent recovery plant. To a large extent this will be governed by the type of process upstream of the solvent recovery plant. Wherever possible, and economically achievable, recirculation of a portion of the solvent laden air should be considered. This will significantly increase the efficiency of solvent recovery across the carbon bed. The examples given below illustrate the typical improvement in efficiency of recovery of solvent which can be achieved:

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- (a) Inlet concentration 1000 ppm
Outlet concentration 35 ppm

$$\eta = \frac{1000-35}{1000} \times 100\% = 96.5\%$$

- (b) Inlet concentration 2000 ppm
Outlet concentration 35 ppm

$$\eta = \frac{2000-35}{2000} \times 100\% = 98.25\%$$

The side benefits from ensuring maximum concentration of solvent vapor that is presented to the solvent recovery plant are that it will reduce the size of the solvent recovery plant required and with it the capital outlay, space needed, and services required to run it. The sensors applicable to this area of plant operation are typically LEL meters coupled to variable damper controls which will vary proportion of solvent laden air from the process which is recirculated and diluted with fresh air.

Let us now consider in some detail the adsorption/desorption cycles which are at the heart of the solvent recovery plant's operation.

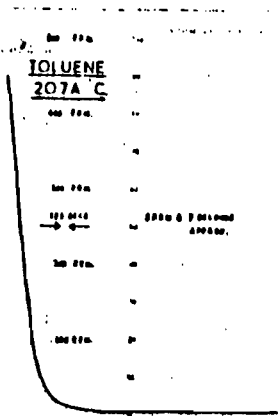


Figure 1

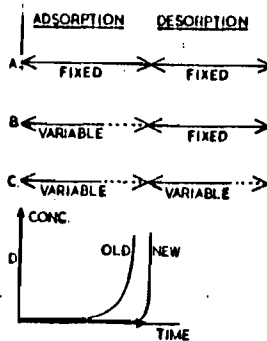


Figure 2

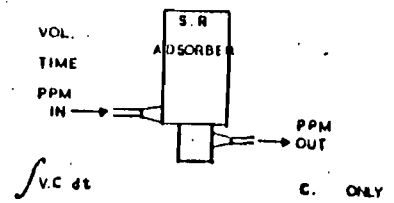


Figure 3

In Figure 1 we see the typical rise in solvent concentration as solvent (in this example - Toluene) starts to "slip" through the bed of activated carbon as an adsorption cycle nears its

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end. It can be seen that there is a rapid rise in solvent concentration - about 100 ppm in around 135 sec. With the particular sensor favored by Sutcliffe Speakman a rise of 5 ppm can readily be detected, and when a pre-selected concentration is reached (usually around 50 - 75 ppm) the sensor produces a signal which is used to initiate an automatic cycle controller which changes over the particular adsorber vessel being monitored from "Adsorb" to "Desorb" mode by opening/closing appropriate valves.

Let us now examine this process more closely. In Figure 2, on Line A, we have an "ideal" situation where a constant concentration of solvent in a constant airflow is presented to the solvent recovery plant. Provided constant conditions are maintained and the activated carbon's adsorptive properties also remain constant, then it is merely a matter of timing the length of the adsorption and steaming cycles to achieve optimum operating efficiency. This ideal situation is rare. What is frequently encountered in practice is that a variable concentration of solvent is presented to the solvent recovery plant. Now, if the adsorption cycle has been time controlled for the full designed load of solvent, and if, for example, half that load has been adsorbed on to the carbon bed in the preset time, not only will the carbon not be utilized fully, but an excessive amount of steam also will be consumed in the desorption cycle. As much as one third of the steam used for desorption could, for example, be wasted.

In Line B of Figure 2 we have the situation where we must be able to sense when the active carbon has adsorbed the optimum amount of solvent, and solvent "slip" is starting to occur through the carbon bed. In this case the length of the adsorbing cycle will be a variable and this is represented by the dotted part of the line in the figure.

However, what will be the situation if, in addition to the solvent concentration varying, we have a process in which there are airborne contaminants with the solvent which cause an unavoidable and progressive deterioration with time of the activated carbon (irrespective of the design or manufacture of the plant or source of carbon)? In some large, multi-adsorber plants associated with such processes it is often the practice to overhaul one adsorber at a time and to replace the poisoned carbon with a charge of new carbon. In this situation one can have an adsorber containing new carbon working in conjunction with one containing old carbon.

On Line D of Figure 2 we see a graphical representation of how the "slip" curves for old and new carbon compare.

In Figure 3 we see the two basic approaches used to measure when an adsorption cycle should be terminated. On the left side

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as solvent laden air enters the adsorber we can measure solvent concentration and flow rate, and compute, by integration with respect to time, the total quantity of solvent passed to the adsorber. When a predetermined amount of solvent has been passed a signal is generated which is used to change over the adsorber from Adsorb to Desorb mode. This approach, as used by some companies, seems to be a fundamentally less reliable approach in that it assumes a given carbon performance at all times. Furthermore, it can never give any useful information about what is happening inside the adsorber.

For example, an unusually high "slip" of solvent could indicate that valve seatings require renewal. It is considered that the fundamentally correct approach is to sense what is happening on the exhaust side of the adsorber and use this information for control purposes. It should be noted that this entails measuring only one parameter (solvent concentration in the air stream) against time. Now, whereas the inlet concentration is typically in the 1000 - 3000 ppm range we are now looking at a 0 - 100 ppm range on the outlet of the adsorber. For optimum efficiency we need a sensor which will be capable of measuring very low ppm's, have a fast response, be stable, be accurate, have high reproducibility, not be affected by water vapor and by other extraneous factors which are frequently encountered in the industrial situation. This tends to exclude the use of LEL meters (too insensitive), gas chromatography (slow response and not continuous), and flame ionization detectors, and brings us to the infra-red gas analyser.

Many gases absorb energy in the infra-red region because of the resonant molecular vibrations within the molecules of gas or solvent vapor. These resonances are highly specific and give rise to I-R spectra which are characteristic of the substances in question. The I-R spectrum of a molecule has been likened to the human finger print by means of which one specific wavelength depends on the concentration of absorbing molecules placed between the infra-red source and the detector, and this fact is utilized to determine the concentration of a selected component in a gas or vapor mixture.

Because of the complexity of typical gaseous infra-red spectra, some overlapping of absorption bands frequently occurs, and it is usually necessary to select a very narrow wavelength band to avoid cross sensitivity. Figure 4 shows the 1-8 microns region of the I-R spectrum of toluene and shows several characteristic absorption bands for this solvent.

There are several types of I-R gas analysers potentially available for use, and broadly they can be divided into single wavelength/single beam; single wavelength/double beam; and double wavelength/single beam instruments. In my experience all two-beam energy absorbing systems suffer from the inherent defect of

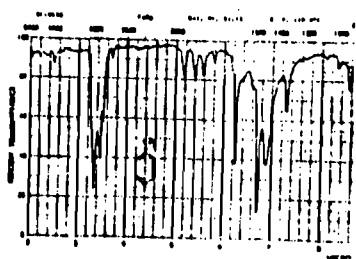


Figure 4

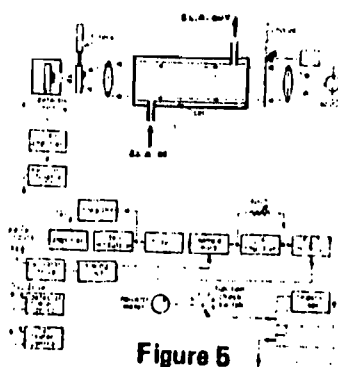


Figure 5

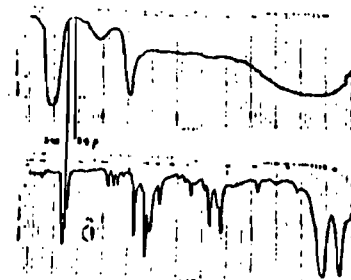


Figure 6

being sensitive to background absorptions caused by fouling of the sample cell windows, broad band absorptions in the sample gas (such as that caused by water vapor), and temperature differences between sample and reference beams. However, most of the disadvantages of the two beam system can be eliminated without losing the advantage of a continuous reference measurement by examining, alternately in time, the I-R energy transmission at the peak absorption wavelength of the measured component and that at a nearby wavelength which represents the "background" energy, but which is itself unaffected by the measured component. This can be achieved with a two wavelength/single beam instrument.

Such a system is shown in Figure 5. By having the source and detector units in optical alignment with each other, and arranging for a pair of very narrow band pass interference filters to be switched alternately in time into the beam (actually at 6 Hz) it is possible to produce a sensitive, accurate, reliable, I-R gas analyser which is immune from many of the extraneous factors which can plague other types of I-R gas analysers.

Let us now turn to the critical area of deciding what wavelengths to select for the solvent(s) in question. This is an area to which, I believe, too little consideration is often given - probably because it calls for expert knowledge in the field of infra-red spectroscopy.

If the solvent is toluene there are, as we can see from the I-R spectrum for this solvent (lower spectrum in Figure 6) several bands to choose from. Or are there? If we look at the spectrum of water (upper spectrum in Figure 6) we can see that there are several regions where water is absorbing energy strongly and has the potential ability to interfere. We must remember also that we are wanting to detect a 50 ppm level (say) which is only a 0.005% v/v concentration whereas if there is a 40% R.H. situation with respect to water vapor that corresponds to a 2.9% v/v concentration of water vapor - i.e. 580 times greater! The possibility of interference from water is obvious. However, if we select wavelengths 3.45 microns for toluene and 3.9 microns for the reference, then we can see that we shall meet the best criteria for having the solvent and reference wavelengths near to each other, but with the latter at a point of least interference from water. In this way we can ensure that even 100% RH will not affect this sensor's performance in controlling the solvent recovery plant.

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Figure 7



Figure 8



Figure 9



To illustrate just how complex the I-R spectrum of water vapor really is and how the effect of raised temperature can cause the molecules to become more agitated and resonate more strongly, Figure 7 shows the I-R spectrum for water vapor at 100° C, using an expanded scale between 5 and 8 microns. It will readily be appreciated that it is most important that the designer of a solvent recovery plant must have access to such specialized knowledge if the best sensor control design is to be achieved.

Let us now consider some other commonly used solvents: Figure 8 shows the I-R spectra for acetone and methyl ethyl ketone. Here the CH stretching absorption band at around 3.5 microns is relatively weak and it is usual to select the C=O (carbonyl) band at about 5.8 microns instead.

Figure 9 shows the I-R spectra for methyl alcohol and ethyl alcohol. Here one can use the CH band at around 3.5 microns as both alcohols have strong enough absorption bands.



Figure 10



Figure 10 shows the I-R spectra for ethyl acetate and hexane. Here we have a good illustration of the relative weakness of the CH band at 3.5 microns for ethyl acetate and the corresponding strength of the CH band for hexane at the same point in the spectrum.

In conclusion, in order to get the best out of a solvent recovery plant the following objectives should be achieved:

- (1) Maximize the solvent concentration and minimize the airflow within safe/economic limits.

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(2) Use I-R Gas Analyser control of the adsorption cycle to nullify the effects of varying solvent load and possibly also varying air flow in order to achieve the fullest use of the charge of active carbon in any particular adsorber.

(3) Select a 2 wavelength/single beam instrument which can guarantee:

- (a) immunity from water vapor
- (b) Virtual immunity from all normal extraneous factors such as voltage variation, temperature change, etc.
- (c) fast and accurate response
- (d) is fully automatic and requires virtually no attention.

(4) If the carbon is likely to have its activity modified by the process gas/vapor stream, then control the steaming cycle also by sensing a falling concentration of solvent in the steam/solvent mixture coming from an individual adsorber.

As manufacturers of their own activated carbon for the past 75 years plus solvent recovery plant which embraces the sensor technology, I have described, I believe, that Sutcliffe Speakman have one of the best systems for optimizing efficiency and minimizing the use of energy.



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~998029~~ 240189

YOUR REF:

OUR REF:

DATE: 4 August 1983

DESIGN FOR EXPANDABILITY

Mr. Michael A. Ware
Environmental Coordinator
Wheelabrator-Frye Incorporated
Graphic Supplies Division
Chemicals and Coatings group
2010 Indiana Street
Racine, Wisconsin 53405

Dear Mr. Ware:

We have pleasure in referring to the discussions with your colleagues and yourself during the visit to David M. Company, Longwood, last Friday, and we appreciated the opportunity to discuss the project for the recovery of toluene from the three rubber spreading machines.

For your preliminary information we have pleasure in enclosing our Preliminary Proposal No. S 7212, in triplicate, for a plant to recover up to 50 gallons per hour of toluene from 25,000 scfm airflow, as you requested, for your initial studies. We understand that consideration is being given to the possibility of making changes to the process operations to reduce the peak evaporation rates of solvent so that the capacity of the recovery plant might be reduced.

The proposed plant is quoted on the basis of supply only, with the addition of a preliminary proposal, Appendix 'A', for the turnkey installation of the plant which we would be pleased to firm up after further study of site conditions.

The proposed plant is very similar to a considerable number of installations that we have supplied for operation with rubber spreading machines in which particular industry we have over 40 years experience.

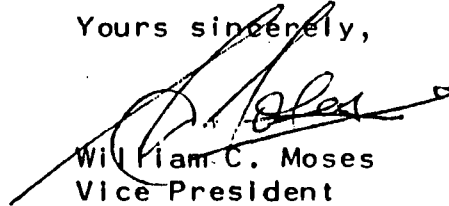
We will be pleased to quote also for the supply of suitable hoods for the spreading machines and a collection system when you so require.

David M. Company

4 August 1983

We trust that the enclosed information is sufficient for your immediate requirements but we will be very pleased to discuss any aspects further with you at your convenience.

Yours sincerely,



William C. Moses
Vice President

WCM/pmd



COMPANY

Wheelabrator-Frye Inc.

Chemicals & Coatings Group

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

10/13 Bill Willard

October 10, 1983

Mr. C / H. Fancy, P.E.
Deputy Chief
Bureau of Air Quality Management
Twin Towers Office Building
9000 Blair Stone Road
Tallahassee, FL 32301-3241

DER
OCT 13 1983
BAQM

Dear Mr. Fancy:

David 'M' Company is in receipt of your correspondence requesting additional information concerning its recent application to construct a lithographic printing blanket manufacturing facility. We will move as expeditiously as possible to gather the necessary information and will consult Mr. Willard Hanks of your staff concerning any questions we may have during the course of compiling the information you have requested.

Sincerely,

Michael A. Ware

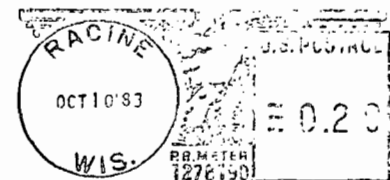
Michael A. Ware
Graphics Div; Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

MAW:bjw

cc: Wayne Brady
R. H. Baddeley
J. Spangler
H. Mycroft
T. Lucas
J. Seabury
C. Collins



201 VALENTINE WAY
LONGWOOD, FLORIDA 32750



Mr. C. H. Fancy, P.E.
Deputy Chief
Bureau of Air Quality Management
Twin Towers Office Building
9000 Blair Stone Road
Tallahassee, FL 32301-3241



DUST COLLECTOR

DUCT WORK -

TALC APPLICATOR

DAVID M
ORLANDO

SCALE: 1"=1'-0" 9/1/83 RHE

DWG # 2 OF 3 REV 9-2-83

REV 10/1/83

BLAST
GATE
(2)
PER'D

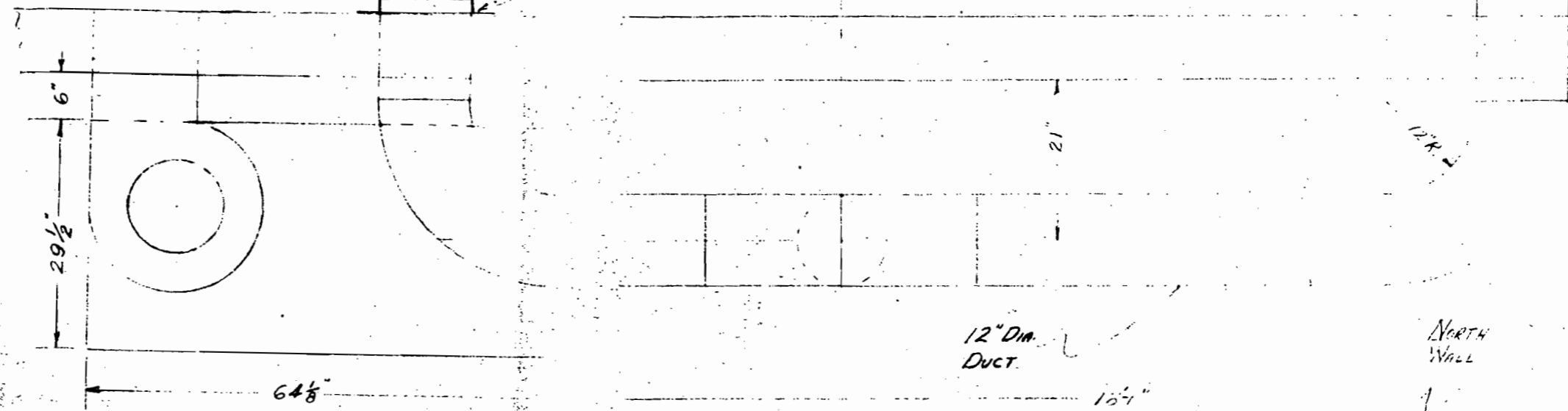
15'-2"

31

WALL FLANGE (4) REQ'D

4'-3"

10'-6"

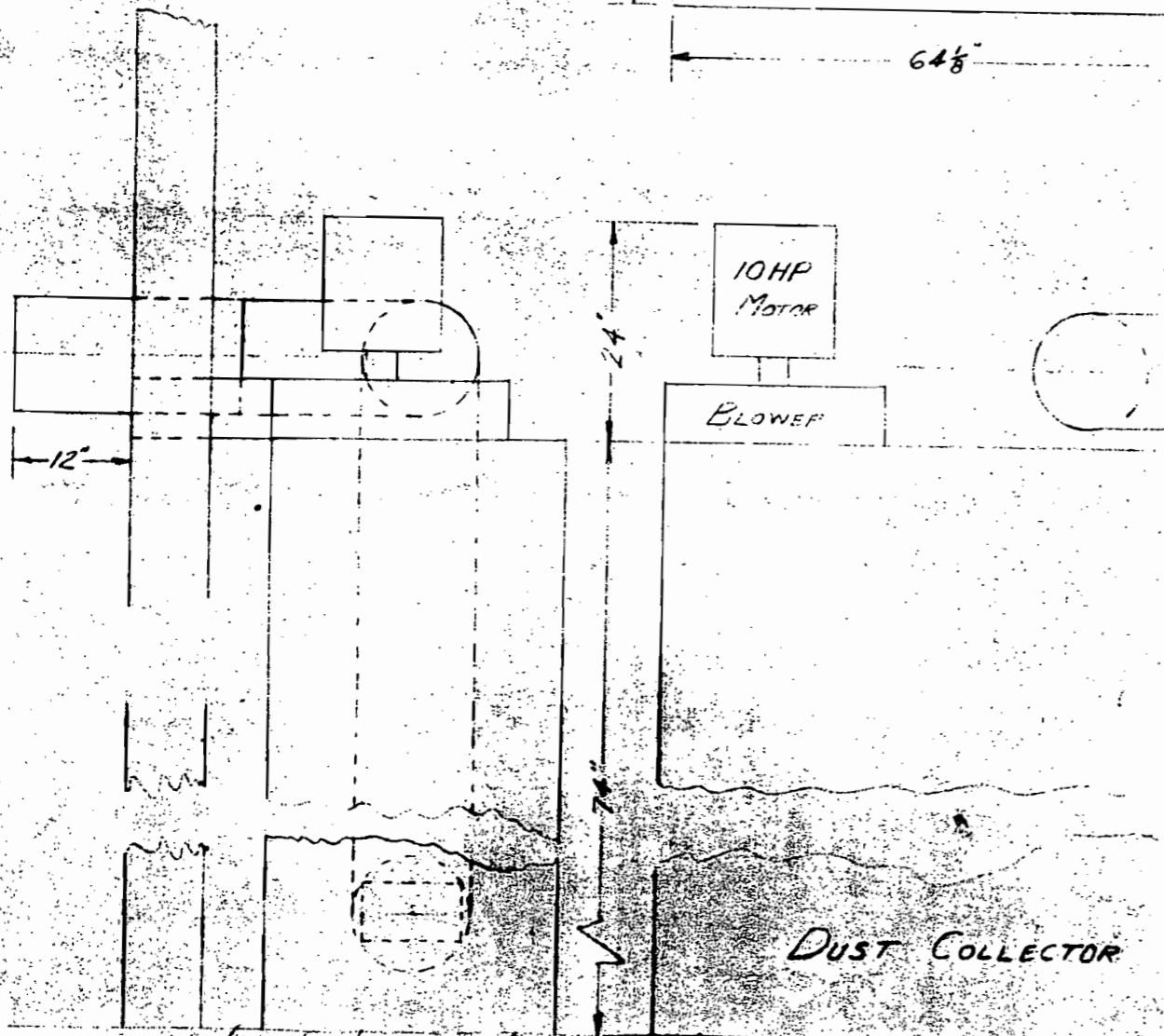


12" DIA.
DUCT

NORTH
WALL

10'-4"

6'-4 1/8"



10HP
MOTOR

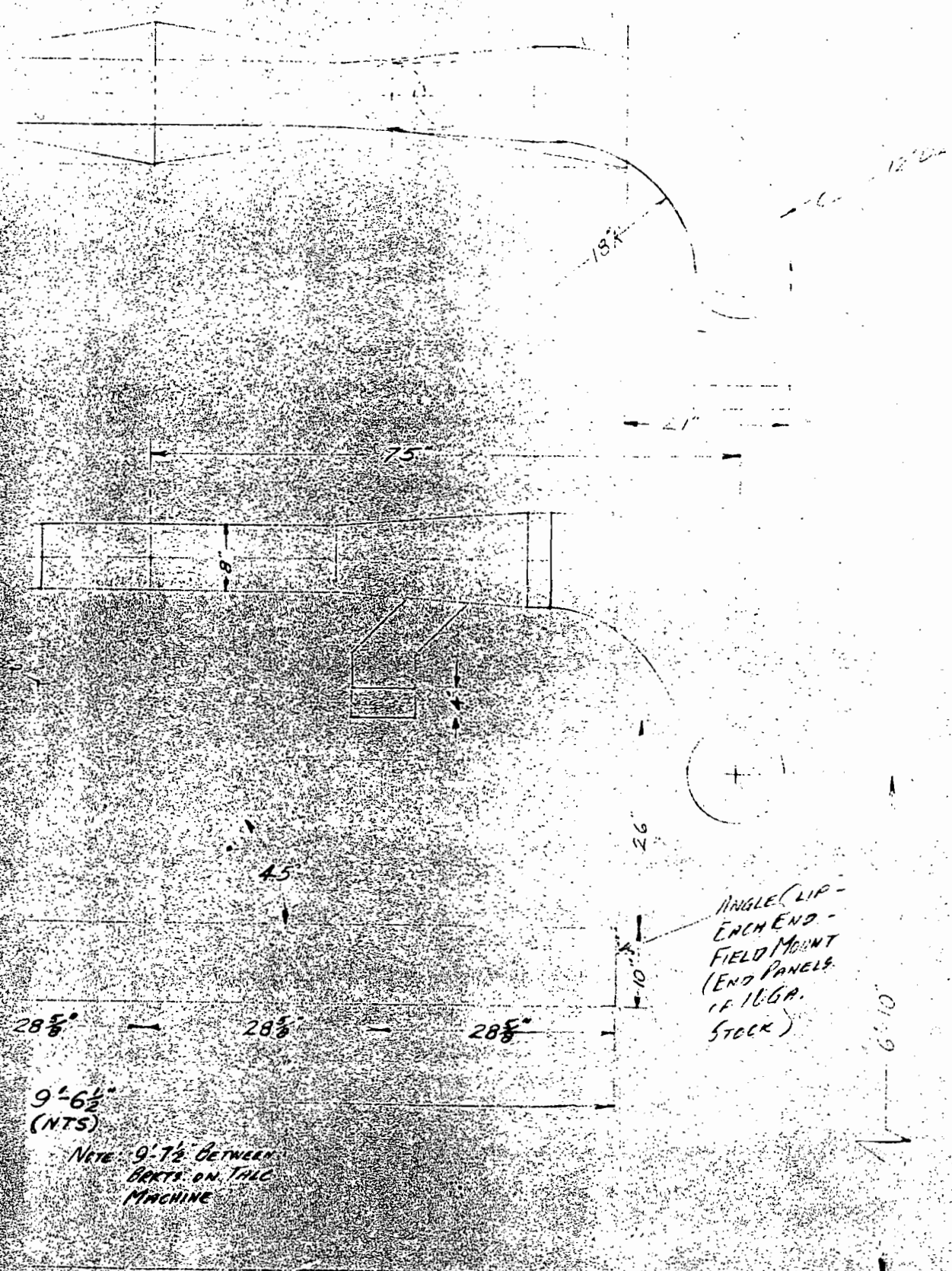
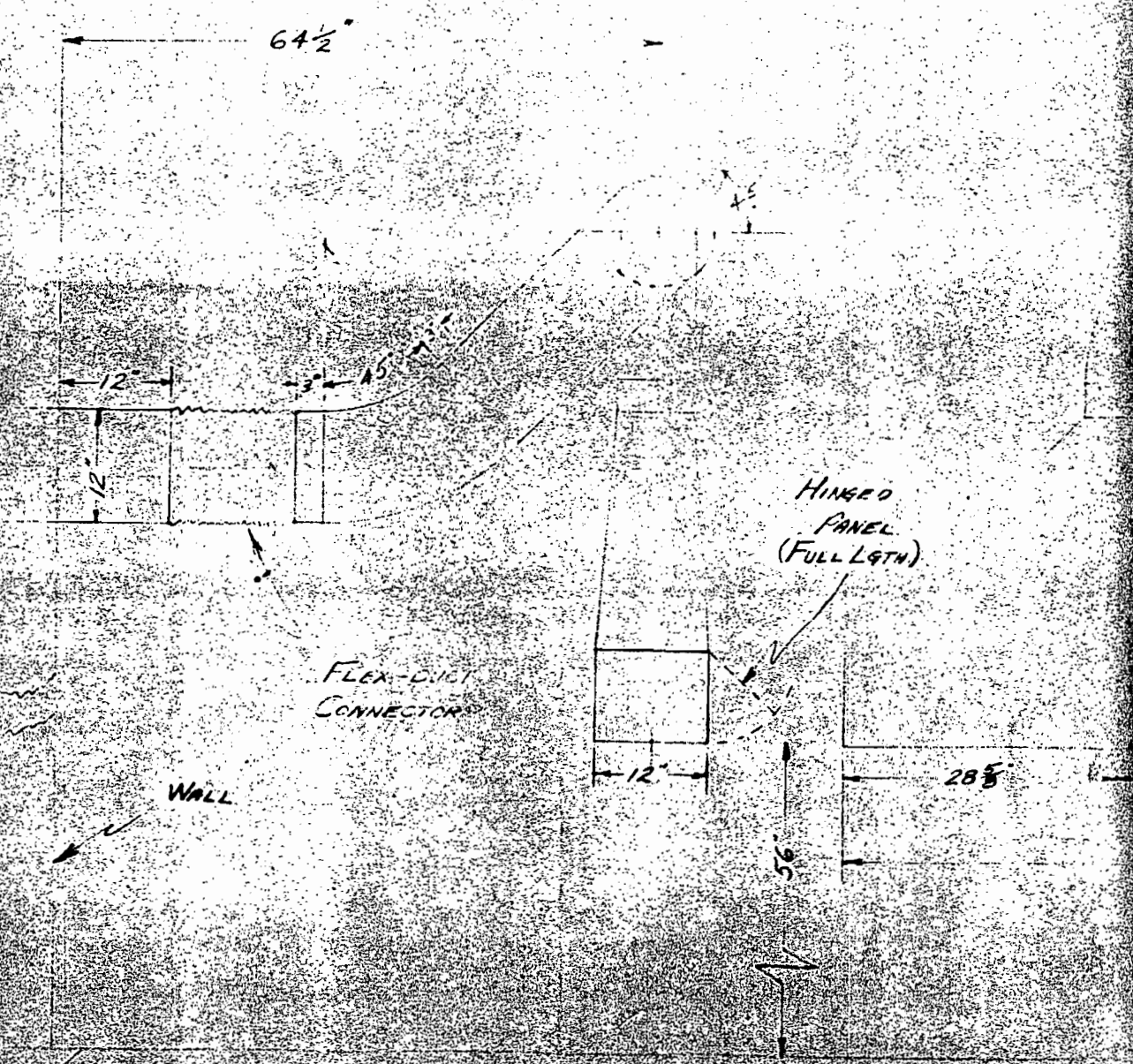
BLOWER

DUST COLLECTOR

DUST COLLECTOR
HOOD & DUCT -
TALC APPLICATOR -
 DAVID M
 ORLANDO

SCALE: 1"=1'-0"
 DWG. # 1 OF 3

9/19/83 RHB
 REV. 3/22/83
 REV. 10/1/83



9'-6 1/2"
 (NTS)
 NOTE: 9'-7 1/2" BETWEEN
 PARTS ON TALC
 MACHINE

ANGLE (LIP -
 EACH END -
 FIELD MOUNT
 (END PANELS.
 IF 16 GA.
 STOCK))



Wheelabrator-Frye Inc. Chemicals & Coatings Group

201 VALENTINE WAY • LONGWOOD, FLORIDA 32750 • (305) 321-0945

October 10, 1983

Mr. L. T. Kozlov, P.E.
District Enforcement Chief
State of Florida
Department of Environmental Regulation
St. Johns River District
3319 Maguire Blvd.
Suite 232
Orlando, FL 32803

DER
OCT 12 1983
BAQM

SUBJECT: Your Memorandum of August 12, 1982 Concerning VOC Permits,
Boiler Permits and Settlement Fees

Dear Mr. Kozlov:

Following our original meeting with Mr. Chuck Collins of your department in late February of 1983, the David 'M' Company undertook a project to quantify the VOC content of its many rubber cements used in the blanket making process. The purpose of the study was to assess our ability to comply with existing low solvent technology guidelines for the fabric coating industry which were referred to by Mr. Collins during the course of our February meeting. The results of our study which encompassed ten of our rubber cements representing high, median and low VOC formulations, yielded solvent contents in coatings ranging from a low of 3.6 lbs. solvent/gallon of coating applied to 6.1 lbs. solvent/gallon of coating applied. The afore-mentioned results were obtained in early May and indicated that David 'M' Company could not meet the 2.9 lb. VOC/gallon of coating applied limitation established by Florida Administrative Code Rule 17-2.650(1)(f)(4) with our existing process parameters.

We subsequently initiated a project to assess the feasibility of lowering our existing VOC leadings in order to achieve compliance with the previously referenced emission limitation. In July of 1983, our study was completed and we concluded that although a few of our compounds could be reformulated to meet the existing guidelines, the vast majority of them could not be formulated into compliance without inducing density and viscosity problems which would preclude their use on our coating lines.

While conducting the previous survey, we concurrently evaluated the possibility of utilizing waterborne or 100% solids coatings. The results of this survey indicated that both of these approaches would require extensive research and development over a long period of time and

promised no guaranty that compliance would be achieved while maintaining the ability to produce a viable and marketable product.

Having exhausted the alternatives previously discussed, David 'M' Company began evaluating the cost and feasibility of installing three types of end of pipe control devices designed to yield compliance with F.A.C. Rule 17-2.650(1)(f)(4) by removing 90% of the VOC delivered to the control device. The technologies preliminarily investigated were thermal oxidation, catalytic oxidation and carbon adsorption.

At the time of this writing, we have received representative quotations from suppliers of the above technologies for the purpose of cost comparison. Our engineering, environmental and process personnel are currently engaged in the establishment of the specifications and operational parameters required for the installation of such control devices.

We have also completed and forwarded to your office, permit and applications covering the existing VOC operations at the suggestion of both yourself and Mr. Chuck Collins of your staff during the course of our July 29, 1983 meeting concerning this subject.

I sincerely hope that I have described to your satisfaction our efforts to comply with regulations of your department. In the event that you should require further information and/or explanation of our efforts, either past or present, please don't hesitate to contact me directly.

Sincerely,



Michael A. Ware
Graphics Div. Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

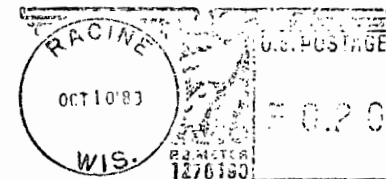
MAW:bjw

cc: Wayne Brady
Charles Collins
Willard Hanks
Herb Mycroft
Jerry Spangler



PRINTING DEVELOPMENTS, INC.

2010 INDIANA STREET
RACINE, WIS. 53405



Mr. Willard Hanks
State of Florida
Department of Environmental Regulation
Twin Towers Office Building
9600 Blair Stone Road
Tallahassee, FL 32301

P 408 530 302

RECEIPT FOR CERTIFIED MAIL

NO INSURANCE COVERAGE PROVIDED—
NOT FOR INTERNATIONAL MAIL

(See Reverse)

Sent to <i>Wayne Brady</i>	
Street and No. <i>201 Valentine Way</i>	
P.O., State and ZIP Code <i>Longwood, FL 32750</i>	
Postage	\$
Certified Fee	
Special Delivery Fee	
Restricted Delivery Fee	
Return Receipt Showing to whom and Date Delivered	
Return Receipt Showing to whom, Date, and Address of Delivery	
TOTAL Postage and Fees	\$
Postmark or Date	

PS Form 3800, Feb. 1982

PS Form 3811, Jan. 1979

RETURN RECEIPT, REGISTERED, INSURED AND CERTIFIED MAIL

SENDER: Complete items 1, 2, and 3. Add your address in the "RETURN TO" space on reverse.

1. The following service is requested (check one.)
 Show to whom and date delivered. \$
 Show to whom, date and address of delivery. \$
 RESTRICTED DELIVERY
 Show to whom and date delivered. \$
 RESTRICTED DELIVERY.
 Show to whom, date, and address of delivery. \$

(CONSULT POSTMASTER FOR FEES)

2. ARTICLE ADDRESSED TO:
Wayne Brady
201 Valentine Way
Longwood, FL 32750

3. ARTICLE DESCRIPTION:
 REGISTERED NO. CERTIFIED NO. INSURED NO.
P408530302

(Always obtain signature of addressee or agent)

I have received the article described above.
 SIGNATURE Addressee Authorized agent
Wayne Brady

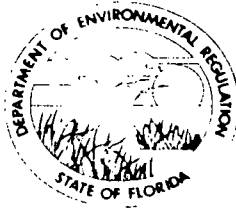
4. DATE OF DELIVERY POSTMARK
9-6-83

5. ADDRESS (Complete only if requested)

6. UNABLE TO DELIVER BECAUSE: CLERK'S INITIALS
CS

DEPARTMENT OF ENVIRONMENTAL REGULATION

TWIN TOWERS OFFICE BUILDING
2600 BLAIR STONE ROAD
TALLAHASSEE, FLORIDA 32301-8241



BOB GRAHAM
GOVERNOR

VICTORIA J. TSCHINKEL
SECRETARY

September 2, 1983

Mr. Wayne L. Brady, General Manager
David "M" Company
201 Valentine Way
Longwood, Florida 32750

Dear Mr. Brady:

The Department has made a preliminary review of your application for a permit to construct a lithographic printing blanket manufacturing facility. The additional information listed below is required before we can process your application.

1. What will be the maximum production (weight/time) and operation time (hr/d) of the plant? This information is required to establish the maximum allowable production of the plant and will be a condition in any permit to construct that is issued for the plant.
2. What percent of the maximum production was the emission and raw material data in the application based on?
3. Will solvents other than MEK and toluene be used at this plant?
4. How many gallons per year of each solvent will be consumed at the maximum production of the plant?
5. How will the solvents be shipped to the plant? Submit a drawing showing the storage area and piping to the mixing area.
6. How many storage tanks will be built at this plant? What air pollution control equipment will be used to minimize VOC emission from the tanks? What will the emission from the tanks be? Please submit a copy of the emission calculations.
7. What will be the mixing ratio of solvent to rubber (lb/lb)?
8. How many pounds of volatile organic compounds (VOC) will be in a gallon of applied coating?

Mr. Wayne L. Brady
Page Two
September 2, 1983

9. How will talc be received, stored, and conveyed to the process? What air pollution control equipment or operation practices will be used to minimize talc emissions? How much talc will be emitted to the atmosphere?
10. Will the solvent laden air discharged from the plant contain any particulate matter or oil mist?
11. What will be the emissions (particulate matter and VOC) from air emission point No. 2 (mills)? What control equipment will be used to limit particulate emission from the mills? Will product scrap be recycled to the mills? If so, how much VOC will be emitted by the scrap?
12. Are the flow rates listed in the application from air emissions points 4 and 5 correct? A flow of 1415 ACFM through two 1.33 ft x 1.33 ft ducts would have a velocity of 400 FPM.
13. What will be the percent of the lower explosive limit (LEL) at peak solvent evaporation rates for each exhaust stream?
14. What will be the percent of the threshold limit values (TLV) near the discharge points (approx. 20') outside the plant during peak solvent evaporation rates?
15. How many employees will work at this plant and what impact will the emissions from the plant have on the ambient air quality, vegetation and visibility near the plant?
16. If the solvents evaporated from the process were recovered, could they be reused by the plant or sold to a reclamation plant?
17. What is the current delivered cost of the solvents used at the plant?
18. Has the Company made any studies on recovering and reusing the solvents at the plant and, if so, what conclusions were reached?
19. On page 10 of 12 of the application, the capital cost of carbon absorbers is shown as \$600,000 - \$700,000.

Mr. Wayne L. Brady
Page Three
September 2, 1983

Reference sources available to us estimate the cost of a carbon absorber is \$300,000. This cost includes process and equipment design, foundations, erection, electrical wiring, controls and thermal insulation. Please explain what is included in your estimate.

20. How many gallons of VOC per year will be sent to Emille, Alabama for disposal?

If the information requested confirms the proposed plant is a new major source of VOC, (annual emissions greater 250 TPY), you will also need a federal permit to construct the plant. To apply for a federal permit, you need only to request the Department to process the application for both federal and state permits.

Emission standards for major VOC sources in attainment areas are established by the best available control technology (BACT) procedures described in Florida Administrative Code Rule 17-2.500. Although I cannot predict what standard will be allowed for your process, regulations require the emission standard to be no more than 2.9 lb VOC/gal of coating applied or 90 percent removal of VOC from the exhaust stream (Fla. Admin. Code Rule 17-2.650(1)(f)4). If the plant cannot meet this standard with low solvent content coating technology, then it will be necessary for you to propose the control equipment that the plant will use to meet the standard.

Please furnish the information requested by November 1, 1983, so that we can resume processing your application. If you have any questions on the data requested, Willard Hanks (904/488-1344) will discuss them with you.

Sincerely,



C. H. Fancy, P.E.
Deputy Chief
Bureau of Air Quality
Management

CHF/WH/s

cc: Charles Collins
John Seabury

DER PERMIT APPLICATION TRACKING SYSTEM MASTER RECORD

FILE#000000073966 COE# DER PROCESSOR:CLAIR FANCY DER OFFICE:TLH
FILE NAME:DAVID "M" COMPANY DATE FIRST REC: 08/12/83 APPLICATION TYPE:AC
APPL NAME:DAVID M/SPREADERS #1,2,3 APPL PHONE:(305)324-0945 PROJECT COUNTY:59
ADDR:204 VALENTINE WAY CITY:LONGWOOD ST:FLZIP:
AGNT NAME:SEABURY-BOTTORF ASSOC AGNT PHONE:(305)298-0846
ADDR:3702 SILVER STAR ROAD CITY:ORLANDO ST:FLZIP:32808

ADDITIONAL INFO REQ: / / / / / / REC: / / / / / /
APPL COMPLETE DATE: / / COMMENTS NEC:Y DATE REQ: / / DATE REC: / /
LETTER OF INTENT NEC:Y DATE WHEN INTENT ISSUED: / / WAIVER DATE: / /

HEARING REQUEST DATES: / / / / / /
HEARING WITHDRAWN/DENIED/ORDER -- DATES: / / / / / /
HEARING ORDER OR FINAL ACTION DUE DATE: / / MANUAL TRACKING DESIRED:N
*** RECORD HAS BEEN SUCCESSFULLY UPDATED *** 08/22/83 14:29:50
FEE PD DATE#1:08/12/83 \$1000 RECEIPT#00074416 REFUND DATE: / / REFUND \$
FEE PD DATE#2: / / \$ RECEIPT# REFUND DATE: / / REFUND \$
APPL:ACTIVE/INACTIVE/DENIED/WITHDRAWN/TRANSFERRED/EXEMPT/ISSUED:TR DATE:08/12/83
REMARKS:ORL. CHECK ALREADY SENT TO PURCHASING OFFICE.

DER
AUG 24 1983
BAQM



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~000020~~ 240189

COPY

YOUR REF:

OUR REF: E 0379
S 7212

DATE: 2 August 1983

P R O P O S A L

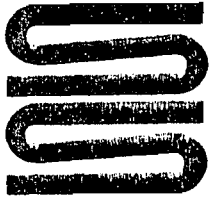
F O R

S O L V E N T R E C O V E R Y S Y S T E M

TO BE LOCATED AT:

DAVID M. COMPANY

201 VALENTINE WAY
LONGWOOD, FLORIDA



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~900020~~ 240189

YOUR REF:

OUR REF: S 7212

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

SOLVENT RECOVERY PLANT

ACTIVATED CARBON ADSORPTION UNIT

AIRFLOW.....25,000 scfm.

AIR TEMPERATURE.....140° F.

SOLVENT CAPACITY.....50 gph (360 lb/hr)

SOLVENT.....Toluene

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL

598 $\frac{176}{42}$ $\frac{2000\#}{7}$ $\frac{7}{16 \times 250h}$ = 299 #/h
 300 #/h
 (304 #/h
 2000 #/h)

David M. Company

Proposal No. S 7212

PLANT DESCRIPTION

The solvent laden air would be filtered to remove dust, cooled to approximately 95° F by the air cooler and then delivered by the fan to the three adsorbers.

Two of the three adsorbers would normally be handling the solvent laden air whilst the other would be regenerating with steam to extract the solvent from the carbon or would be held on "stand-by" following "steaming" and a short dry/cool period.

The adsorbers would cycle consecutively and automatically with override of the cycle time and steam input by gas analyser control. The proposed adsorbers are of annular bed design, the solvent air after filtering and cooling passes through an automatic inlet valve into the outer annulus of the adsorber, through the carbon bed and then exhausts to atmosphere through the inner annulus and the automatic exhaust valve.

At the end of an adsorbing period the air inlet and outlet valves close and an automatic steam valve allows steam to enter the inner annulus of the adsorber and pass through the carbon bed to the outer annulus from which point the steam together with solvent vapour extracted from the carbon passes to the condenser. During steaming some condensation occurs on the outer shell of the adsorber and is drained through filters to join the main condensate flow from the condenser to the decanter.

At the end of the steaming period the adsorber automatic steam valve shuts and the air inlet and outlet valves open to allow solvent laden air to again pass through the adsorber for a short period to dry and cool the bed (during this dry/cool period the solvent laden air is shut off from the other adsorber). The inlet valve of the dried/cooled adsorber would then close and the adsorber would be held on "stand-by" until the gas analyser, sampling the exhaust of the other adsorber, or the override timer, gives the signal for that adsorber to change to the steaming cycle, the stand-by adsorber being first brought back into service, so that there is no interruption to the adsorber cycle.

Any incondensibles from the condenser return through the condenser vent pipe to the inlet duct for recycling through the adsorbers. The condenser vent is fitted with high temperature alarm to indicate overloading from excessive steam flow or shortage or failure of the cooling water supply.

Safety features include adsorber pressure and vacuum relief valves and liquid seals.

David M. Company

Proposal No. S 7212

We would supply the following:

1. AIR FILTER - 25,000 scfm capacity

a) Pre-Filter Section

With removable and disposable fiberglass panels to protect and extend the life of the main filter.

b) Main Filter Section

With disposable filter elements of the HEPA type of fiberglass material, with an efficiency of better than 99% for particles down to 0.3 microns per standard tests.

The whole of the above would be incorporated in a galvanized steel case with access doors to facilitate servicing.

A differential resistance gauge would be fitted.

2. AIR COOLER - 25,000 scfm capacity.

Would be of finned tube design with copper tubes and aluminum fins to cool the inlet airstream from 140° F to approximately 95° F when supplied with cooling water at a temperature of maximum 86° F.

The air cooler coils would be housed in a galvanized steel case and would be drainable.

3. FAN AND MOTOR

The fan would be of the high efficiency type with airfoil impeller capable of handling 25,000 scfm of solvent laden air, when at an actual temperature of approximately 95° F, at a total pressure of approximately 18" w.g., allowing for 2" w.g. suction before the air filters.

The casing would be of substantial carbon steel construction and the unit would have spark-proof features to AMCA standards.

The impeller would be driven through a flexible coupling by a 125 HP explosion proof motor to Class 1, Group D, Division 1, 460 volt, 3 phase, 60 cycles.

A manually operated radial leaf damper would be fitted at the fan inlet for regulation of the airflow.

David M. Company

Proposal No. S 7212

4. THREE ADSORBERS - 6'8" diameter x approximately 10'3" high shell.

Each capable of handling 12,500 scfm of solvent laden air calculated at 70° F, when at an actual temperature of approximately 95° F.

The adsorbers would be of the annular carbon bed type with the carbon contained between stainless steel panels formed up into cyclinders and reinforced by carbon steel stiffening bands. The activated carbon bed and screens would be supported upon a carbon steel tray to which would be connected the solvent laden air inlet valve, vapour piping to condenser and liquor drains. The air exhaust valve and the steam valve would be connected to the outlet branch of the adsorber.

The carbon bed and screens would be contained within a removable carbon steel shell, fitted with access and inspection holes with bolted cover plates. Flanged connections would be fitted to the tray so that the carbon could be easily and quickly run off into drums.

Vacuum and pressure relief valves would be provided for each adsorber.

Handrails would be fitted round the tops of the adsorbers, with a walkway over and between the adsorbers, and with an access ladder from ground level, in accordance with the requirements of OSHA.

5. ACTIVATED CARBON

We would provide the initial charge of approximately 6400 lbs of high quality carbon for each of the three adsorbers. The carbon would be manufactured by Sutcliffe Speakman and would be of the grade most suited to the required duty.

6. INLET AND OUTLET AIR VALVES - 26" diameter

We would provide an inlet and outlet valve for each adsorber 26" diameter of the mushroom type of substantial construction arranged for pneumatic operation through the automatic control mechanism.

Renewable seats would be fitted in the valve bodies and access holes with bolted cover plates provided. An indicator would be fitted to each valve so that the attitude of the valve could be easily ascertained.

David M. Company

Proposal No. S 7212

7. CONDENSER

To condense the steam and solvent vapours leaving an adsorber during the steaming period.

The unit would be of the tubular surface type with the shell in 304 stainless steel and the tubes in 316 stainless steel. The water boxes and tubeplates would be in carbon steel.

The condenser vent pipe (fitted with thermometer with high temperature alarm contacts) would be connected to the inlet duct so that any incondensibles displaced from the adsorbers would recycle.

8. ONE DECANTER - 3'0" diameter x 4'6" high

It would be of the self-adjusting decanting type to separate automatically the condenser water and solvent. The tank would be of carbon steel welded construction fitted with a bolted cover plate and all the necessary connections. A sight glass would be fitted in the solvent outlet.

9. TRUNKING

We would provide the interconnecting trunking between the air filter, air cooler, fan and adsorbers, but do not allow for any trunking prior to the inlet manifold of the air filter. The trunking would be of carbon steel of welded construction flanged at suitable intervals and be adequately supported.

10. THREE EXHAUSTS

An exhaust stack would be provided for each adsorber, of carbon steel welded construction, galvanized and would terminate about 6' above the adsorber platform.

11. PIPING AND VALVES

We would provide pipework to interconnect the above and manifold the service pipes to terminal flanges at the battery limits.

Vapour and liquor piping would be in 304 stainless steel, utility piping in carbon steel.

We include for all isolating valves, steam traps, pipe supports, etc.

David M. company

Proposal No. S 7212

12. PLANT CONTROL UNIT

Would consist of the following:

a) Gas Analyser (Infra-Red Type)

To measure the solvent content in the exhaust airstream of the adsorber next due for desorbing and to delay steaming until the set point was achieved. However, an adjustable override timer would be incorporated into the program to limit the period of such a delay.

A strip recorder for the exhaust gas concentration would be included and the unit would have a "high level" alarm.

b) Programmable Logic Controller

Of the solid state type which on receipt of the appropriate signals would cause the adsorber valves to sequence as appropriate. The program logic would be such that the next step would not be commenced until the previous one had been proven completed.

In the unlikely event of failure of the P.L.C. the solvent recovery plant could be operated manually by the use of pilot air solenoid valves inbuilt into the panel.

c) Annunciation Section

Of the 'Panalarm' type or similar to alarm:
"high level" - gas analyser output
"high temperature" - condenser vent

d) Control Panel

The above items would be mounted in a free-standing enclosure, suitable for indoor location in a non-hazardous area, assumed within 50' of the adsorbers.

The panel would be pre-wired and tubed at terminal points on the frame. We exclude the impulse tubing from the panel to the plant items.

David M. Company

Proposal No. S 7212

13. INSTRUMENTS

We would provide the following indicating type instruments which would be locally mounted on the plant:

- 4 Dial type thermometers
- 1 Steam pressure gauge
- 1 Differential pressure gauge (resistance through air filter)
- 1 4-Point pressure gauge (air pressure and suction within the plant)

14. STRUCTURES

We would provide structures to support items of the plant.

15. PAINTING

Where applicable, plant items would be primer coated before shipment.

16. COMPRESSED AIR

You to supply compressed air, clean, dry and oil free at a pressure of approximately 80 psig.

17. DRAWINGS AND INSTRUCTIONS

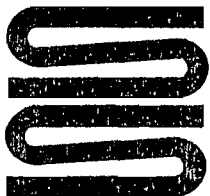
We would provide three (3) copies of each of the following:

- Foundation drawing
- Flowsheet
- Plant arrangement drawing
- Major assemblies

Plus two (2) copies of:

- Erection drawings and instructions
- Operating and Maintenance Manual

A list of recommended spares would also be provided.



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: 908020 240189

YOUR REF:

OUR REF:

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

TERMS, PRICE AND CONDITIONS

PRICE OF PROPOSAL

PROPOSAL NO. S 7212

For the supply of the whole of the items as specified in this proposal, delivered only to your works in Longwood, Florida (offloading from the transport and installation by others) would be the sum of approximately.....\$ 370,000.00

(THREE HUNDRED SEVENTY THOUSAND DOLLARS)

Any Federal, State or local taxes that may be applicable are excluded.

TERMS OF PAYMENT

25% of quoted price with order.

45% of quoted price in three (3) equal payments pursuant to invoices on the following basis:

- a) upon submission of preliminary drawings comprising plant arrangement and foundation drawings and flowsheet.
- b) upon submission of confirmation that the manufacture is at least 40% complete.
- c) upon submission of confirmation that the manufacture is at least 70% complete.

25% of quoted price upon despatch or when ready for despatch if despatch date is delayed by you.

5% of quoted price on start-up and successful operation, at latest three (3) months after despatch, or when ready for despatch, whichever is sooner.

Continued.....

David M. Company

Proposal No. S 7212

TERMS, PRICE AND CONDITIONS...continued

DESPATCH

Approximately six (6) months from receipt of order assuming that there are no delays due to changes in specifications or approval of drawings.

EXCLUSIONS

We do not include for any motor controls, electrical wiring, or any special cold weather protection or insulation, or any parts, materials, work for services of any kind whatsoever except those specifically mentioned in the body of the proposal.

STANDARD OF MANUFACTURE

The goods described in this proposal would be manufactured to normal commercial standards and be suitable for the duty specified. Should you desire manufacture to any specific standards, codes or inspection schedules other than those allowed for then we will be pleased to comply with your requests, but any extra charges so incurred would be in addition to the prices quoted.

VALIDITY OF PROPOSAL

This proposal is open for acceptance for a period of thirty (30) days from the date of this quotation and after this period confirmation must be obtained that the terms and conditions contained herein are still valid.

SUTCLIFFE SPEAKMAN INCORPORATED



William C. Moses
Vice President

E. & O. E.

WCM/pmd

David M. Company

Proposal No. S 7212

UTILITY REQUIREMENTS

The following are the approximate utility requirements of the plant when operating at design load conditions. The figures are subject to confirmation by our final design study.

STEAM.....1190 lb/hr, dry saturated quality
at 40 psig.

COOLING WATER.....170 gpm, inlet 86^o, maximum,
return 115^o.

POWER.....100 Kwh.

COMPRESSED AIR.....20 scfh.

SPACE

The equipment, we estimate, would require a floor area of approximately 45' x 20'.

The height of the equipment would be approximately 18', but if it is to be located inside, then provision should be made for lifting of the adsorber shells.

APPENDIX 'A'

OUR PRELIMINARY PROPOSAL FOR THE TURNKEY INSTALLATION

OF THE PLANT QUOTED IN MAIN PROPOSAL NO. S 7212

1. FOUNDATIONS

We would provide concrete foundations for the appropriate items of our equipment.

Our price is conditional on the understanding that the ground is of suitable load bearing capacity, that no rock blasting or piling would be required and that all work would be above the water table.

2. MOTOR CONTROL CENTER

We include the starters for the fan motor, the water cooling tower fan and the recirculating water pumps, and which would be suitable for location in a non-hazardous area which we assume will be by the control panel.

3. WATER COOLING TOWER

unknown Inhibitor S? Chemicals

We would supply and install a water cooling tower designed to cool the required amount of water from 115° F to approximately 86° F. The tower would be of induced draft type and complete with fan and motor. The system would include two pumps with motors (one normally operating, one stand-by) and the water circulating piping and valves between the water cooling tower and the air cooler and condenser of the recovery plant, the water cooling tower to be at ground level and located adjacent to the recovery plant.

4. INSTALLATION

We include for the necessary labour, tools, moving and lifting gear including cranes, for the installation of the items of equipment specified in this proposal onto the foundation.

4. INSTALLATION...Continued

It is understood that the work would be undertaken during our normal working hours and that we and our sub-contractors or agents would be allowed free and unrestricted access to the site.

Any overtime worked at your request would be chargeable extra at the rate in force at the time.

You would be responsible for offloading from the transport any items delivered prior to the start of the installation work and provide suitable storage and protection for all parts close to the battery limits.

All temporary site support facilities including but not limited to water, electricity, steam, drain, office and telephone would be provided by you at your expense, adjacent to the site.

Note that the quoted price assumes we will have continuity of work on site.

5. SUPERVISION

We include for the services of our own suitably qualified supervisor to supervise and give instructions to the labour provided by our sub-contractors, to secure the installation of the equipment in a competent manner.

6. IMPULSE TUBING

The pneumatic tubing from the appropriate plant items to the control panel would be supplied and installed in a workmanlike manner. It is assumed that the tubing run is a maximum of fifty feet.

7. ELECTRICAL WIRING

We would supply and install all necessary electrical wiring, cable trays, etc. to interconnect the various items of plant to the motor control center and the control panel, assuming a maximum cable run of fifty feet from the adsorbers. Where appropriate, the wiring would be to explosion proof, classification.

David M. Company

Proposal No. S 7212

8. INSULATION/WINTERIZATION

Insulation and heat tracing of appropriate equipment within our battery limits effected, assuming a minimum ambient temperature of 25° F.

9. START-UP

We include for the services of a skilled engineer for a period of two (2) consecutive weeks on site to start-up the plant and provide instructions to your personnel on its operation.

Any extra time required by you or non-continuity in attendance would be at additional cost.

10. INSURANCE

We would provide and maintain the following insurances and would require any and all sub-contractors employed by us at your site to procure and maintain the same type and amount of insurance.

- a) Workman's Compensation and employer's Liability
- \$100,000.00 each accident and aggregate disease.
- b) Comprehensive General Liability
- Bodily injury and Property damage of up to \$1,000,000 combined single limit.
- c) Comprehensive Automobile Liability
- To cover owned, non-owned, leased and hired cars, combined single limit \$1,000,000.

Certificates of the above shall be provided to you prior to the start of our on site work.

In addition to the above and in order to insure the equipment from its delivery to your site, through installation to your acceptance of the plant, we would take out "all risks" insurance plus "builders risk" and "installation floaters" at the appropriate time and at your expense.

David M. Company

Proposal No. S 7212

BUDGETARY PRICE

Our approximate additional price to that quoted in Main Proposal No. S 7212 for the turnkey installation of the plant as detailed in this Appendix 'A' would be the sum of\$ 160,000.00.

(ONE HUNDRED SIXTY THOUSAND DOLLARS)

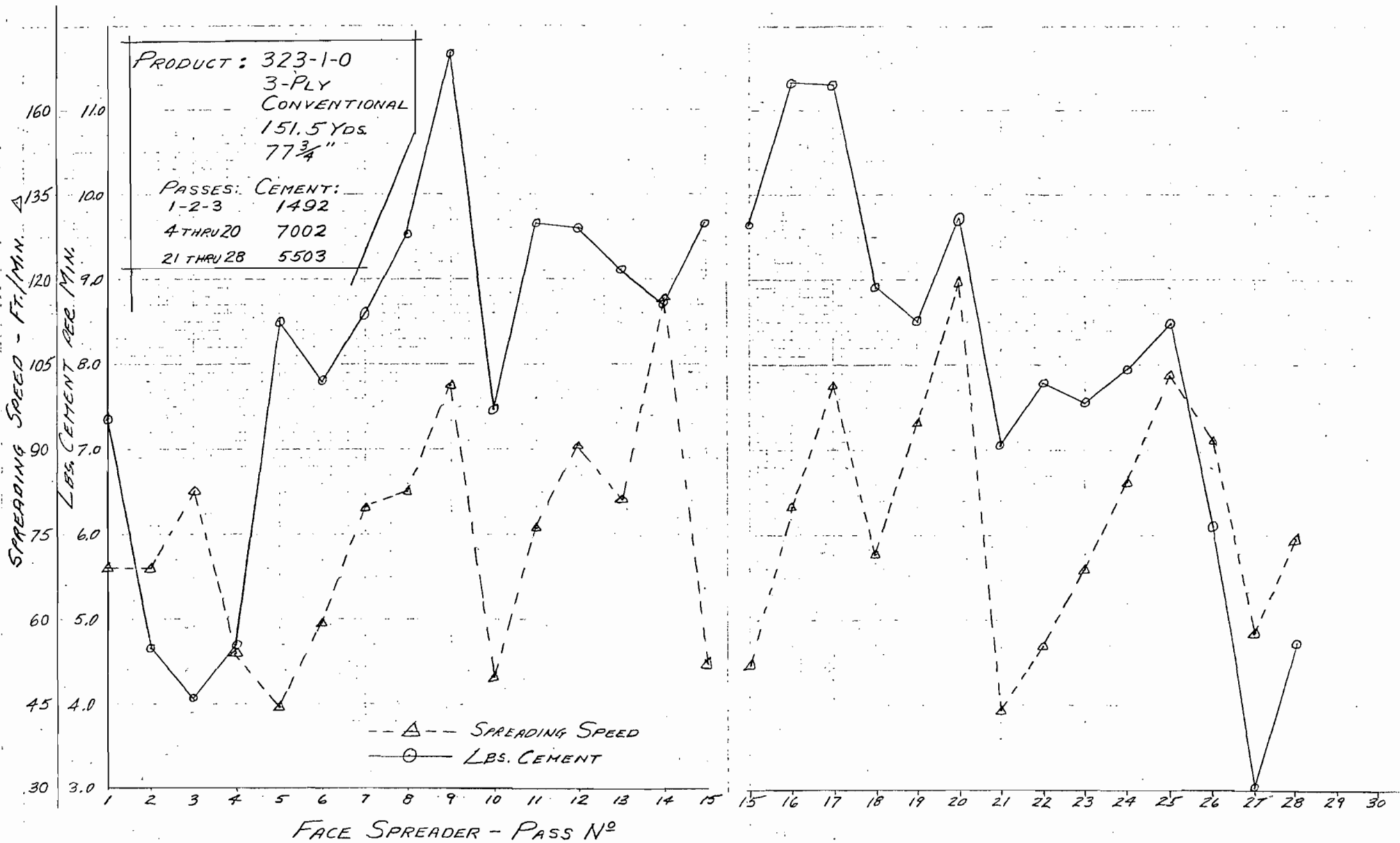
COMPLETION

We would expect to supply and install the specified plant in approximately nine (9) months from receipt of order.

TERMS AND CONDITIONS

Otherwise as Main Proposal No. S 7212.

A handwritten signature in black ink, appearing to be 'D. M. Company', is located in the lower right quadrant of the page.



CHARTED ^{9/1/63} 5/19/63 RHB

AC 59-073966
\$1,000 fee paid

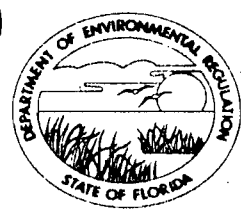
STATE OF FLORIDA

DEPARTMENT OF ENVIRONMENTAL REGULATION

ST. JOHNS RIVER DISTRICT

3319 MAGUIRE BOULEVARD
SUITE 232
ORLANDO, FLORIDA 32803

P A I D
AUG 12 1983
SAINT JOHNS RIVER DISTRICT



BOB GRAHAM
GOVERNOR
VICTORIA J. TSCHINKEL
SECRETARY
ALEX SENKEVICH
DISTRICT MANAGER

APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCES

DER

SOURCE TYPE: Blanket Spreading Operation New Existing **AUG 24 1983**

APPLICATION TYPE: Construction Operation Modification

COMPANY NAME: DAVID "M" COMPANY COUNTY: SEMINOLE **BAOM**

Identify the specific emission point source(s) addressed in this application (i.e. Lime Spreaders #1, #2, #3)
Kiln No. 4 with Venturi Scrubber; Peaking Unit No. 2, Gas Fired) Pestooning and Mixing area

SOURCE LOCATION: Street 201 Valentine Way City Longwood

UTM: East 17-470106 North 3177160

Latitude 28° 43' 26" N Longitude 81° 18' 18" W

APPLICANT NAME AND TITLE: Wayne L. Brady, General Manager

APPLICANT ADDRESS: 201 Valentine Way, Longwood, Florida 32750

SECTION I: STATEMENTS BY APPLICANT AND ENGINEER

A. APPLICANT

I am the undersigned owner or authorized representative* of David "M" Company

I certify that the statements made in this application for a construction permit are true, correct and complete to the best of my knowledge and belief. Further, I agree to maintain and operate the pollution control source and pollution control facilities in such a manner as to comply with the provision of Chapter 403, Florida Statutes, and all the rules and regulations of the department and revisions thereof. I also understand that a permit, if granted by the department, will be non-transferable and I will promptly notify the department upon sale or legal transfer of the permitted establishment.

*Attach letter of authorization

Signed: [Signature]
Wayne L. Brady, General Manager
Name and Title (Please Type)

Date: 8/ /83 Telephone No. 305/321-0945

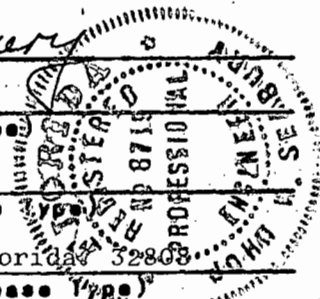
B. PROFESSIONAL ENGINEER REGISTERED IN FLORIDA (where required by Chapter 471, F.S.)

This is to certify that the engineering features of this pollution control project have been designed/examined by me and found to be in conformity with modern engineering principles applicable to the treatment and disposal of pollutants characterized in the permit application. There is reasonable assurance, in my professional judgment, that

¹ See Florida Administrative Code Rule 17-2.100(57) and (104)

the pollution control facilities, when properly maintained and operated, will discharge an effluent that complies with all applicable statutes of the State of Florida and the rules and regulations of the department. It is also agreed that the undersigned will furnish, if authorized by the owner, the applicant a set of instructions for the proper maintenance and operation of the pollution control facilities and, if applicable, pollution sources.

Signed John W. Seabury
John W. Seabury
Name (Please Type)
Seabury-Bottorf Associates, Inc.
Company Name (Please Type)
3702 Silver Star Rd., Orlando, Florida 32808
Mailing Address (Please Type)



Florida Registration No. 8719 Date: August 11, 1983 Telephone No. 305/298-0846

SECTION II: GENERAL PROJECT INFORMATION

A. Describe the nature and extent of the project. Refer to pollution control equipment, and expected improvements in source performance as a result of installation. State whether the project will result in full compliance. Attach additional sheet if necessary.

The project will involve the installation of a lithographic printing blanket manufacturing facility and will include the following sources which are part of a common exhaust system. Spreaders #1, #2, #3, Festooning area, and Mixing area. Please refer to emission points 1-5 on the following process layout diagrams to ascertain particular emission points and exhaust ventilation configurations.

B. Schedule of project covered in this application (Construction Permit Application Only)

Start of Construction upon arrival Completion of Construction 1 month post approval

C. Costs of pollution control system(s): (Note: Show breakdown of estimated costs only for individual components/units of the project serving pollution control purposes. Information on actual costs shall be furnished with the application for operation permit.)

Not applicable

D. Indicate any previous DER permits, orders and notices associated with the emission point, including permit issuance and expiration dates.

None known

E. Requested permitted equipment operating time: hrs/day 16 ; days/wk 5 ; wks/yr 50 ;
if power plant, hrs/yr N.A. ; if seasonal, describe: _____

F. If this is a new source or major modification, answer the following questions.
(Yes or No)

1. Is this source in a non-attainment area for a particular pollutant? No
a. If yes, has "offset" been applied? No
b. If yes, has "Lowest Achievable Emission Rate" been applied? No
c. If yes, list non-attainment pollutants. No

2. Does best available control technology (BACT) apply to this source?
If yes, see Section VI. Yes

3. Does the State "Prevention of Significant Deterioration" (PSD)
requirement apply to this source? If yes, see Sections VI and VII. No

4. Do "Standards of Performance for New Stationary Sources" (NSPS)
apply to this source? No

5. Do "National Emission Standards for Hazardous Air Pollutants"
(NESHAP) apply to this source? No

H. Do "Reasonably Available Control Technology" (RACT) requirements apply
to this source? No

a. If yes, for what pollutants? N.A.

b. If yes, in addition to the information required in this form,
any information requested in Rule 17-2.650 must be submitted.

Attach all supportive information related to any answer of "Yes". Attach any justifi-
cation for any answer of "No" that might be considered questionable.

F. 2. (Yes) Florida Air Pollution Regulation 17-2.500(5)(c) Application of best
available control technology to pollutants subject to NSR requirements as set forth
in 17.2.5(2)(f).

SECTION III: AIR POLLUTION SOURCES & CONTROL DEVICES (Other than Incinerators)

A. Raw Materials and Chemicals Used in your Process, if applicable:

Description	Contaminants		Utilization Rate - lbs/hr	Relate to Flow Diagram
	Type	% Wt		
Rubber	None Known	-	Variable	Used to formulate cements 2 Mixing (2), Spreading (3, 4, 5), Festooning (6) (At step #6 99%+ of V.O.C. has been flashed off)
Rubber Cement	V.O.C.	30-70%	Variable	
	MEK			
	Toluene			

*The process utilizes rubber cements which are mixtures of various rubber compounds i.e. (nitrile, etc) and hydrocarbon diluents such as toluene or MEK. VOC content of cement may vary from 30 to 70% of

B. Process Rate, if applicable: (See Section V, Item 1) coating weight depending upon type.

- Total Process Input Rate (lbs/hr): Variable
- Product Weight (lbs/hr): Variable

C. Airborne Contaminants Emitted: (Information in this table must be submitted for each emission point, use additional sheets as necessary)

Name of Contaminant	Emission ¹		Allowed ² Emission Rate per Rule 17-2	Allowable ³ Emission lbs/hr	Potential ⁴ Emission ^{16 hrs}		Relate to Flow Diagram
	Maximum lbs/hr	Actual T/yr			lbs/yr	T/yr	
VOC - MEK/	103	84	BACT	To be determined	420,000	210	1
" Toluene	5	4	BACT	To be determined	20,000	10	3
"	94.5	75.6	BACT	To be determined	378,000	189	4
"	94.5	75.6	BACT	To be determined	378,000	189	5
	<u>297.0</u>				<u>major source</u> 598		

* Potential emissions are based upon a two shift (16 hour) operation, five days per week and 250 days per year. Actual emissions are based upon a one shift (8 hour) operation, five days per week, and 250 days per year.

¹ See Section V, Item 2.

² Reference applicable emission standards and units (e.g. Rule 17-2.600(5)(b)2. Table II, E. (1) - 0.1 pounds per million Btu heat input)

³ Calculated from operating rate and applicable standard.

⁴ Emission, if source operated without control (See Section V, Item 3).

2 ?
Missing

↑
these must be listed
11/1/83
300 TPY
VOC emissions
(uncontrolled)

D. Control Devices: (See Section V, Item 4)

Name and Type (Model & Serial No.)	Contaminant	Efficiency	Range of Particles Size Collected (in microns) (if applicable)	Basis for Efficiency (Section V Item 3)

E. Fuels

Type (Be Specific)	Consumption*		Maximum Heat Input (MMBTU/hr)
	avg/hr	max./hr	

*Units: Natural Gas--MMCF/hr; Fuel Oils--gallons/hr; Coal, wood, refuse, other--lbs/hr.

Fuel Analysis:

Percent Sulfur: N.A. Percent Ash: N.A.

Density: N.A. lbs/gal Typical Percent Nitrogen: N.A.

Heat Capacity: N.A. BTU/lb N.A. BTU/gal

Other Fuel Contaminants (which may cause air pollution): N.A.

F. If applicable, indicate the percent of fuel used for space heating.

Annual Average N.A. Maximum N.A.

G. Indicate liquid or solid wastes generated and method of disposal.

Rubber cement waste is generated by the process and consists of rubber and varying amounts of rubber mixed with VOC. The waste is packaged in 55 gallon open head drums and disposed of in a secure hazardous waste land disposal site in Emille, Alabama.

H. Emission Stack Geometry and Flow Characteristics (Provide data for each stack); (See following page)
 Stack Height: _____ ft. Stack Diameter: _____ ft.
 Gas Flow Rate: _____ ACFM _____ DSCFM Gas Exit Temperature: _____ °F.
 Water Vapor Content: _____ % Velocity: _____ FPS

SECTION IV: INCINERATOR INFORMATION

Not Applicable

Type of Waste	Type 0 (Plastics)	Type I (Rubbish)	Type II (Refuse)	Type III (Garbage)	Type IV (Pathological)	Type V (Liq. & Gas By-prod.)	Type VI (Solid By-prod.)
Actual lb/hr Incinerated							
Uncontrolled (lb/hr)							

Description of Waste: _____
 Total Weight Incinerated (lb/hr) _____ Design Capacity (lb/hr) _____
 Approximate Number of Hours of Operation per day _____ day/wk _____ wks/yr. _____
 Manufacturer: _____
 Date Constructed: _____ Model No. _____

	Volume (ft) ³	Heat Release (BTU/hr)	Fuel		Temperature (°F)
			Type	BTU/hr	
Primary Chamber					
Secondary Chamber					

Stack Height: _____ ft. Stack Diameter: _____ Stack Temp. _____
 Gas Flow Rate: _____ ACFM _____ DSCFM* Velocity: _____ FPS

*If 50 or more tons per day design capacity, submit the emissions rate in grains per standard cubic foot dry gas corrected to 50% excess air.

Type of pollution control devices: Cyclone Wet Scrubber Afterburner
 Other (specify) _____

H. Emission stack geometry and flow characteristics: From estimates

Emission Point 1 (SEE process floor diagram):

Stack Height: 18 ft. Stack Dimensions: 4.5 x 4.5 square (ft.)
Gas Flow Rate: 12,600 ACFM 9960 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content Ambient (variable)% Velocity: 600 FPS 622.2 FPM

20.25 FT

Emission Point 3 (See process flow diagram):

Stack Height: 12 ft. Stack Dimensions: 1.5 x 1.5 square (ft.)
Gas Flow Rate: 1575 ACFM 1410 DSCFM Gas Exit Temp.: 95 (°F)
Water Vapor Content Ambient (variable)% Velocity: 700 FPS 700 FPM

Emission Point 4 (See process flow diagram):

Stack Height: 12.5 ft. Stack Dimensions: 2(1.33 x 1.33 outlets)(ft.)
Gas Flow Rate: 1415 ACFM 1120 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content: Ambient (variable)% Velocity: 800 FPS ? 400 FPS

Emission Point 5 (See process flow diagram):

Stack Height: 12.5 ft. Stack Dimensions: 2(1.33 x 1.33 outlets)(ft.)
Gas Flow Rate: 1415 ACFM 1120 DSCFM Gas Exit Temp.: 150 (°F)
Water Vapor Content: Ambient (variable)% Velocity: 800 FPS ?

$$\text{Velocity} = \frac{12,600 \text{ FT}^3}{\text{MIN } 20.25 \text{ FT}^2} = 622.2 \text{ FPM}$$

Discharge point 2 (Calls) missing

Brief description of operating characteristics of control devices: Not Applicable

Ultimate disposal of any effluent other than that emitted from the stack (scrubber water, ash, etc.):

Not Applicable

NOTE: Items 2, 3, 4, 6, 7, 8, and 10 in Section V must be included where applicable.

SECTION V: SUPPLEMENTAL REQUIREMENTS

Please provide the following supplements where required for this application.

1. Total process input rate and product weight -- show derivation [Rule 17-2.100(127)]
NOT APPLICABLE
2. To a construction application, attach basis of emission estimate (e.g., design calculations, design drawings, pertinent manufacturer's test data, etc.) and attach proposed methods (e.g., FR Part 60 Methods 1, 2, 3, 4, 5) to show proof of compliance with applicable standards. To an operation application, attach test results or methods used to show proof of compliance. Information provided when applying for an operation permit from a construction permit shall be indicative of the time at which the test was made.
SEE APPENDIX 1 ATTACHED
3. Attach basis of potential discharge (e.g., emission factor, that is, AP42 test).
SEE APPENDIX 1 ATTACHED
4. With construction permit application, include design details for all air pollution control systems (e.g., for baghouse include cloth to air ratio; for scrubber include cross-section sketch, design pressure drop, etc.)
NOT APPLICABLE
5. With construction permit application, attach derivation of control device(s) efficiency. Include test or design data. Items 2, 3 and 5 should be consistent: actual emissions = potential (1-efficiency).
NOT APPLICABLE
6. An 8 1/2" x 11" flow diagram which will, without revealing trade secrets, identify the individual operations and/or processes. Indicate where raw materials enter, where solid and liquid waste exit, where gaseous emissions and/or airborne particles are evolved and where finished products are obtained.
SEE APPENDIX 2
7. An 8 1/2" x 11" plot plan showing the location of the establishment, and points of airborne emissions, in relation to the surrounding area, residences and other permanent structures and roadways (Example: Copy of relevant portion of USGS topographic map).
SEE APPENDIX 4
8. An 8 1/2" x 11" plot plan of facility showing the location of manufacturing processes and outlets for airborne emissions. Relate all flows to the flow diagram.
SEE APPENDIX 3

NO
Control
Device

9. The appropriate application fee in accordance with Rule 17-4.05. The check should be made payable to the Department of Environmental Regulation. ATTACHED
10. With an application for operation permit, attach a Certificate of Completion of Construction indicating that the source was constructed as shown in the construction permit. NOT APPLICABLE

SECTION VI: BEST AVAILABLE CONTROL TECHNOLOGY

A. Are standards of performance for new stationary sources pursuant to 40 C.F.R. Part 60 applicable to the source?

Yes No

Contaminant	Rate or Concentration
N.A.	N.A.

B. Has EPA declared the best available control technology for this class of sources (If yes, attach copy)

Yes No

Contaminant	Rate or Concentration
BACT has not been declared for this class of sources.	

C. What emission levels do you propose as best available control technology?

Contaminant	Rate or Concentration
VOLATILE ORGANIC COMPOUNDS	AVERAGE COATING VOC CONTENT OF 5.4 lbs./gal.
	VOC/gal. OF COATING DELIVERED TO COATING APPLICATOR.

D. Describe the existing control and treatment technology (if any).

- | | |
|---------------------------|--------------------------|
| 1. Control Device/System: | 2. Operating Principles: |
| 3. Efficiency:* | 4. Capital Costs: |

*Explain method of determining

5. Useful Life:

6. Operating Costs:

7. Energy:

8. Maintenance Costs:

9. Emissions:

Contaminant	Rate or Concentration

10. Stack Parameters

- a. Height: ft.
- b. Diameter: ft.
- c. Flow Rate: ACFM
- d. Temperature: °F.
- e. Velocity: FPS

E. Describe the control and treatment technology available (As many types as applicable, use additional pages if necessary).

1. LOW SOLVENT TECHNOLOGY

- a. Control Device: FORMULATE COATING TO GIVEN SPECIFICATION
- b. Operating Principles: CONTROL VOC EMISSION BY FORMULATION
- c. Efficiency:¹ NOT APPLICABLE
- d. Capital Cost: ESTIMATE, \$10,000.00
- e. Useful Life: CONTINUING
- f. Operating Cost: \$20,000.00
- g. Energy:² NOT APPLICABLE
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE
- j. Applicability to manufacturing processes: APPLICABLE
- k. Ability to construct with control device, install in available space, and operate within proposed levels: ABLE TO INITIATE

2. CATALYTIC INCINERATION

- a. Control Device: CATALYTIC INCINERATOR
- b. Operating Principles: CATALYZED INCINERATION
- c. Efficiency:¹ 90% +
- d. Capital Cost: \$500,000.00
- e. Useful Life: 10-15 YEARS
- f. Operating Cost: NOT AVAILABLE AT THIS WRITING
- g. Energy:² NOT AVAILABLE FROM BUDGET
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE

¹ Explain method of determining efficiency.

² Energy to be reported in units of electrical power - KWH design rate.

- j. Applicability to manufacturing processes: APPLICABLE WITH SOME PROCESS MODIFICATIONS
- k. Ability to construct with control device, install in available space, and operate within proposed levels: NO FORSEEABLE DIFFICULTIES

3. SOLVENT RECOVERY

- a. Control Device: CARBON ADSORBER
- b. Operating Principles: PHYSICAL ADSORPTION
- c. Efficiency:¹ 85-90%
- d. Capital Cost: \$600,000.00-\$700,000.00
- e. Useful Life: 10-15 YEARS
- f. Operating Cost: NOT AVAILABLE AT THIS WRITING
- g. Energy:² NOT AVAILABLE FROM BUDGET QUOTATION
- h. Maintenance Cost: NOT AVAILABLE
- i. Availability of construction materials and process chemicals: AVAILABLE
- j. Applicability to manufacturing processes: APPLICABLE WITH PROCESS MODIFICATIONS
- k. Ability to construct with control device, install in available space, and operate within proposed levels: NO FORSEEABLE MAJOR OBSTACLES TO INSTALLATION

4.

- a. Control Device:
- b. Operating Principles:
- c. Efficiency:¹
- d. Capital Costs:
- e. Useful Life:
- f. Operating Cost:
- g. Energy:²
- h. Maintenance Cost:
- i. Availability of construction materials and process chemicals:
- j. Applicability to manufacturing processes:
- k. Ability to construct with control device, install in available space, and operate within proposed levels:

F. Describe the control technology selected: LOW SOLVENT TECHNOLOGY

- 1. Control Device: FORMULATE TO GIVEN SPECIFICATION
- 2. Efficiency:¹ NOT APPLICABLE
- 3. Capital Cost: \$10,000.00
- 4. Useful Life: CONTINUING
- 5. Operating Cost: \$20,000.00
- 6. Energy:² NOT APPLICABLE
- 7. Maintenance Cost: NOT APPLICABLE
- 8. Manufacturer: NOT APPLICABLE
- 9. Other locations where employed on similar processes: NONE KNOWN

a. (1) Company: NOT APPLICABLE

(2) Mailing Address:

(3) City:

(4) State:

¹ Explain method of determining efficiency.

² Energy to be reported in units of electrical power - KWH design rate.

(5) Environmental Manager:

(6) Telephone No.:

(7) Emissions:¹

Contaminant

Rate or Concentration

(8) Process Rate:¹

b. (1) Company:

(2) Mailing Address:

(3) City:

(4) State:

(5) Environmental Manager:

(6) Telephone No.:

(7) Emissions:¹

Contaminant

Rate or Concentration

(8) Process Rate:¹

10. Reason for selection and description of systems:

¹Applicant must provide this information when available. Should this information not be available, applicant must state the reason(s) why.

SECTION VII - PREVENTION OF SIGNIFICANT DETERIORATION

NOT APPLICABLE

A. Company Monitored Data

1. _____ no. sites _____ TSP _____ () SO₂ _____ Wind spd/dir

Period of Monitoring _____ / _____ / _____ to _____ / _____ / _____
month day year month day year

Other data recorded _____

Attach all data or statistical summaries to this application.

*Specify bubbler (B) or continuous (C).

2. Instrumentation, Field and Laboratory

- a. Was instrumentation EPA referenced or its equivalent? [] Yes [] No
- b. Was instrumentation calibrated in accordance with Department procedures?
[] Yes [] No [] Unknown

B. Meteorological Data Used for Air Quality Modeling

- 1. _____ Year(s) of data from _____ / _____ / _____ to _____ / _____ / _____
month day year month day year
- 2. Surface data obtained from (location) _____
- 3. Upper air (mixing height) data obtained from (location) _____
- 4. Stability wind rose (STAR) data obtained from (location) _____

C. Computer Models Used

- 1. _____ Modified? If yes, attach description.
- 2. _____ Modified? If yes, attach description.
- 3. _____ Modified? If yes, attach description.
- 4. _____ Modified? If yes, attach description.

Attach copies of all final model runs showing input data, receptor locations, and principle output tables.

D. Applicant's Maximum Allowable Emission Data

Pollutant	Emission Rate
TSP	_____ grams/sec
SO ²	_____ grams/sec

E. Emission Data Used in Modeling

Attach list of emission sources. Emission data required is source name, description of point source (on NEDS point number), UTM coordinates, stack data, allowable emissions, and normal operating time.

F. Attach all other information supportive to the PSD review.

G. Discuss the social and economic impact of the selected technology versus other applicable technologies (i.e., jobs, payroll, production, taxes, energy, etc.). Include assessment of the environmental impact of the sources.

H. Attach scientific, engineering, and technical material, reports, publications, journals, and other competent relevant information describing the theory and application of the requested best available control technology.

Maximum Potential Emissions

Spreader #1

Maximum VOC loading in lbs./min. during spreading: 4.3 lbs./min. = 258 lbs./hr

Actual spreading time: 15% of operating time (from time study)

Potential operating parameters: 16 hours/day
250 days/year
5 days/week

Use Calculation:

16 hrs./day x .15 = 2.4 hrs./day in spread mode

2.4 hrs./day x $\frac{60 \text{ min.}}{1 \text{ hr.}}$ x 4.3 lbs./min. during spreading = 619.20 lbs./day emission
(maximum potential)

619.20 lbs./day x 250 days/yr. = 154,800 lbs./yr. = **77.4 tons/yr.**

Spreaders #2 and #3

Maximum VOC loading in lbs./min. during spreading: 14 lbs./min. = 840 lbs./hr

Actual spreading time: 15% of operating time (from time study)

Potential operating parameters: 16 hours/day per spreader
250 days/year per spreader
5 days/week per spreader

Use Calculation:

16 hrs./day x .15 = 2.4 hrs./day in spread mode

2.4 hrs./day x $\frac{60 \text{ min.}}{1 \text{ hr.}}$ x 14 lbs./min. = 2016 lbs./day emission-maximum potential

2016 lbs./day x 250 days/yr. = 504,000 lbs./yr. = **252 tons/yr.** per spreader

2 spreaders x 252 tons/year = **504 tons/yr.** for spreaders #2 and #3 - Maximum potential

Festooning Area

The festooning area routinely flashes off one pass worth of solvent per day/blanket. Four blankets would be placed into festooning each day on a two shift operation. Process records indicate that a maximum of 10 lbs. of solvent per blanket enters the festooning area. The blankets leave nearly solvent free. Therefore maximum potential solvent emissions would approximate.

4 blankets/day x 250 days/yr. x 10 lbs. solvent/blanket = 10,000 lbs./yr. = **5 tons/yr.**

Mixing Area

Emissions from the mixing area can be expected to approximate 12 tons/yr. maximum potential by estimate. This would include both local and general room exhausts. 10 T/yr. can be associated with local mixing tank exhaust and 2 T/yr. can be associated with general room exhaust.

Emission Point Allocation

Emission Point 1 :

Emission point 1 captures 25% of the VOC exhaust from spreaders #2 and #3, 100% of the VOC exhaust emissions from spreader #1, 100% of the festooning area VOC exhaust and 100% of general room exhaust from the mixing area.

$$\text{Emission Point 1 Total} = (\text{spreader (\# 1 : 77.4 T/yr.)} + (\text{spreaders (2 and 3) 504 T/yr.} \times .25) + (\text{festooning 5 tons/yr.}) + (\text{mixing general 2.0 T/yr.}) = \boxed{210 \text{ T/yr.}}$$

Emission Point 3 :

Emission Point 3 captures 100% of the VOC exhaust from the mixing room tanks. This emission can be expected to approximate a maximum potential emission of 10 tons/yr.

Emission Point 4 :

Emission point 4 captures 75% of spreader #2's exhaust. Therefore maximum potential emission from emission point 4 would be

$$252 \text{ T/yr.} \times .75 = \boxed{189 \text{ tons/yr.}}$$

Emission Point 5 :

Emission point 5 captures 75% of spreader #3's exhaust. Therefore maximum potential emission from emission point 5 would be

$$252 \text{ T/yr.} \times .75 = \boxed{189 \text{ tons/yr.}}$$

Maximum Lbs./Hr. Calculation

Emission Point 1 :

- a) Spreader #1 maximum spreading time/hr. = 9.0 min./hr.
Spreader #1 maximum solvent use/min. = 4.3 lbs./min.
Spreader #1 emission point collection % = 100%
 $9.0 \text{ min./hr.} \times 4.3 \text{ lbs./min} \times 1.0 = \boxed{38.7 \text{ lbs./hr.}}$
- b) Spreader #2 maximum spreading time/hrs. = 9.0 min./hr.
Spreader #2 maximum solvent use/min. = 14.0 lbs./min.
Spreader #2 emission point collection % = 25%
 $9.0 \text{ min./hr.} \times 14.0 \times .25 = \boxed{31.5 \text{ lbs./hr.}}$
- c) Spreader #3 maximum spreading time/hr. = 9.0 min./hr.
Spreader #3 maximum solvent use/min. = 14.0 lbs./min.
Spreader #3 emission point collection % = 25%
 $9.0 \text{ min./hr.} \times 14.0 \times .25 = \boxed{31.5 \text{ lbs./hr.}}$

d) Mixing general exhaust 2 T/yr. = 4000 lbs./hr. ÷ 250 days/yr. = 16 lbs./day ÷ 16 hrs./day = 1 lb./hr.

TOTAL MAXIMUM LBS./HR. = a + b + c + d
 = 103 lbs./hr.

Emission Point 3 :

Total annual emission = 10 T/yr. = 20,000 lbs./yr.

20,000 lbs./yr. ÷ 250 days/yr. = 80 lbs./day ÷ 16 hrs./day = 5.0 lbs./hr.

Emission Point 4 :

Spreader #2
 Maximum spreading time/hr. = 9.0 min./hr.
 Spreader #3
 Maximum solvent use/min. = 14.0 lbs./min.
 Spreader #2
 Emission point collection % = 75%

9.0 min./hr. x 14.0 lbs./min. x .75 = 94.5 lbs./hr.

Emission Point 5 :

Spreader #3 maximum spreading time/hr. = 9.0 min./hr.
 Spreader #3 solvent use/min. = 14.0 lbs./min.
 Spreader #3 emission point collection % = 75%

9.0 min./hr. x 14.0 lbs./min. x .75 = 94.5 lbs./hr.

Actual Tons/Yr.

The facility currently operates at approximately 40% of its emitting potential. Therefore, potential T/yr. are multiplied by .40 to obtain actual emissions.

<u>Emission Point</u>	<u>Potential Emissions</u>	x	<u>Actual Emissions</u> (.40)	
<u>1</u>	210 T/yr.		84 T/yr.	
<u>3</u>	10 T/yr.		4 T/yr.	
<u>4</u>	189 T/yr.		75.6 T/yr.	
<u>5</u>	189 T/yr.		75.6 T/yr.	TOTAL ACTUAL TONS/YR. = 239.2
	598			

Total VOC/yr. figures correlate well with 1982/83 purchasing records for VOC raw materials. Current 1983 use projections would approximate 240 T/yr.

APPENDIX II CONT.
PROCESS FLOW EXPLANATION

Step ①

Rubber is milled into sheets for use in cement making. No calculation of emissions is included in this application since only particulate emissions would be regulated and this source is a minor one which will be dealt with separately from VOC emissions.

How PM Controlled

Step ②

Rubber is combined with VOC (MEK/Toluene) and mixed to the desired viscosity for spreading.

AP How STORAGED & TRANSPORTED?

Step ③

Rubber cement is spread on cotton fabric in repeated passes of varying thicknesses utilizing a knife type applicator. This step forms the blanket carcass. VOC flashes off due to steam heated coils and is captured by overhead hoods.

How STEAM PRODUCED?

Steps ④ & ⑤

Carcass is finished and moved in roll form to face spreaders 2 and 3 where top or face coats are applied to the blanket, solvent flashes off as in Step 3. The final pass is not dried, rather it is talced and recoiled with interleaving paper at which point it moves to Step 6.

Step ⑥

The blanket is then unrolled and hung open in the festooning area where the final pass dries. The blanket is virtually solvent free after 16 hours of air drying time.

Step ⑦

The blanket is once again rolled up and is put into a vulcanizer which is actually an autoclave which puts the final cure on the blanket.

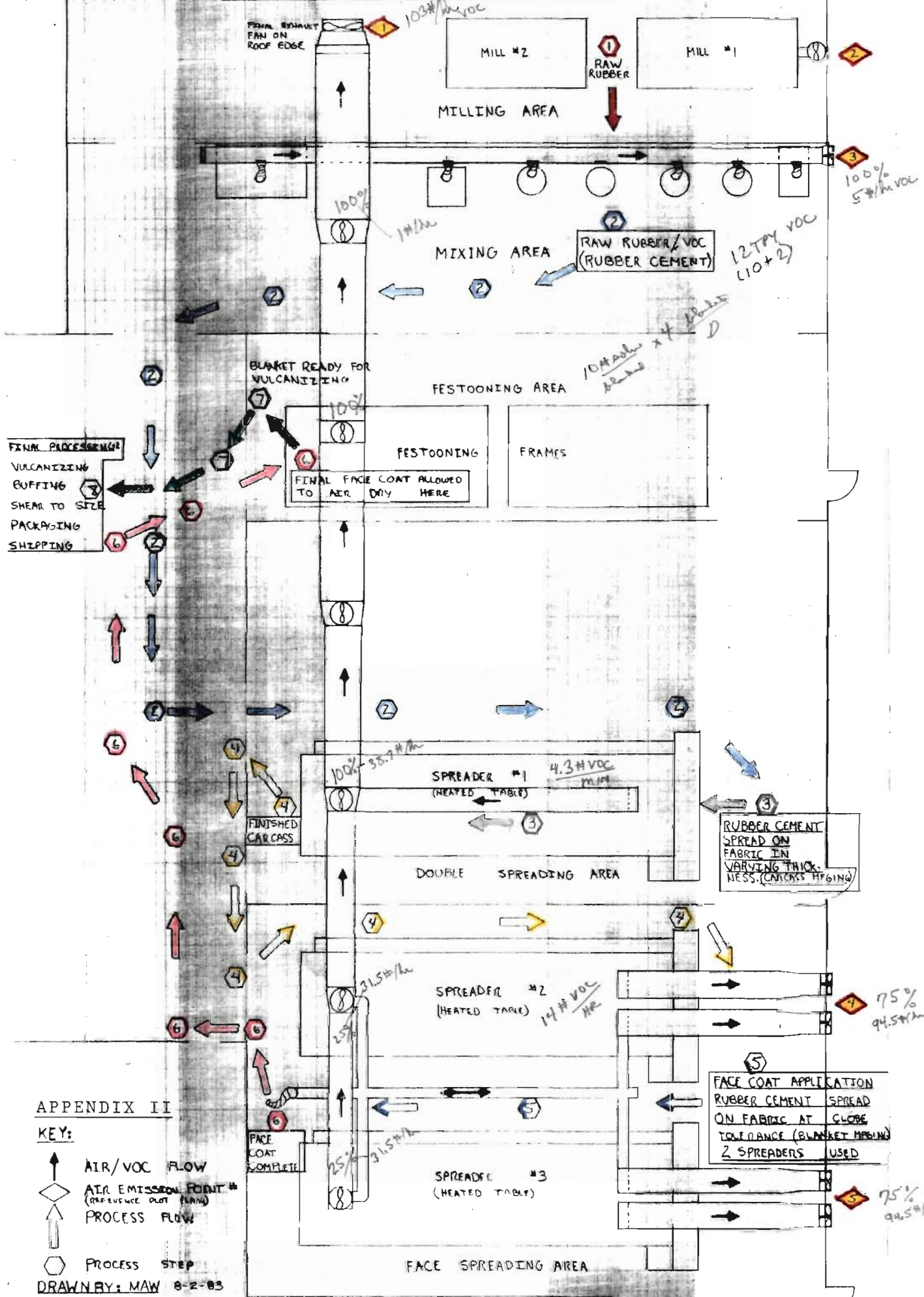
SOURCE OF HEAT

Step ⑧

Following final cure the blanket may be buffed or sheared and then packaged for shipping.

PM Control

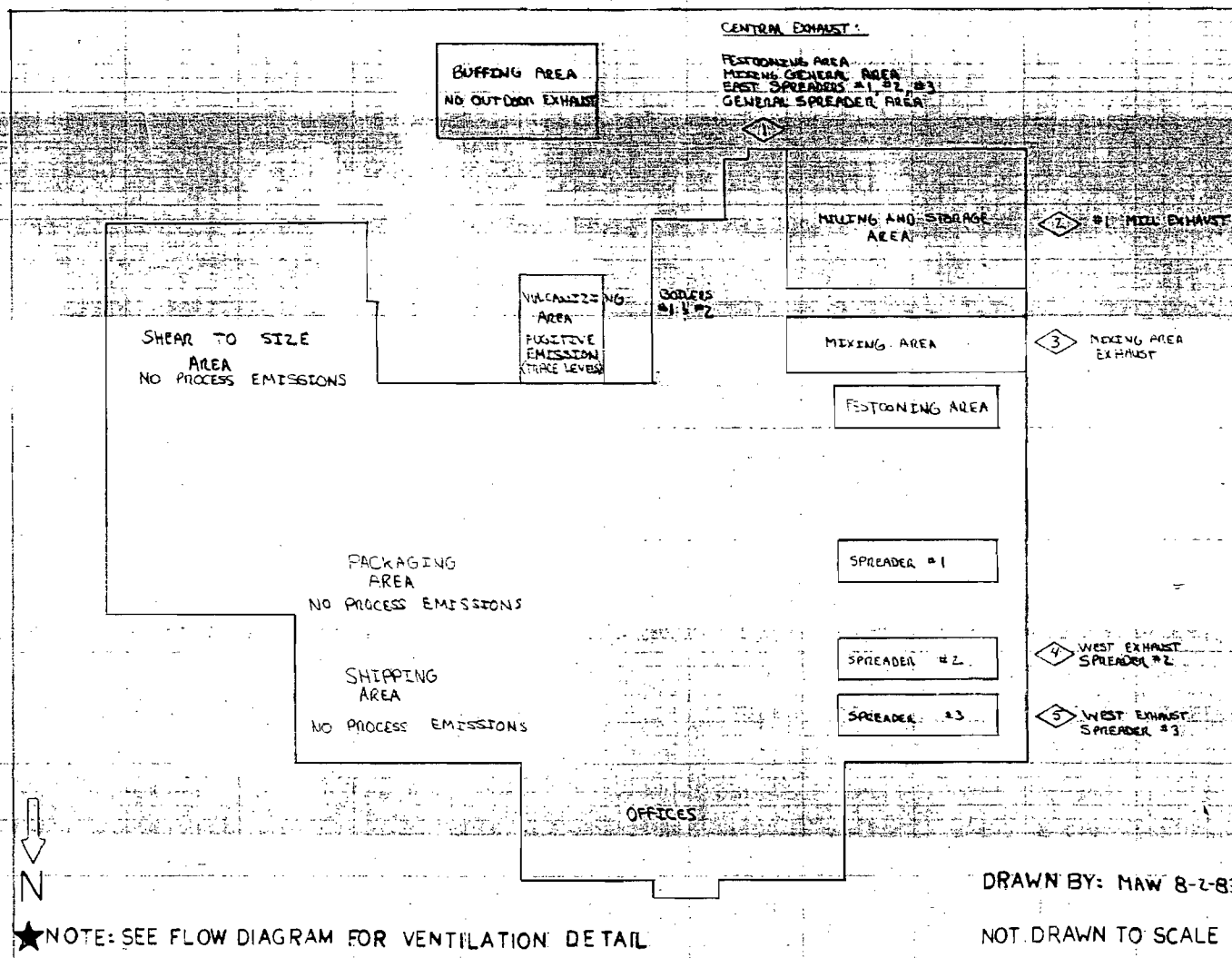
PROCESS LAYOUT AND FLOW DIAGRAM: VOC DATA



Best Available Copy

APPENDIX III

PROCESS LAYOUT AND EMISSION POINTS: \diamond



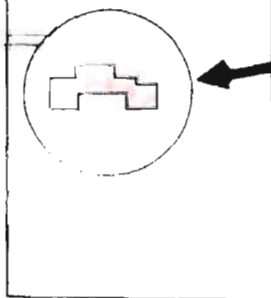
DAVID M COMANY: FACILITY LOCATION

US 17-92

SCS R.R.



MAIN FACILITY



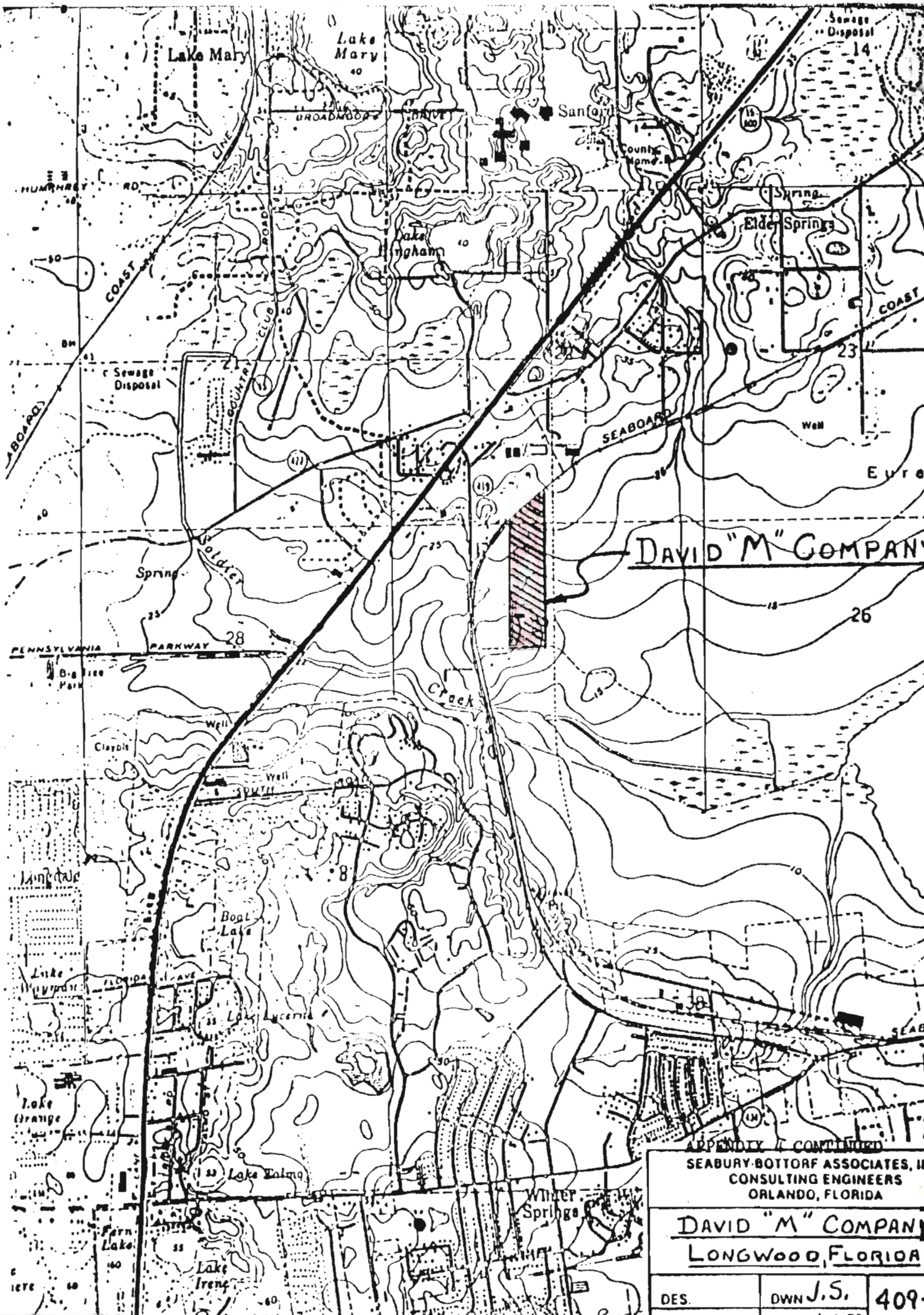
★ NOTE: SEE ATTACHED DRAWINGS FOR SOURCE LOCATIONS

SR 419

APPENDIX IV

DRAWN BY: MAW 8-2-83

NOT DRAWN TO SCALE



DAVID "M" COMPANY

APPENDIX A CONTINUED
 SEABURY-BOTTORF ASSOCIATES, INC.
 CONSULTING ENGINEERS
 ORLANDO, FLORIDA

DAVID "M" COMPANY
 LONGWOOD, FLORIDA

DES.	DWN J.S.	409-
------	----------	------

<u>COMPOUND</u>	<u># SAMPLES ANALYZED</u>	<u>AVG LBS. TOLUENE/100 LBS.</u>	<u>AVG LBS RUBBER/100 LBS.</u>
7002	48	58.9	41.1
5503	22	63.7	36.3
1101	48	47.8	52.2
6607	40	63.4	36.6
1100	23	49.3	50.7
1492	20	61.2	38.8
6609	31	66.5	33.5
106E	1	63.3	36.7
7001	2	62.1	37.9
108	12	32.7	67.3
1280	12	57.1	42.9
106	9	65.6	34.4
105	13	33.8	66.2
3303	2	64.9	35.1
7007	7	62.0	38

*0% VOC
in raw
matl.*

Note: Encompasses all but 3 compounds currently used in production and would be a suitable cross section of those compounds used most frequently.

8. How many pounds of volatile organic compounds will be in a gallon of applied coating?

The chart below delineates lbs. solvent/gallon of applied coating for the most frequently used compounds.

<u>COMPOUND</u>	<u>HI</u>	<u>LO</u>	<u>AVERAGE</u>	<u>NUMBER TESTED</u>
7007	5.6	4.8	5.3	16
5503	6.1	5.4	5.7	8
6607	5.8	4.6	5.2	21
1492	5.4	4.6	5.2	4
3303	5.3	5.6	5.4	2
1280	5.1	4.9	5.0	2
1100	5.6	4.6	4.9	9
1101	5.0	4.3	4.7	16
105	4.1	3.8	3.9	9

*HVOC
gal*

* The above examples are typical of those compounds current in use, however, they do not include all compounds currently in use.

9. a) How will Talc be received, stored and conveyed to the process?

Talc is received, stored and conveyed to the process in 55 pound bags.

- b) What air pollution control equipment or operation practices will be used to minimize Talc emissions?

Appendix III contains a drawing of the bag house and ancillary piping and collectors to be used in the talcing operation for the purpose of dust collection.

- c) How much Talc will be emitted to the atmosphere?

Talc emissions will be fugitive since the collector will be located in-doors. The dust collector supplier is being contacted in order to ascertain estimated collection and removal efficiencies for the purpose of a fugitive emissions estimate. We will transmit that information to you as soon as it becomes available.

10. Will solvent laden air discharged from the plant contain any particulates or oil mist?

The solvent laden air will contain only fugitive particulates and no oil mist. Should solvent recovery be the selected control device, air discharged will be filtered at an efficient removal level of 99.0%+ of particles \gg .3 microns in diameter.

11. a) What will be the particulate and VOC emissions from emission point No. 2 mills?

Particulate and VOC emissions combined from the milling operation would approximate 1.7 - 8.0 lbs/day. These numbers were generated by weighing all the individual components to be added to a batch before milling and then weighing the total batch post milling and noting the lost material. The plasticizers used in the milling process are not appreciably volatile. Appendix IV contains several example data sheets on plasticizers commonly used in the milling operation and Appendix V contains volatility data on many plasticizers used in the rubber industry.

talc

S.E. of plant description

I/c talc emissions?

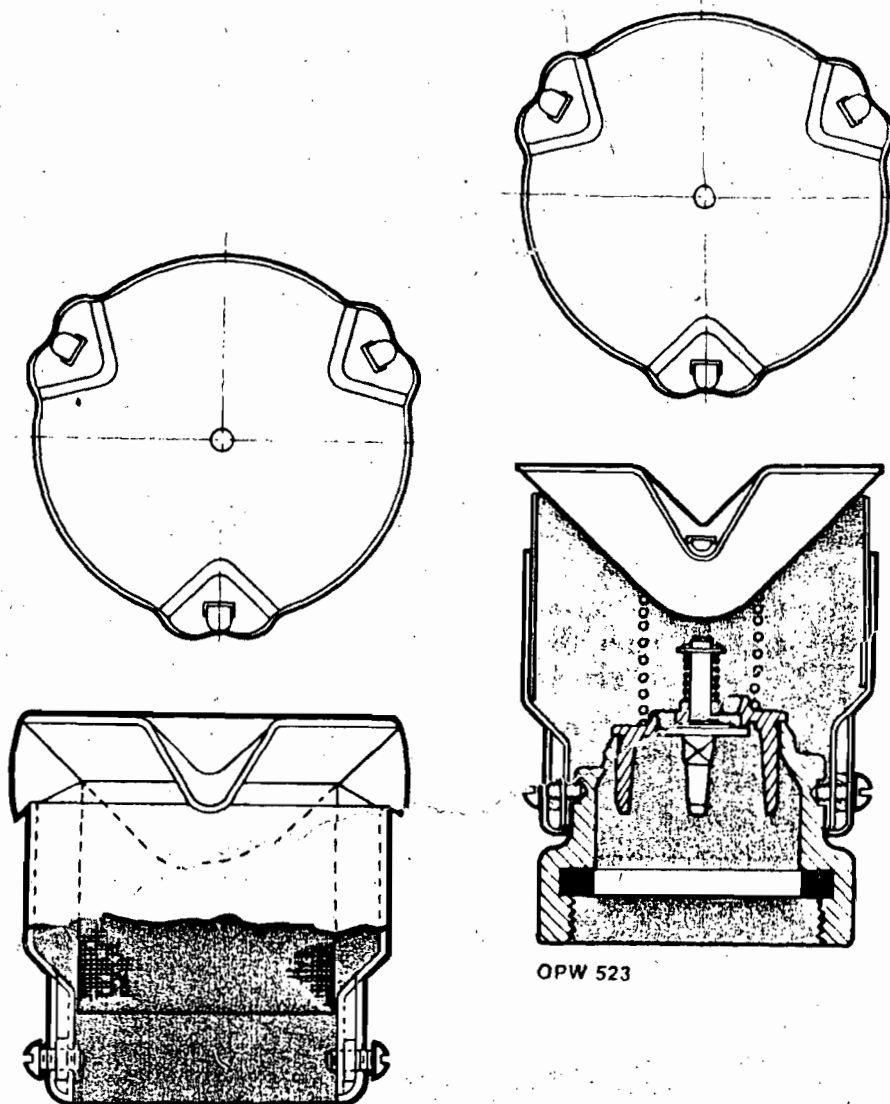
PM
I/c How much PM

No. 2 Mill Emissions

- b) What control equipment will be used to limit particulate emissions from the mills?

Since nearly all rubber compounds are compounded from master-batches, particulate emissions are quite low and as such no control device is currently proposed for the operation, see (11a) above.^c

no PM from
mills
(limit VE)



OPW 23

OPW 523

OPW 523

OPW 23, 523, 523-S VENTS

The OPW 23 Vent has an aluminum body and a 40-mesh brass screen. The OPW 23 is an open vent and directs vapors upward in accordance with NFPA Code 30. Set screws make installation easy. Available in 1½", 2" and 3" sizes.

The OPW 523 Pressure Vacuum Vent is an upward vapor discharge vent and is available with either an 8 oz. or 12 oz. pressure setting and a ½ oz. vacuum setting.

The 523 is available in 2" size, and is attached to the 2" threaded vent pipe.

The OPW 523-S Pressure Vacuum Vent is similar to the 523 above, except it is attached to the vent line with set screws rather than pipe threads. A gasket is provided.

Available in 2" size.

The rated maximum flow pressure drop for all OPW 523 Vents is 28 oz. per square inch at 7000 SCFH.



Wheelabrator-Frye Inc.

Graphic Supplies Division

7520 University

Des Moines, Iowa 50311

Tel. (515) 274-4901

November 1, 1983

Mr. Willard Hanks
Engineer
State of Florida
Department of Environmental Regulation
2600 Blair Stone Road
Twin Towers Building
Tallahassee, FL 32301

DER
NOV 02 1983
DAQM

Dear Mr. Hanks:

I'd like to take this opportunity to thank you, Bill Thomas and Nancy Wright for the courtesies extended myself and Herb Mycroft during the course of our recent meeting in Tallahassee. In accordance with our conversations at that time, please find enclosed additional data gathered pursuant to Mr. C.H. Fancy's request for same dated September 2, 1983.

Should you require further information and/or assistance in processing David "M" Company's air pollution permit application, please don't hesitate to contact me directly.

Sincerely,

Michael A. Ware

Michael A. Ware
Graphics Div. Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

MAW;bjw
Encl.

cc: Wayne Brady
Jerry Spangler
Herb Mycroft
Bob Baddeley
J. H. Gamble
T. J. Lucas

SUPPLEMENT TO DAVID 'M' COMPANY
APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCES

Prepared By: Michael A. Ware

1. What will be the maximum production and operation time (hour/day) of the plant?

The maximum uncontrolled emissions of the plant will be 300 tons/year based upon a two shift, 16 hours/day operation. The potential emissions incorporating an end of pipe control device capable of delivering 90% removal efficiency would approximate 30 tons/year. The 300 ton/year two shift operation is based upon the following historical information:

may be wrong 300 T/yr for my code

no process insight

<u>YEAR</u>	<u>OPERATING HOURS/DAY</u>	<u>ESTIMATED % NORMAL BUSINESS VOLUME</u>	<u>ANNUAL EMISSIONS</u>
1982	10	90%	271 Tons
1983	16	80%	255 Tons (Annualized Estimate)

2. What percent of the maximum production was the emission and raw material data in the application based upon?

The application was based upon 80% of the historically derived 300 tons/year uncontrolled emission number and reflected current economic conditions which approximate 70-80% of the usual demand for lithographic printing blankets. Should blanket demand return to 100% of the normal level, 300 tons uncontrolled and/or 30 tons controlled would approximate annual emissions without subtracting rubber cement containing VOC's which are disposed of at Emille, Alabama.

30 Tpy VOC Emission

3. Will solvents other than MEK and Toluene be used at this plant?

Yes, small amounts of Naptha and Isopropyl Alcohol will be used at this facility for the purpose of manually cleaning blankets during inspection. The additional fugitive type emissions from the use of these materials will approximate

5,000 lbs/yr Ispropyl Alcohol
2,000 lbs/yr Naptha

Limit on other VOC

4. How many gallons per year of each solvent will be consumed at the maximum production of the plant?

84000 Gallons
MAX
PROD
of
plant

87,000 gal Toluene x 7.2#/gal = 298.8 TONS

Approximately 75,000 gallons Toluene and 9,000 gallons MEK will be used at the maximum production of the plant. Were the use of MEK to be discontinued in order to facilitate efficient recovery of Toluene, approximately 83,000 gallons of Toluene would be used annually at the maximum production of the plant.

5. How will the solvents be shipped to the plant? Submit a drawing showing the storage area and piping to the mixing area.

The solvents will be shipped via bulk tanker and Appendix I is a conceptualization of the storage area and piping to the mixing area.

6. a) How many storage tanks will be built at the plant?

Storage tanks

Five, two for #2 diesel fuel, two for Toluene storage and one for MEK storage.

- b) What air pollution control equipment will be used to minimize VOC emissions from the tanks?

S.C. or jet description
Storage tank
controls

A coaxial vapor recovery system of the type 1 or 2 nature will be installed to minimize VOC emissions during off loading. In addition, an OPW 523, or 523-S pressure vacuum vent will be installed on each tank vent line to minimize VOC emissions during storage. Appendix II contains example information on coaxial vapor recovery and OPW 523 and 523-S vents like those proposed for installation.

- c) What will be emissions be from the tanks? Submit emissions calculation.

I/C
emission from
Storage tanks?

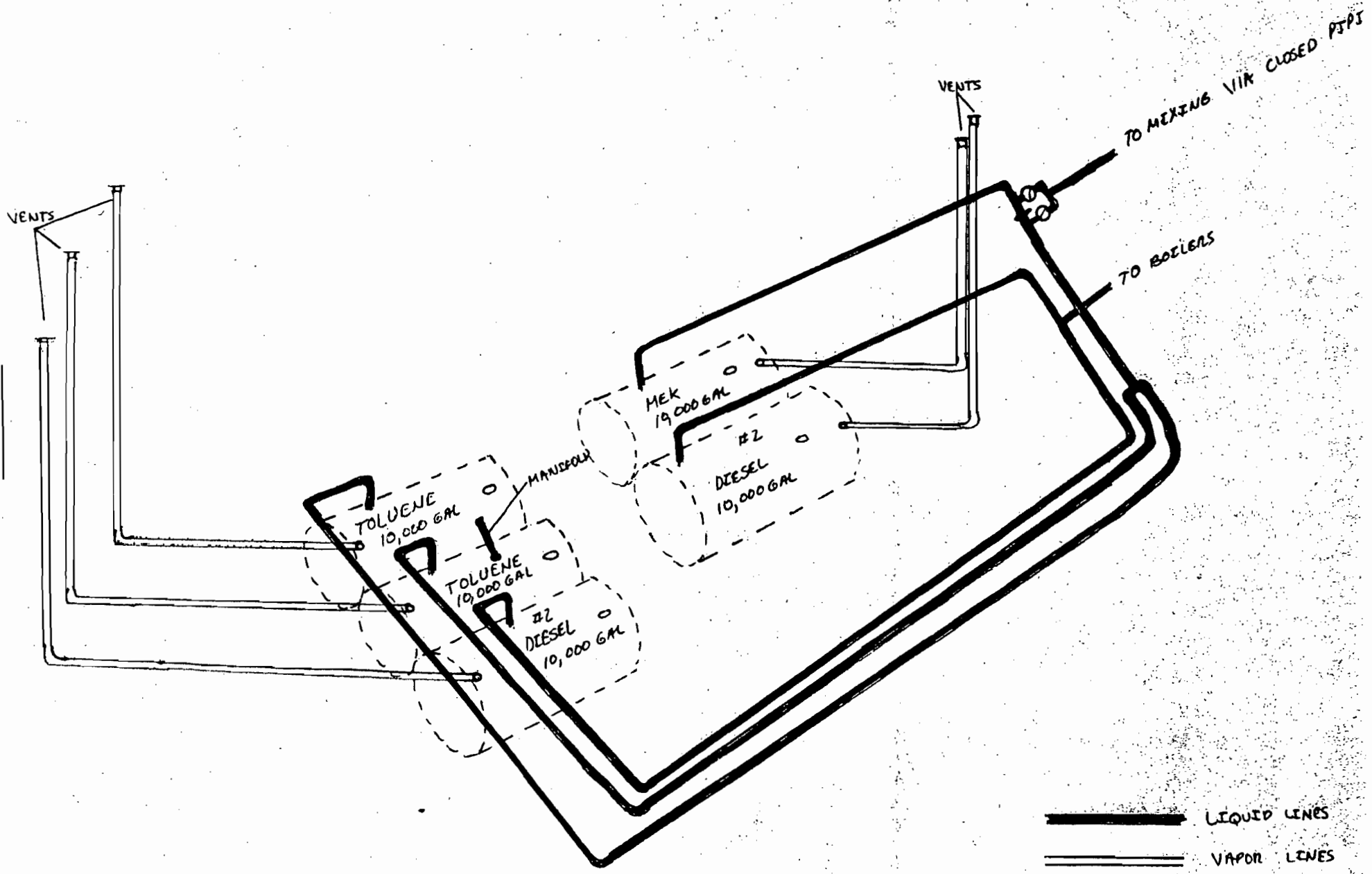
The equipment supplier has not as yet provided information concerning expected emissions from tanks equipped with coaxing recovery and pressure vents. We will endeavor to obtain this information as soon as possible.

7. What will be the mixing ratio of solvent to rubber in lbs/lb?

The chart below outlines the mixing ratio of solvent to rubber on a 100 lb. batch basis.

EXISTING SOLVENT STORAGE / DELIVERY SYSTEM

APPENDIX I



————— LIQUID LINES
————— VAPOR LINES

TANKS: UNDERGROUND
PLUMBING: UNDERGROUND TO PUMP

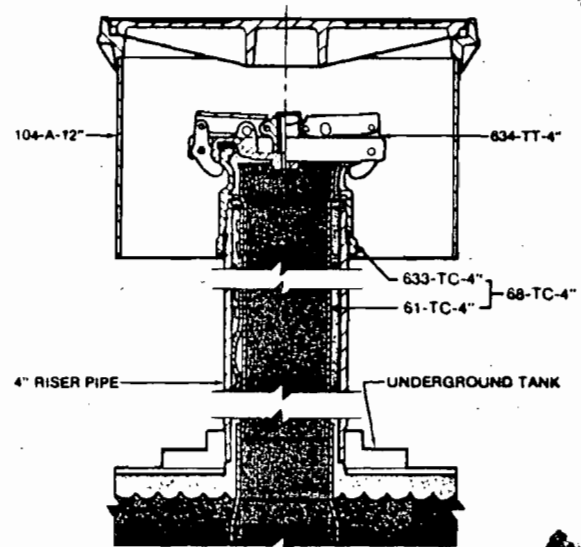
COAXIAL

SECTION B**Coaxial Vapor Recovery Equipment
FOR USE IN STAGE I AND STAGE II SYSTEMS****System C****TYPE 1 - 68-TC - 3" & 4" Series Fittings
(Non Poppeted)****TYPE 2 - 68-TCP - 3" & 4" Series Fittings
(Poppeted)**

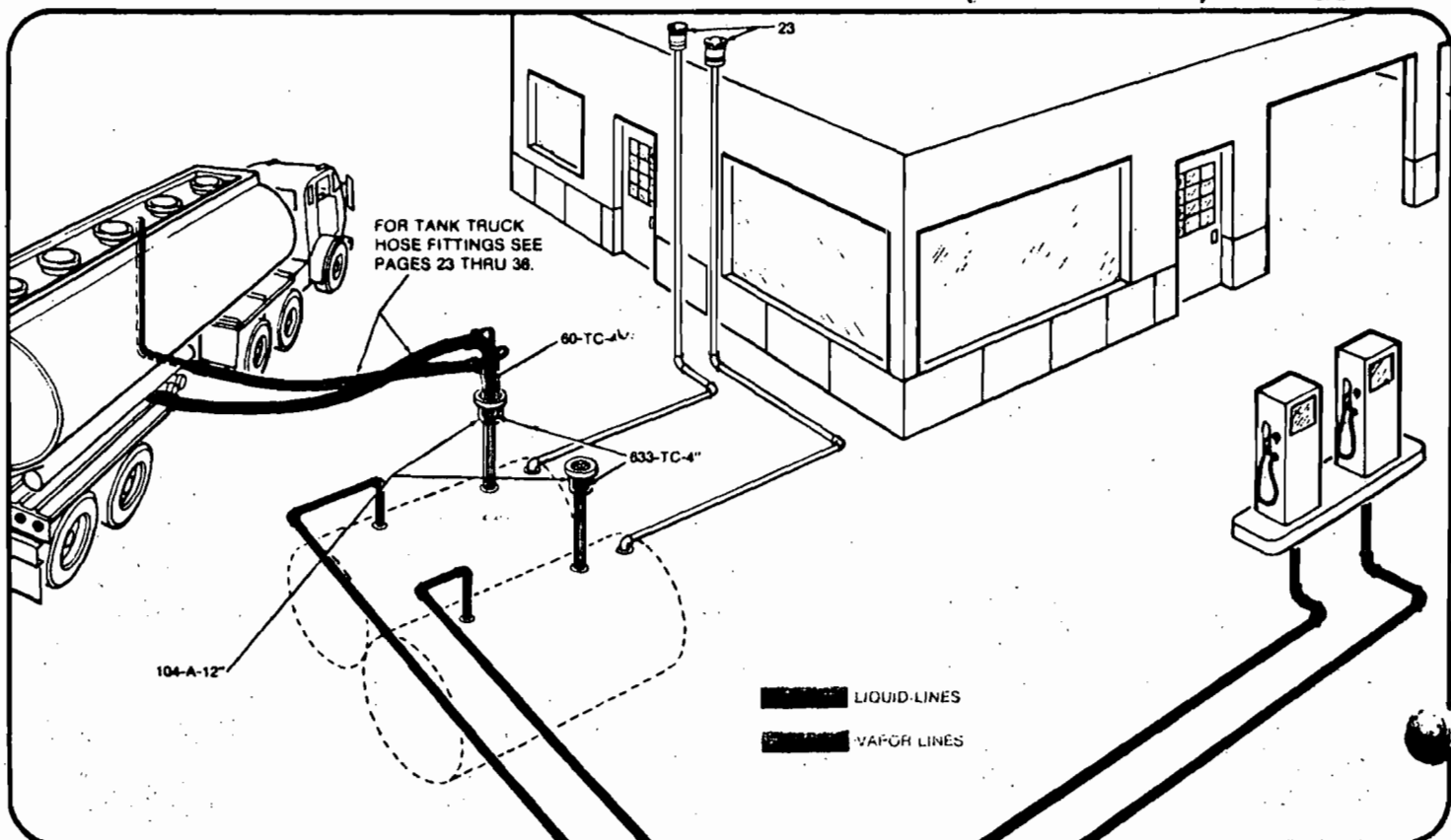
An easy, inexpensive means to convert existing conventional 3" & 4" size underground storage tank fill pipes to coaxial installations. Normally little to no digging or tearing up of concrete is necessary. Simply remove existing tight fill adaptor and fill pipe (if installation is so equipped), inserting Type 1 or Type 2 coaxial fill tube and adaptor as required.

NOTE: Type 2 poppeted fittings are certified by California Air Resources Board (CARB)

SHOWN BELOW IS A TYPE 1 - 68TC-4" TYPICAL INSTALLATION

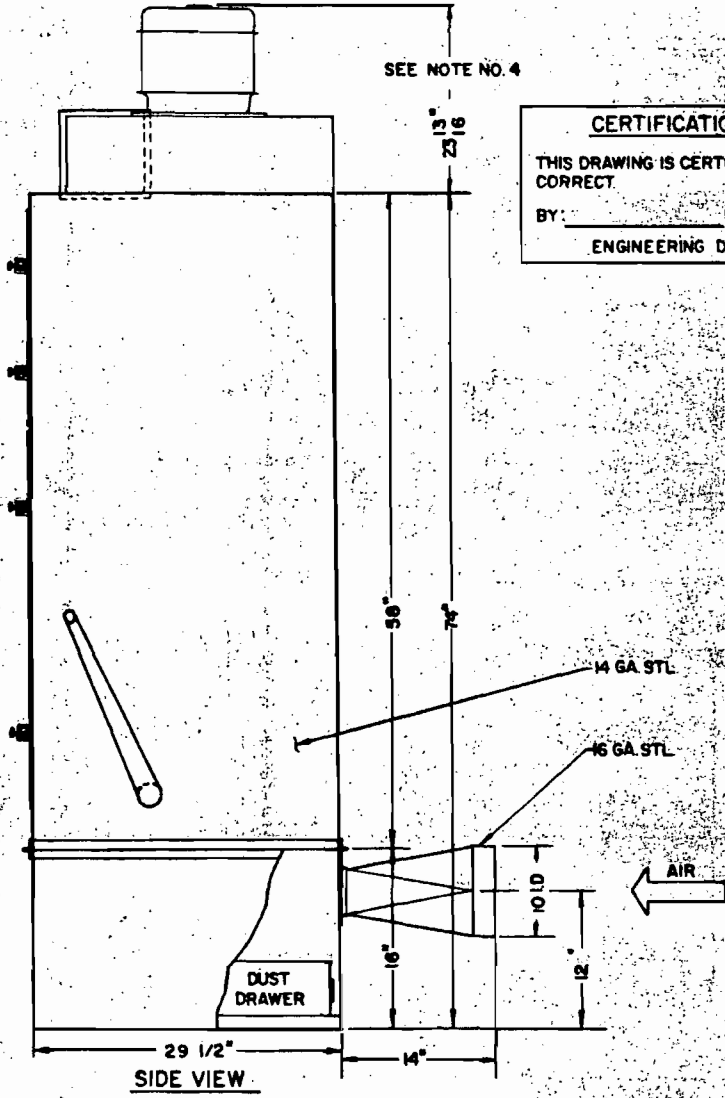
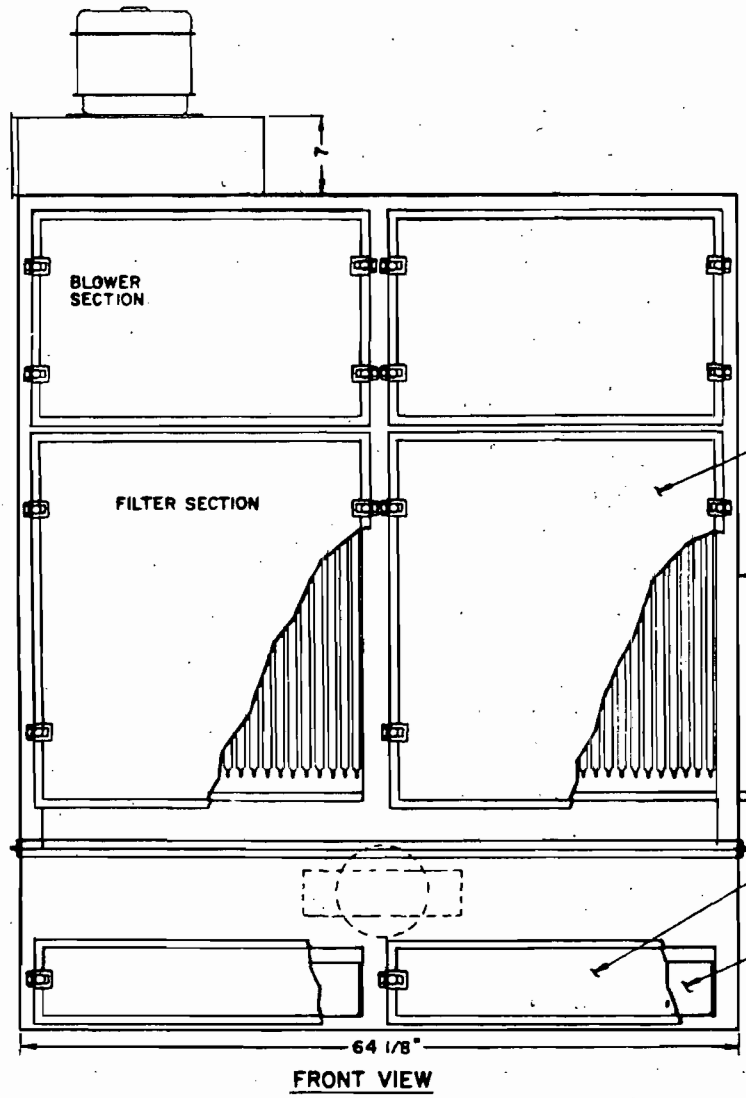


**TYPE 1 68-TC - 4" SERIES
(NON-POPPETED) FITTINGS**



Talc bag Collector

APPENDIX III



SEE NOTE NO. 4

CERTIFICATION

THIS DRAWING IS CERTIFIED TO BE CORRECT.

BY: _____ DATE: _____

ENGINEERING DEPT.

FRONT VIEW

SIDE VIEW

... MULTIPLE RATING TABLES ...

C.F.M.	F.P.M.	EXT. S.P.	C.F.M.	F.P.M.	EXT. S.P.
3625	6654	4.65	1912	3351	12 15
3365	6179	5.70			
2995	5491	7.38			
2390	4386	9.95			

- SPECIFICATIONS**
1. MOTOR SHALL BE 10 HP ON NO. 124-20 5-10 3600 RPM, 230-460/90/3.
 2. DUST STORAGE CAPACITY SHALL BE 4.4 CU. FT.
 3. FILTER AREA SHALL BE 400 SQ. FT. (40 FILTERS 30" x 24" EACH.)
 4. NET WT. 1100 SHIPPING WT. 1475

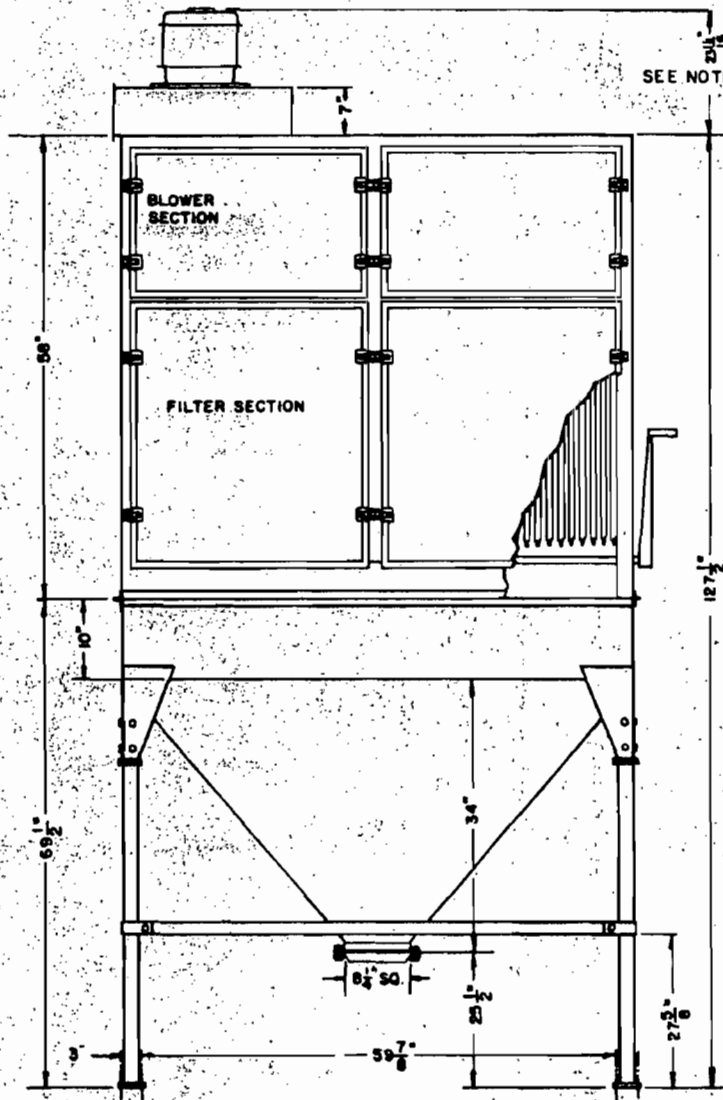
- NOTES**
1. THIS DRAWING IS FOR A 124-20-5-10 DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST.
 2. THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
 3. ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR & WHEEL ASS'Y.
 4. MOTOR HEIGHT BASED ON STOCK, CAN VARY WITH MFRS.

TORIT DIVISION, DONALDSON COMPANY, INC.
P.O. BOX 1289 - MINNEAPOLIS, MINNESOTA 55440

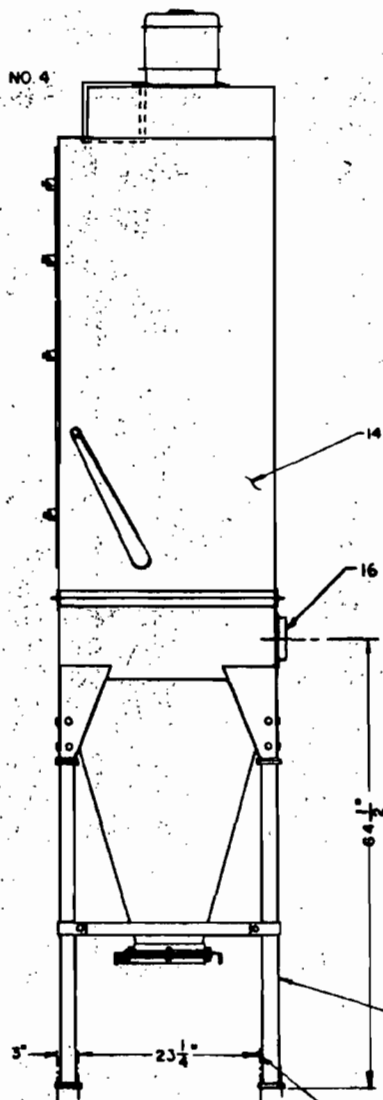
SPECIFICATION CONTROL DRAWING
NO. 124-20-5-10 DUST COLLECTOR

DRAWN: T.H.H.
APP'D: J.H.H.

DRWG. NO: **124-020**
REV. E 2-25-72

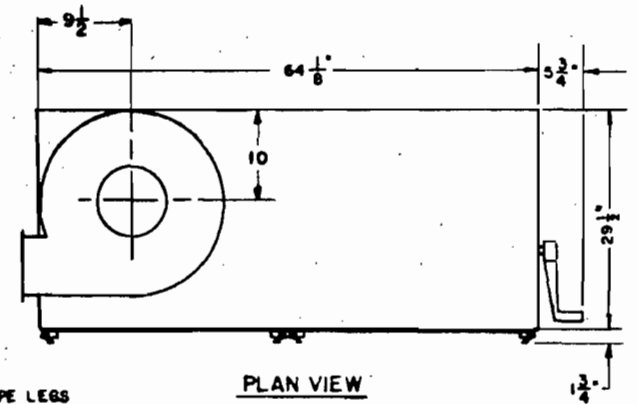
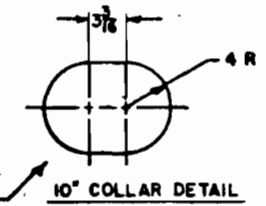


FRONT VIEW



SIDE VIEW

CERTIFICATION
 THIS DRAWING IS CERTIFIED TO BE CORRECT.
 BY: _____ DATE: _____
 ENGINEERING DEPT.



PLAN VIEW

MULTIPLE RATING TABLES					
C.F.M.	F.P.M.	EXT. S.P.	C.F.M.	F.P.M.	EXT. S.P.
3625	6654	4.65	1912	3351	12.15
3365	6179	5.70			
2995	5491	7.38			
2390	4386	9.95			

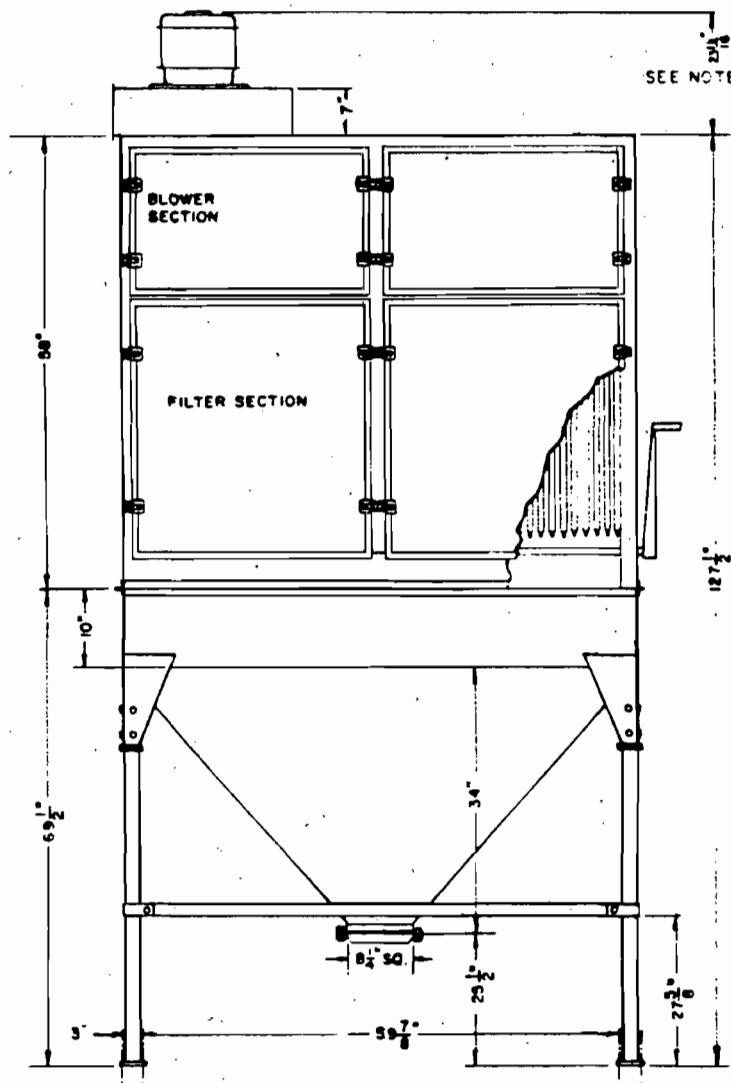
SPECIFICATIONS	
1. MOTOR SHALL BE 10 HP 3600 RPM, 230-460/60/3.	
2. HOPPER CAPACITY SHALL BE 16.0 CU. FT.	
3. FILTER AREA SHALL BE 400 SQ. FT. (40 FILTERS 30" X 24" EACH)	
4. INLET LOCATION: CENTERED ON BACK OR CENTERED ON EITHER END	
5. NET WT. 1070 SHIPPING WT. 1365	
	1100 1405

NOTES
1. THIS DRAWING IS FOR A 124-20-5-10H DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST
2. THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
3. ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR AND WHEEL ASY
4. MOTOR HEIGHT BASED ON STOCK CAN VARY WITH MFRS.

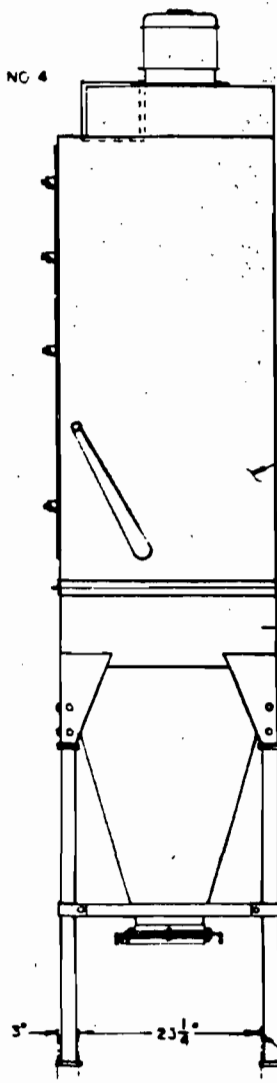
TORIT DIVISION, DONALDSON COMPANY, INC.
 P.O. BOX 1289 · MINNEAPOLIS, MINNESOTA 55440

SPECIFICATION CONTROL DRAWING
 NO. 124-20-5-10H DUST COLLECTOR

DRAWN: T. [Signature] DRWG. NO. 124-021
 APP'D: [Signature] REV. C 1-82

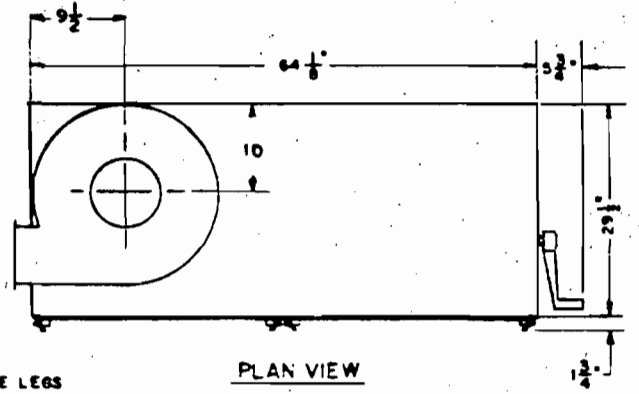
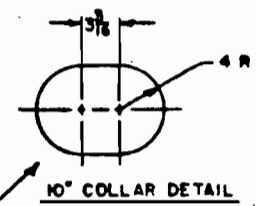


FRONT VIEW



SIDE VIEW

CERTIFICATION
 THIS DRAWING IS CERTIFIED TO BE CORRECT.
 BY: _____ DATE: _____
 ENGINEERING DEPT.



PLAN VIEW

LOCATING DIMS FOR 9/16" DIA. MTG HOLES

MULTIPLE RATING TABLES					
C.F.M.	F.P.M.	EXT. S.P.	C.F.M.	F.P.M.	EXT. S.P.
3625	6654	4.65	1912	3351	12.15
3365	6179	5.70			
2995	5491	7.38			
2390	4386	9.95			

SPECIFICATIONS	
1. MOTOR SHALL BE 10 HP 3500 RPM, 230-460/60/3.	
2. HOPPER CAPACITY SHALL BE 160 CU. FT	
3. FILTER AREA SHALL BE 400 SQ. FT (40 FILTERS 30" X 24" EACH)	
4. INLET LOCATION, CENTERED ON BACK OF CENTERED ON OTHER END	
5. NET WT. 1070 SHIPPING WT. 1365 1100 1405	

- NOTES**
1. THIS DRAWING IS FOR A 124-20-5-10H DUST COLLECTOR. ANY DEPARTURE FROM THIS DESIGN SHALL REQUIRE SPECIAL FABRICATION AND AN INCREASE IN COST
 2. THIS DRAWING IS NOT CERTIFIED FOR CONSTRUCTION PURPOSES UNLESS PROPERLY SIGNED BY TORIT.
 3. ALLOW A MIN. CLEARANCE OF 5" HEAD ROOM FOR REMOVAL OF MOTOR AND WHEEL ASY
 4. MOTOR HEIGHT BASED ON STOCK CAN VARY WITH MFRS.

TORIT DIVISION, DONALDSON COMPANY, INC.
 P.O. BOX 1299 - MINNEAPOLIS, MINNESOTA 55440

SPECIFICATION CONTROL DRAWING
NO. 124-20-5-10H DUST COLLECTOR

DRAWN: T. W. J. DRWG. NO. 124-021
 APP'D: [Signature] REV. C 1-88

transfer agent; fungistat
plant disease control;
tant for polyesters.

OOH.
crystals; b.p. sublimes;
in hot water, alcohol.

(C₆H₅)₂CHCN.
e powder; m.p. 73-
very soluble in alcohol.
cetic acid; synthesis of

e. See diphenadione.

aniline (C₆H₅)₂NH.
sh crystals. Soluble in
ohol, and ether; insolu-
p. 52.85°C; b.p. 302°C;
on temp. 1173°F; com-

rmula weights of aniline
in autoclave. The prod-
chloric acid to remove
residue is distilled.
ce and fused.

ce and fused.
-lined paper bags; fiber
skin. Tolerance, 10 mg

and accelerators; solid
dyes; pharmaceuticals;
preservation of apples;
nalytical chemistry.

adamsite; phenarsazine
H(C₆H₅)₂.

stials. Sublimes readily.
410°C (dec); insoluble
e, xylene, carbon tetra-

enylamine with arsenic
halation and ingestion:

od treating.
Poison label. Not ac-
Rail) Tear gas label.

(C₆H₅)₂.
48-250°C; insoluble in
soluble in toluene.

length shifter in solution

enyl.
C₆H₅NHC₆H₅.
soluble in water; slightly
atic hydrocarbons. Sen-

d fuming sulfuric acid.

See benzidine.
d nitrites.

diphenylbromide (C₆H₅)₂AsBr.
Properties: White crystals. M.p. 54-56°C.
Derivation: (a) Hydrobromic acid and diphenylarse-
nious oxide are heated together for about 4 hours at
115-120°C; (b) by action of arsenic tribromide on
triphenyl arsine at 300-350°C.
Hazard: Highly toxic; strong irritant.
Shipping regulations: (Rail, Air) Arsenical compounds,
solids, n.o.s.. Poison label.

1,3-diphenyl-2-buten-1-one. See dyprnone.

diphenylcarbazine (C₆H₅)₂NHNH₂CO.
Properties: White crystals or flakes. Insoluble in
water; soluble in alcohol and benzene. M.p. 173°C.
Decomposes in light.
Derivation: Phenylhydrazine and urea.
Use: Determination of copper and other metals.

diphenylcarbinol. See benzhydrol.

diphenyl carbonate (C₆H₅O)₂CO.
Properties: White, crystalline solid. Can be halogen-
ated and nitrated in characteristic manner. Readily
undergoes hydrolysis and ammonolysis. Soluble in
acetone, hot alcohol, benzene, carbon tetrachloride,
ether, glacial acetic acid and other organic solvents;
insoluble in water. B.p. 302°C; m.p. 78°C; sp. gr.
1.1215 (87.4°C).
Grade: Technical.
Uses: Plasticizer and solvent; synthesis of polycar-
bonate resins.

diphenylchloroarsine (C₆H₅)₂AsCl.
Properties: Colorless crystals or dark-brown liquid,
which slowly becomes semi-solid. Decomposed by
water (slowly). Soluble in carbon tetrachloride,
chloropicrin, phenyldichloroarsine; practically insolu-
ble in water. Sp. gr. 1.363 (40°C) (solid), or 1.358
(45°C) (liquid); b.p. 333°C (in CO₂ atmosphere);
m.p. 41°C.
Derivation: Benzene and arsenic trichloride are heated
in presence of aluminum chloride.
Grade: Technical.
Hazard: Highly toxic by inhalation; strong irritant to
tissue.
Use: Military poison gas.
Shipping regulations: (Rail) Tear Gas label (Air)
Poison label. Not accepted on passenger planes.

diphenyldecyl phosphite (C₆H₅O)₂POC₁₀H₂₁.
Properties: Nearly water-white liquid; sp. gr. 1.200
(25/15.5°C); m.p. 18°C; refractive index 1.5160
(25/D). Combustible.
Uses: Chemical intermediate; stabilizer for poly-
olefin and polyolefin resins.

diphenyldichlorosilane (C₆H₅)₂SiCl₂.
Properties: Colorless liquid; b.p. 305°C; f.p. 25°C;
sp. gr. 1.19 (20°C); refractive index (n_D²⁵) 1.5150;
flash point (COC) 288°F. Readily hydrolyzed by
moisture, with liberation of hydrochloric acid.
Combustible.
Derivation: (a) Reaction of powdered silicon
chlorobenzene in the presence of copper powder as
catalyst; (b) reaction of phenylmagnesium chloride
with silicon tetrachloride.
Grade: Technical.
Hazard: Highly toxic; strong irritant to tissue.
Use: Intermediate for silicone lubricants.

diphenyldodecyl phosphite (C₆H₅O)₂POC₁₂H₂₅.
Properties: Nearly water-white liquid; sp. gr. 1.200
(25/15.5°C); m.p. 18°C; refractive index 1.5160
(25/D). Combustible.
Uses: Chemical intermediate; stabilizer for poly-
olefin and polyolefin resins.

diphenyldichlorosilane (C₆H₅)₂SiCl₂.
Properties: Colorless liquid; b.p. 305°C; f.p. 25°C;
sp. gr. 1.19 (20°C); refractive index (n_D²⁵) 1.5150;
flash point (COC) 288°F. Readily hydrolyzed by
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Combustible.
Derivation: (a) Reaction of powdered silicon
chlorobenzene in the presence of copper powder as
catalyst; (b) reaction of phenylmagnesium chloride
with silicon tetrachloride.
Grade: Technical.
Hazard: Highly toxic; strong irritant to tissue.
Use: Intermediate for silicone lubricants.

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Properties: Colorless liquid; b.p. 305°C; f.p. 25°C;
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flash point (COC) 288°F. Readily hydrolyzed by
moisture, with liberation of hydrochloric acid.
Combustible.
Derivation: (a) Reaction of powdered silicon
chlorobenzene in the presence of copper powder as
catalyst; (b) reaction of phenylmagnesium chloride
with silicon tetrachloride.
Grade: Technical.
Hazard: Highly toxic; strong irritant to tissue.
Use: Intermediate for silicone lubricants.

Phosphorogens

APPENDIX IV

diphenyldichlorosilane (C₆H₅)₂SiCl₂.
Derivation: Reaction of didodecyldichlorosilane with
phenyl lithium.
Hazard: Probably toxic.
Use: High-temperature lubricant.
diphenyldimide. See azobenzene
diphenyldimethoxysilane (C₆H₅)₂Si(OCH₃)₂.
Properties: Liquid; sp. gr. 1.080 (25°C); b.p. 191°C
(53 mm); refractive index 1.5404 (25°C). Soluble in
acetone, benzene, methyl alcohol; combustible.
Hazard: Probably toxic.
Uses: Treatment of powders, glass, paper, and fabrics.

diphenyleneimine. See carbazole
alpha-diphenylenemethane. See fluorene.
diphenylene oxide (dibenzofuran) C₁₂H₈O (tricyclic).
Properties: Crystalline solid; m.p. 87°C; b.p. 288°C;
insoluble in water; slightly soluble in alcohol, ether
and benzene.
Derived from coal-tar.
Hazard: Probably toxic.
Use: Insecticide.

1,1-diphenylethane. See uns-diphenylethane.

1,2-diphenylethane. See sym-diphenylethane.

uns-diphenylethane (1,1-diphenylethane)
(C₆H₅)₂CHCH₃.
Properties: Colorless liquid. Soluble in chloroform,
ether, carbon disulfide. B.p. 286°C; sp. gr. 1.004
(20°C); f.p. -21.5°C. Flash point 264°F; combus-
tible.
Derivation: Action of acetaldehyde on benzene in
presence of concentrated sulfuric acid.
Uses: Solvent for nitrocellulose; organic synthesis.

**sym-diphenylethane (bibenzyl; dibenzyl; 1,2-diphenyl-
ethane) C₆H₅CH₂CH₂C₆H₅.**
Properties: White, crystalline needles or small plates.
Soluble in alcohol, chloroform, ether, carbon di-
sulfide; insoluble in water. Sp. gr. 0.9782; b.p.
284°C; m.p. 52°C.
Derivation: (a) By treating benzyl chloride with met-
allic sodium. (b) Action of benzyl chloride on ben-
zylmagnesium chloride.
Use: Organic synthesis.

diphenyl ether. See diphenyl oxide.

diphenylethylene. See stilbene.

N,N-diphenylethylenediamine (ethyl diphenyldiamine)
C₆H₅NHCH₂CH₂NHC₆H₅.
Properties: Cream-colored solid; sp. gr. 1.14; soften-
ing point 54°C; insoluble in water; soluble in ace-
tone, ethylene dichloride, benzene, and gasoline.
Use: Antioxidant in rubber compounding.

diphenylglycolic acid. See benzoic acid.

diphenylguanidine (DPG; melaniline) HN₂C(NHC₆H₅)₂.
Properties: White powder; bitter taste; slight odor;
sp. gr. 1.13; m.p. 147°C; decomposes above 170°C;
soluble in ethyl alcohol, carbon tetrachloride, chlo-
roform, hot benzene and toluene; slightly soluble in
water.
Derivation: Treatment of aniline with cyanogen chlo-
ride

diphenylmethane (benzylbenzene) (C₆H₅)₂CH₂.
Properties: Long colorless needles. Soluble in alcohol
and ether; insoluble in water. Sp. gr. 1.0056; m.p.
26.5°C; b.p. 264.7°C. Flash point 266°F; combus-
tible.
Derivation: Condensation of benzyl chloride and ben-
zene in presence of aluminum chloride.
Hazard: Probably irritant and narcotic in high con-
centrations.
Uses: Organic synthesis; dyes; perfumery.

diphenyl isophthalate (DPIP)
C₆H₄(OCC₆H₄COOC₆H₅)₂.
Properties: White solid; m.p. 138-139°C. Combustible.
Use: Manufacture of polybenzimidazoles, high tem-
perature-resistant polymers.

diphenylketone. See benzophenone.

diphenylmethane (benzylbenzene) (C₆H₅)₂CH₂.
Properties: Long colorless needles. Soluble in alcohol
and ether; insoluble in water. Sp. gr. 1.0056; m.p.
26.5°C; b.p. 264.7°C. Flash point 266°F; combus-
tible.
Derivation: Condensation of benzyl chloride and ben-
zene in presence of aluminum chloride.
Hazard: Probably irritant and narcotic in high con-
centrations.
Uses: Organic synthesis; dyes; perfumery.

**diphenylmethane-4,4'-diisocyanate (MDI; methylene-di-
paraphenylene isocyanate; methylenebis(phenyl iso-
cyanate) CH₂(C₆H₄NC=O)₂.**
Properties: Light-yellow, fused solid; solidification
point 37°C; sp. gr. (70°C) 1.197; soluble in acetone,
benzene, kerosine and nitrobenzene; combustible.
Derivation: para, para'-Diaminodiphenylmethane and
phosgene.
Hazard: Highly toxic by inhalation of fumes. Strong
irritant. Tolerance, 0.02 ppm in air.
Uses: Preparation of polyurethane resin and spandex
fibers; bonding rubber to rayon and nylon.

diphenylmethanol. See benzhydrol.

diphenylmethyl bromide (benzhydryl bromide)
BrCH(C₆H₅)₂.
Properties: Solid; m.p. 45°C; b.p. 193°C (26 mm).
Decomposes in hot water; soluble in alcohol; very
soluble in benzene.
Hazard: Strong irritant to eyes and skin.
Use: Organic synthesis.
Shipping regulations: (Air) Corrosive label. (Rail)
Corrosive material, n.o.s., White label.

diphenylmethylchlorosilane (C₆H₅)₂(CH₂)₂SiCl.
Properties: Colorless liquid; sp. gr. 1.107 (25°C); b.p.
295°C; flash point 135°F. Combustible.
Derivation: Grignard reaction of diphenyldichlorosi-
lane with methylmagnesium chloride.
Hazard: Moderate fire risk. Probably toxic.
Use: Intermediate; end stopper for silicone oils.

diphenylnaphthylenediamine C₁₀H₆(NHC₆H₅)₂.
Properties: Silvery, crystalline plates. Slightly soluble
in alcohol; insoluble in water. M.p. 164°C.
Derivation: By heating 2,7-dihydroxynaphthalene
with aniline and aniline hydrochloride.
Use: Organic synthesis.

diphenylnitrosamine. See N-nitrosodiphenylamine.

2,5-diphenyloxazole (DPO) OOC₆H₄CC₆H₅.
Properties: White, fluffy solid; m.p. 70-72°C.
Grade: Scintillation.

Superior numbers refer to Manufacturers of Trade Mark Products. For page number see Contents.

types: curing and hardening of polymers (e.g., core-binding resins); corrosion inhibitor; propellant.
Shipping regulations: (Rail) Flammable liquids, n.o.s., Red label. (Air) Flammable Liquid label.

triethylborane (triethylborine; boron triethyl).
(C₂H₅)₃B.

Properties: Colorless liquid. Sp. gr. (25°C); flash point -32°F; f.p. -93°C; b.p. 95°C; refractive index 1.3971; heat of combustion 20,000 Btu/lb. Miscible with most organic solvents; immiscible with water.

Derivation: Reaction of triethylaluminum and boron halide, or diborane and ethylene.

Hazard: Flammable, dangerous fire risk. Ignites spontaneously in air. Highly toxic by inhalation; strong irritant. Reacts violently with water and oxidizing materials.

Uses: Igniter or fuel for jet and rocket engines; fuel additive; olefin polymerization catalyst; intermediate.

Shipping regulations: (Rail) Pyrophoric liquid, n.o.s., Red label. (Air) Not acceptable.

triethyl borate (ethyl borate) (C₂H₅)₃BO₃.

Properties: Colorless liquid; mild odor. Hydrolyzes rapidly, depositing boric acid in finely divided crystalline form. B.p. 120°C; sp. gr. 0.863-0.864 (20/20°C); flash point 51.8°F (C.C.); wt./gal 7.20 lb. (20°C); refractive index 1.37311 (20°C).

Uses: Antiseptics; disinfectants; antiknock agent.

Hazard: Flammable, dangerous fire risk. Moderately toxic.

Shipping regulations: (Rail) Flammable liquid, n.o.s., Red label. (Air) Flammable Liquid label. (Air) Legal label name, ethyl borate.

triethylborine. See triethylborane.

triethyl citrate (ethyl citrate) C₃H₇(COOC₂H₅)₃.

Properties: Colorless, mobile liquid. Bitter taste; b.p. 294°C; b.p. (1 mm) 126-127°C; sp. gr. 1.136 (25°C); refractive index 1.4405 (24.5°C); pour point -50°F; solubility in water 6.5 g/100 cc; solubility in oil 0.8 g/100 cc. Flash point 303°F (COC); combustible. Low toxicity.

Derivation: Esterification of citric acid.

Grades: Technical; refined; F.C.C.

Containers: Metal drums and cans; tank cars.

Uses: Solvent and plasticizer for nitrocellulose and natural resins; softener; paint removers; agglutinant; perfume base; food additive (not over 0.25%).

triethylenediamine N(CH₂CH₂)₃N. Catalyst used in the production of polyurethanes. Combustible.

triethylene glycol (TEG) HO(C₂H₄O)₃H.

Properties: Colorless, hygroscopic, practically odorless liquid. Sp. gr. 1.1254 (20/20°C); b.p. 287.4°C; vapor pressure less than 0.01 mm (20°C); flash point 350°F (C.C.); wt./gal 9.4 lb (20°C); freezing point -7.2°C; viscosity 0.478 poise (20°C). Autoignition temp. 700°F. Soluble in water; immiscible with benzene, toluene and gasoline. Combustible; low toxicity.

Derivation: From ethylene and oxygen, as a by-product of ethylene glycol manufacture.

Grades: Technical; C.P.

Containers: 1-, 5-gal cans; 55-gal drums; tank cars.

Uses: Solvent and plasticizer in vinyl, polyester and polyurethane resins; dehydration of natural gas; humectant in printing inks; extraction solvent ("Udex" process).

triethylene glycol diacetate

CH₃COOCH₂CH₂OCH₂CH₂OCH₂CH₂OOCCH₃.

Properties: Colorless liquid. Sp. gr. (25°C) 1.112; re-

fractive index n_D(25°C) 1.437; b.p. 300°C; f.p. less than -60°C. Combustible; low toxicity.
Use: Plasticizer.

triethylene glycol dibenzoate

C₆H₅CO(OCH₂CH₂)₃OOC₆H₅.

Properties: Crystals; b.p. 210-223°C; m.p. 46°C; flash point 457°F (TOC); sp. gr. 1.168. Combustible. Low toxicity.

Uses: Plasticizer for vinyl resins; adhesives.

triethylene glycol dicaprylate (triethylene glycol dioc-
toate) C₁₇H₃₃COO(CH₂CH₂O)₃OCC₁₇H₃₃.

Properties: Clear liquid; sp. gr. 0.973 (20°C); acidity 0.3% max. (caprylic); moisture 0.05% max.; f.p. -3°C; b.p. 243°C (5 mm). Soluble in most organic solvents. Combustible; low toxicity.

Uses: Low-temperature plasticizer for elastomers.

triethylene glycol dichloride. See triglycol dichloride.

triethylene glycol didecanoate

C₁₀H₁₉COO(C₂H₄O)₃OCC₁₀H₁₉.

Properties: Colorless liquid. B.p. 237°C (2.0 mm); sp. gr. 0.9584 (20/20°C); viscosity 28.6 cp (20°C). Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol di(2-ethylbutyrate)

C₁₁H₂₁OCOCH₂(CH₂OCH₂)₃CH₂OCC₁₁H₂₁.

Properties: Light-colored liquid; sp. gr. 0.9946 (20/20°C); 8.3 lb/gal (20°C); b.p. 196°C (5 mm); vapor pressure 5.8 mm Hg (200°C); solubility in water 0.02% by wt (20°C); viscosity 10.3 cp (20°C). Flash point 385°F. Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol di(2-ethylhexoate)

C₁₃H₂₅OCOCH₂(CH₂OCH₂)₃CH₂OCC₁₃H₂₅.

Properties: Light-colored liquid; sp. gr. 0.9679 (20/20°C); 8.1 lb/gal (20°C); b.p. 219°C (5 mm); vapor pressure 1.8 mm Hg (200°C); insoluble in water; viscosity 15.8 cp (20°C). Flash point 405°F. Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol dihydroabietate

C₁₉H₃₁COO(C₂H₄O)₃OCC₁₉H₃₁.

Properties: Liquid. Sp. gr. (25°C) 1.080-1.090; refractive index (20°C) 1.5180; vapor pressure (225°C) 2.5; flash point 226°C; insoluble in water. Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol dimethyl ether

CH₃(OCH₂CH₂)₃OCH₃.

Properties: Water-white liquid; mild ether odor; sp. gr. (20/20°C) 0.9862; refractive index 1.4233 (n_D 20/D); flash point 232°F; b.p. (760 mm) 216.0°C; (100 mm) 153.6°C; f.p. -46°C. Autoignition temp. 1166°F. Completely soluble in water and hydrocarbons at 20°C. May contain peroxides. Combustible. Low toxicity.

Containers: Glass bottles; cans; 55-gal drums.

Uses: Solvent for gases; coupling immiscible liquids.

triethylene glycol dioctoate. See triethylene glycol dicaprylate.

triethylene glycol dipelargonate

C₁₇H₃₃COO(C₂H₄O)₃OCC₁₇H₃₃.

Properties: Clear liquid; sp. gr. 0.964 (20/20°C); b.p. 251°C (5 mm); f.p. +1 to -4°C; refractive index 1.4470 (23°C); flash point 410°F. Almost insoluble in water; soluble in most organic solvents. Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol dipropionate

C₇H₁₃CO(OCH₂CH₂)₃OOC₇H₁₃.

Properties: Colorless liquid. Sp. gr. (25°C) 1.066; refractive index (25°C) 1.436; b.p. (2 mm) 138-142°C; f.p. less than -60°C; solubility in water, 6.70% by weight. Combustible. Low toxicity.

Use: Plasticizer.

triethylene glycol monobutyl ether. See butoxytri-
glycol.

triethylenemelamine (tretamine; TEM; 2,4,6-tris(1-azi-
ridinyl)-s-triazine)

NC[N(CH₂)₂]NC[N(CH₂)₂]NC[N(CH₂)₂].

Properties: White, crystalline, odorless powder; m.p. 160°C (polymerizes); polymerizes readily with heat or moisture; soluble in alcohol, water, methanol, chloroform, and acetone.

Grade: N.F.

Hazard: Highly toxic.

Uses: Medicine (see nitrogen mustards); insecticide; chemosterilant.

triethylenephosphoramidate (tepa; tris-(1-aziridinyl)phos-
phine oxide; APO) (NCH₂CH₂)₃PO.

Properties: Colorless crystals; m.p. 41°C; soluble in water, alcohol and ether. Combustible.

Derivation: From ethyleneimine.

Hazard: Highly toxic. Strong irritant to skin and tis-
sue.

Uses: Medicine (see nitrogen mustards); insect steril-
ant. Also used with tetrakis(hydroxymethyl)phos-
phonium chloride (THPC) to form a condensation
polymer suitable for flameproofing cotton. See also
tris[1-(2-methyl)aziridinyl]phosphine oxide.

Shipping regulations: (Rail) White label. (Air) Corro-
sive label. Legal label name: tris(1-aziridinyl)-
phosphine oxide.

See also tepa.

triethylenetetramine NH₂(C₂H₄NH)₃C₂H₄NH₂.

Properties: Moderately viscous yellowish liquid. Less
volatile than diethylenetriamine but resembles it in
many other properties. Soluble in water. B.p. 277.5°C;
sp. gr. 0.9818 (20/20°C); m.p. 12°C; flash point
275°F (C.C.); wt 8.2 lb/gal (20°C). Combustible.
Autoignition temp. 640°F.

Grades: Technical; anhydrous.

Containers: Cans; drums; tank cars.

Hazard: Strong irritant to tissue. Causes skin burns
and eye damage.

Uses: Detergents and softening agents; synthesis of
dyestuffs, pharmaceuticals and rubber accelerators.

Shipping regulations: (Air) Corrosive label.

tri(2-ethylhexyl)phosphate [C₈H₁₇CH(C₂H₅)CH₂]₃PO₄.

Properties: Light-colored liquid; sp. gr. 0.9260 (20/
20°C); 7.70 lb/gal (20°C); b.p. 220°C (5 mm); vapor
pressure 1.9 mm Hg (200°C); insoluble in water;
viscosity 14.1 cp (20°C); pour point -74°C; flash
point 405°F. Combustible; low toxicity.

Use: Plasticizer.

tri(2-ethylhexyl) phosphite (C₈H₁₇O)₃P.

Properties: Straw-colored liquid; sp. gr. 0.897 (25/
15°C); m.p., glass at low temperature; refractive in-
dex 1.451 (n_D 25/D); flash point 340°F (COC). Com-
bustible; low toxicity.

Uses: Plasticizer; intermediate.

tri(2-ethylhexyl) trimellitate C₈H₁₇(COOC₂H₅)₃.

Properties: Clear liquid, mild odor; sp. gr. 0.992 (20/
Use

fractive index n (25°C) 1.437; b.p. 300°C; f.p. less than -60°C. Combustible; low toxicity.
Use: Plasticizer.

triethylene glycol dibenzoate

$C_{20}H_{22}O_6$; $CO(OCH_2CH_2)_3OOCOC_6H_5$.
Properties: Crystals; b.p. 210–223°C; m.p. 46°C; flash point 457°F (TOC); sp. gr. 1.168. Combustible. Low toxicity.
Uses: Plasticizer for vinyl resins; adhesives.

triethylene glycol dicaprylate (triethylene glycol dioctoate)

$C_{30}H_{58}O_6$; $COO(CH_2CH_2O)_3COOC_8H_{17}$.
Properties: Clear liquid; sp. gr. 0.973 (20°C); acidity 0.3% max. (caprylic); moisture 0.05% max; f.p. -3°C; b.p. 243°C (5 mm). Soluble in most organic solvents. Combustible; low toxicity.
Uses: Low-temperature plasticizer for elastomers.

triethylene glycol dichloride. See triglycol dichloride.

triethylene glycol didecanoate

$C_{26}H_{46}O_6$; $COO(C_{10}H_{19}O)_3COOC_{10}H_{19}$.
Properties: Colorless liquid. B.p. 237°C (2.0 mm); sp. gr. 0.9584 (20/20°C); viscosity 28.6 cp (20°C). Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol di(2-ethylbutyrate)

$C_{22}H_{40}O_6$; $(CH_3CH_2OCH_2)_2CH_2OCOC_4H_9$.
Properties: Light-colored liquid; sp. gr. 0.9946 (20/20°C); 8.3 lb/gal (20°C); b.p. 196°C (5 mm); vapor pressure 5.8 mm Hg (200°C); solubility in water 0.02% by wt (20°C); viscosity 10.3 cp (20°C). Flash point 385°F. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol di(2-ethylhexoate)

$C_{24}H_{44}O_6$; $(CH_3CH_2OCH_2)_2CH_2OCOC_6H_{13}$.
Properties: Light-colored liquid; sp. gr. 0.9679 (20/20°C); 8.1 lb/gal (20°C); b.p. 219°C (5 mm); vapor pressure 1.8 mm Hg (200°C); insoluble in water; viscosity 15.8 cp (20°C). Flash point 405°F. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol dihydroabietate

$C_{30}H_{50}O_6$; $COO(C_{10}H_{17}O)_3COOC_{10}H_{17}$.
Properties: Liquid. Sp. gr. (25°C) 1.080–1.090; refractive index (20°C) 1.5180; vapor pressure (225°C) 2.5; flash point 226°C; insoluble in water. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol dimethyl ether

$CH_3(OCH_2CH_2)_3OCH_3$.
Properties: Water-white liquid; mild ether odor; sp. gr. (20/20°C) 0.9862; refractive index 1.4233 (n_D 20/D); flash point 232°F; b.p. (760 mm) 216.0°C (100 mm) 153.6°C; f.p. -46°C. Autoignition temp. 1166°F. Completely soluble in water and hydrocarbons at 20°C. May contain peroxides. Combustible. Low toxicity.
Containers: Glass bottles; cans; 55-gal drums.
Uses: Solvent for gases; coupling immiscible liquids.

triethylene glycol dioctoate. See triethylene glycol dicaprylate.

triethylene glycol dipalargonate

$C_{28}H_{48}O_6$; $COO(C_{16}H_{31}O)_3COOC_{16}H_{31}$.
Properties: Clear liquid; sp. gr. 0.964 (20/20°C); b.p. 251°C (5 mm); f.p. +1 to -4°C; refractive index 1.4470 (23°C); flash point 410°F. Almost insoluble in water; soluble in most organic solvents. Combustible. Low toxicity.

triethylene glycol dipropionate

$C_{18}H_{34}O_6$; $CO(OCH_2CH_2)_3OOCOC_3H_7$.
Properties: Colorless liquid. Sp. gr. (25°C) 1.066; refractive index (25°C) 1.436; b.p. (2 mm) 138–142°C; f.p. less than -60°C; solubility in water, 6.70% by weight. Combustible. Low toxicity.
Use: Plasticizer.

triethylene glycol monobutyl ether. See butoxytriethylene glycol.

triethylenemelamine (tretamine; TEM; 2,4,6-tris(1-aziridinyl)-s-triazine)



Properties: White, crystalline, odorless powder; m.p. 160°C (polymers); polymers readily with heat or moisture; soluble in alcohol, water, methanol, chloroform, and acetone.

Grade: N.F.

Hazard: Highly toxic.

Uses: Medicine (see nitrogen mustards); insecticide; chemosterilant.

triethylenephosphoramidate (tepa; tris-(1-aziridinyl)phosphine oxide; APO) $(NCH_2CH_2)_3PO$

Properties: Colorless crystals; m.p. 41°C; soluble in water, alcohol and ether. Combustible.

Derivation: From ethylenimine.

Hazard: Highly toxic. Strong irritant to skin and tissue.

Uses: Medicine (see nitrogen mustards); insect sterilant. Also used with tetrakis(hydroxymethyl)phosphonium chloride (THPC) to form a condensation polymer suitable for flameproofing cotton. See also tris[1-(2-methyl)aziridinyl]phosphine oxide.

Shipping regulations: (Rail) White label. (Air) Corrosive label. Legal label name: tris(1-aziridinyl)phosphine oxide.

See also tepa.

triethylenetetramine $NH_2(C_2H_4NH)_3C_2H_4NH_2$

Properties: Moderately viscous yellowish liquid. Less volatile than diethylenetriamine but resembles it in many other properties. Soluble in water. B.p. 277.5°C; sp. gr. 0.9818 (20/20°C); m.p. 12°C; flash point 275°F (C.C.); wt 8.2 lb/gal (20°C). Combustible. Autoignition temp. 640°F.

Grades: Technical; anhydrous.

Containers: Cans; drums; tank cars.

Hazard: Strong irritant to tissue. Causes skin burns and eye damage.

Uses: Detergents and softening agents; synthesis of dyes, pharmaceuticals and rubber accelerators.

Shipping regulations: (Air) Corrosive label.

tri(2-ethylhexyl)phosphate $[C_8H_{17}CH(C_2H_5)CH_2]_3PO_4$

Properties: Light-colored liquid; sp. gr. 0.9260 (20/20°C); 7.70 lb/gal (20°C); b.p. 220°C (5 mm); vapor pressure 1.9 mm Hg (200°C); insoluble in water; viscosity 14.1 cp (20°C); pour point -74°C; flash point 405°F. Combustible; low toxicity.
Use: Plasticizer.

tri(2-ethylhexyl) phosphite $(C_8H_{17}O)_3P$

Properties: Straw-colored liquid; sp. gr. 0.897 (25/15°C); m.p., glass at low temperature; refractive index 1.451 (n_D 25/D); flash point 340°F (COC). Combustible; low toxicity.
Uses: Plasticizer; intermediate.

tri(2-ethylhexyl) trimellitate $C_8H_{17}(COOC_6H_4)_3$

Properties: Clear liquid, mild odor; sp. gr. 0.992 (20/

20°C); distillation range at 3 mm, 278–284°C (5–95%); f.p., a gel at -35°C; refractive index 1.4846 (23°C); wt/gal 8.26 lb (20°C). Combustible; low toxicity.
Use: Plasticizer.

triethylmethane. See 3-ethylpentane.

triethylmethyl malonaldehyde diacetal. See 1,1,3-triethoxy-3-methoxypropane.

triethyl orthoformate (triethoxymethane) $CH(OC_2H_5)_3$

Properties: Colorless liquid; pungent odor; b.p. 145.9°C; refractive index 1.39218 (18.8°C); sp. gr. 0.895 (20/20°C). Soluble in alcohol, ether; decomp. in water. Flash point 86°F (C.C.). Low toxicity.

Derivation: Reaction of sodium ethylate with chloroform or reaction of hydrochloric acid with hydrogen cyanide in ethyl alcohol solution.

Containers: 55-gal steel drums.

Hazard: Flammable, moderate fire risk.

Uses: Organic synthesis; pharmaceuticals.

triethyl phosphate (TEP) $(C_2H_5)_3PO_4$

Properties: Colorless, high-boiling liquid. Mild odor; very stable at ordinary temperatures. Compatible with many gums and resins. Soluble in most organic solvents; completely miscible in water. When mixed with water is quite stable at room temperature, but at elevated temperatures it hydrolyzes slowly. F.p. -56.4°C; b.p. 216°C. Flash point 240°F; refractive index 1.4055 (20°C); wt/gal 8.90 lb (68°F). Combustible.

Grades: Technical; 97%.

Containers: Drums; tank cars; tank trucks.

Hazard: May cause nerve damage but to less extent than other cholinesterase-inhibiting compounds.

Uses: Solvent; plasticizer for resins, plastics, gums; manufacture of pesticides; catalyst; lacquer remover.

Shipping regulations: (Rail, Air) Organic phosphate, liquid, n.o.s., Poison label. Not acceptable on passenger planes.

triethyl phosphite $(C_2H_5)_3PO_3$

Properties: Colorless liquid; sp. gr. 0.9687 (20°C); b.p. 156.6°C; refractive index 1.413 (n_D 25/D); flash point 130°F; insoluble in water; soluble in alcohol and ether. Combustible.

Containers: Glass bottles; 5-, 55-gal drums.

Hazard: Moderate fire risk. May be toxic.

Uses: Synthesis; plasticizers; stabilizers; lube and grease additives.

O,O,O-triethyl phosphorothioate (triethyl thiophosphate) $(C_2H_5O)_3PS$

Properties: Colorless liquid with characteristic odor. B.p. (10 mm) 93.5–94°C; sp. gr. 1.074; flash point 225°F (COC); combustible.

Containers: Bottles; drums.

Hazard: Toxic by ingestion; cholinesterase inhibitor.

Uses: Plasticizer; lubricant additive; antifoam agent; hydraulic fluid; intermediate.

Shipping regulations: (Rail, Air) Organic phosphate, liquid, n.o.s., Poison label. Not acceptable on passenger planes.

triethyl tricarballylate $(C_2H_5OCOCCH_2)_3CHCOOC_3H_7$

Properties: Colorless liquid. Sp. gr. (20°C) 1.087; refractive index (26°C) 1.4234; b.p. (5 mm) 158–160°C; solubility in water (20°C) 0.62% by weight. Combustible.

Use: Plasticizer.

er intermediate; sol- parts and precision

thyl)trimethylammo- OH]; C₂H₅O- on); drums.

chloroethylene (q.v.).

obaltic oxide.

tricyclamol chloride

soluble in alcohol; in-) (48°C); b.p. 234°C ble.

ers.

OOH. A saturated in natural fats or oils. crystalline solid; m.p. d in medical research as chromatography.

C₂H₅)₂BO₂.

sp. gr. 1.065 (25°C); x 1.5480 (24°C); flash in all proportions in hydrolyzes on contact

sis.

hate; TCP)

f isomers. odorless liquid. Sta- refractive index 1.556); wt/gal 9.7 lb; crys- Miscible with all the

s, also with vegetable point 437°F; autoigni-

osphorus oxychloride.

; tank cars. ingestion and skin ab- highly toxic; its toler- of air.

chloride, polystyrene. for plastics; air filter waterproofing; additive s; hydraulic fluid and

P. slight phenolic odor. (20/4°C) 1.115; flash ble in water; miscible e, ether, and kerosine.

for plastics and resins.

der; faint characteristic odor; soluble in alcohol and in water. Use: Medicine.

tricyclic. An organic compound comprised of three (only) ring structures, which may be the same or different, e.g., anthracene.

sym-tricyclodecane. See adamantane.

tricyclohexyl borate. See boric acid ester.

n-tridecane CH₃(CH₂)₁₁CH₃.

Properties: Colorless liquid. Soluble in alcohol; insoluble in water. Sp. gr. 0.755 (20/4°C); b.p. 225.5°C; f.p. -5.45°C; refractive index 1.4250 (20/D); flash point 175°F. Combustible; low toxicity. Grades: 95%; 99%; research.

Containers: Glass bottles; 1-, 5-gal drums.

Uses: Organic synthesis; distillation chaser.

n-tridecanoic acid (tridecyclic acid; tridecoic acid) CH₃(CH₂)₁₁COOH. A saturated fatty acid usually prepared synthetically.

Properties: Colorless crystals; m.p. 44.5°C; sp. gr. 0.8458 (80/4°C); b.p. 312.4°C, 192.2°C (16 mm); refractive index 1.4328 (50°C). Slightly soluble in water; soluble in alcohol and ether. Combustible; low toxicity.

Grade: 99% pure.

Uses: Organic synthesis; medical research.

tridecanol. See tridecyl alcohol.

tridecoic acid. See n-tridecanoic acid.

tridecyl alcohol (tridecanol). A commercial mixture of isomers of the formula C₁₂H₂₅CH₂OH.

Properties: Low-melting white solid with pleasant odor; b.p. 274°C; m.p. 31°C; sp. gr. (20/20°C) 0.845; wt/gal 7.0 lb; flash point (TOC) 180°F. Combustible; low toxicity.

Derivation: Oxo process (q.v.) from C₁₁ hydrocarbons. Grade: Technical.

Containers: 55-gal drums; tank cars.

Uses: Esters for synthetic lubricants; detergents; anti-foam agent; other tridecyl compounds; perfumery.

tridecylbenzene (1-phenyltridecane) C₆H₅(CH₂)₁₂CH₃.

Properties: Colorless liquid; sp. gr. 0.85-0.86 (60/60°F); refractive index 1.4815-1.4830. Combustible. Use: Detergent intermediate.

tridecyclic acid. See n-tridecanoic acid.

tri(decyl) orthoformate CH(OC₁₀H₂₁)₃.

Properties: Liquid; b.p. 194°C; f.p. -15 to -20°C; refractive index 1.448; insoluble in water; soluble in benzene, naphtha, ether, and alcohol.

Use: To remove small quantities of water from ethers or other solvents where acid catalysts can be employed.

tri(decyl) phosphite (C₁₀H₂₁O)₃P.

Properties: Water-white liquid; decyl alcohol odor; sp. gr. 0.892 (25/15.5°C); m.p. less than 0°C; refractive index 1.4565 (25°C). Flash point 455°F. Combustible.

Containers: 55-gal drums.

Uses: Chemical intermediate; stabilizer for polyvinyl and polyolefin resins.

tridihexethyl chloride

C₆H₁₃C(C₆H₁₃)₂(OH)CH₂CH₂N(C₂H₅)₂C₂H₅Cl. (3-Diethylamino)hexadecyl hexadecylaziridone. etho-

form. and in alcohol. Practically insoluble in ether and in acetone. Melting range 198-202°C.

Grade: N.F.

Use: Medicine.

2,4,6-tri(dimethylaminomethyl)phenol

[(CH₃)₂NCH₂]₃C₆H₂OH.

Properties: Liquid; refractive index 1.5181. Combustible.

Hazard: May be toxic.

Uses: Antioxidants, acid neutralizers, stabilizers, and catalysts for epoxy and polyurethane resins.

tri(dimethylphenyl)phosphite (trixylenyl phosphate) [(CH₃)₂C₆H₄O]₃PO.

Properties: Liquid. Sp. gr. 1.155; refractive index 1.5535; b.p. (10 mm), 243-265°C; flash point 450°F; solubility in water (85°C), 0.002% by weight. Combustible.

Use: Plasticizer.

"Tridione."¹ Trademark for trimethadione, U.S.P.

tridodecyl amine. See tri-lauryl amine.

tridodecyl borate. See boric acid ester.

tridymite SiO₂. A vitreous, colorless or white, native form of pure silica. Found variously but not so commonly as quartz (q.v.). Quartz will change into tridymite with a 16.2% increase in volume at 870°C. Unlike quartz, it is soluble in boiling sodium carbonate solution. Sp. gr. 2.28-2.3; Mohs hardness 7.

trietazine. Generic name for 2-chloro-4-diethylamino-6-ethylamino-s-triazine ClC₂N₃[N(C₂H₅)₂]NHC₂H₅.

Properties: Solid; practically insoluble in water.

Uses: Herbicide; plant growth regulator.

"Tri-Ethane."¹⁷⁷ Trademark for 1,1,1-trichloroethane (q.v.).

triethanolamine (TEA; tri(2-hydroxyethyl)amine) (HOCH₂CH₂)₃N.

Properties: Colorless, viscous, hygroscopic liquid with slight ammoniacal odor; m.p. 21.2°C; b.p. 335°C (dec); vapor pressure <0.01 mm (20°C); sp. gr. 1.126; flash point (open cup) 375°F; wt/gal 9.4 lb; miscible with water, alcohol; soluble in chloroform; slightly soluble in benzene and ether; slightly less alkaline than ammonia. Commercial product contains up to 25% diethanolamine and up to 5% monoethanolamine. Combustible; low toxicity.

Derivation: Reaction of ethylene oxide and ammonia. Grades: Technical; regular; 98%; U.S.P.

Containers: Drums; tank cars.

Uses: Fatty acid soaps used in drycleaning, cosmetics, household detergents, and emulsions; wool scouring; textile antifume agent and water-repellent; dispersion agent; corrosion inhibitor; softening agent, humectant, and plasticizer; insecticide; chelating agent; rubber accelerator.

triethanolamine lauryl sulfate

(HOC₁₂H₂₅)₂NOS(O₂)OC₁₂H₂₅. A liquid or paste.

Containers: Drums, tank cars; tank trucks.

Uses: Detergent; wetting, foaming and dispersing agent for industrial, cosmetic and pharmaceutical applications, especially shampoos.

triethanolamine methanearsonate

CH₃As(O)[ONH(C₂H₅)₂]₂.

Hazard: Highly toxic by ingestion.

Use: Herbicide.

Shipping regulations: (Rail, Air) Arsenical compounds.

triethanolamine oleate. See trihydroxyethylamine oleate.

triethanolamine stearate. See trihydroxyethylamine stearate.

triethanolamine titanate. See titanium chelate.

1,1,3-triethoxyhexane

CH(OC₂H₅)₂CH₂CH(OC₂H₅)C₂H₅.

Properties: Colorless liquid; sp. gr. 0.8746 (20/20°C); b.p. 133°C (50 mm); f.p. -100°C; wt/gal 7.3 lb; flash point 210°F. Insoluble in water. Combustible. Low toxicity.

Use: Synthesis of aldehydes, acids, esters, chloride, amines, etc.

triethoxymethane. See triethyl orthoformate.

1,1,3-triethoxy-3-methoxypropane (triethylmethyl malonaldehyde diacetal)

(CH₃O)(C₂H₅O)CHCH₂CH(OC₂H₅)₂.

Properties: Colorless liquid. Sp. gr. (25/4°C), 0.9300; b.p. (6 mm) 86°C. Combustible.

Grade: 99%.

Uses: Intermediate; crosslinking and insolubilizing agent.

triethylaconitate

C₂H₅OOCCHC(COOC₂H₅)₂CH₂COOC₂H₅.

Properties: Liquid. Sp. gr. (25°C) 1.096; refractive index (26°C) 1.4517; b.p. (5 mm) 154-156°C. Combustible.

Use: Plasticizer.

triethylaluminum (ATE; TEA; aluminum triethyl). (C₂H₅)₃Al.

Properties: Colorless liquid; sp. gr. 0.837; f.p. -52.5°C; b.p. 194°C; specific heat 0.527 (91.4°F). Miscible with saturated hydrocarbons. Flash point -63°F.

Derivation: By introduction of ethylene and hydrogen into an autoclave containing aluminum. The reaction proceeds under moderate temperature and varying pressures.

Grades: 88-94%.

Containers: Cylinders.

Hazard: Highly toxic; destructive to tissue. Flammable, dangerous fire risk; ignites spontaneously in air; reacts violently with water, acids, alcohols, halogens, and amines.

Uses: Catalyst intermediate for polymerization of olefins, especially ethylene; pyrophoric fuels; production of alpha-olefins and long chain alcohols; gas plating of aluminum.

Shipping regulations: (Rail) Red label. (Air) Not acceptable. (Rail) Legal label name; aluminum triethyl.

triethylamine (C₂H₅)₃N.

Properties: Colorless liquid; strong ammoniacal odor. B.p. 89.7°C; f.p. -115.3°C; sp. gr. (20/20°C) 0.7293; wt/gal (20°C) 6.1 lb; flash point (open cup) 20°F. Soluble in water and alcohol.

Derivation: From ethyl chloride and ammonia under heat and pressure.

Containers: 1-, 5-gal cans; 55-gal drums; tank cars.

Hazard: Flammable, dangerous fire risk. Explosive limits in air 1:2 to 8.0%. Tolerance, 25 ppm in air. Toxic by ingestion and inhalation; strong irritant to tissue.

Uses: Catalytic solvent in chemical synthesis; accelerator activators for rubber; wetting, penetrating and

P-AH TP-90B - SPGR, 0.97. High boiling, straw colored liquid; little or no odor; non-toxic in ordinary handling. Visc (27°C) 8 cps. Bp 660-760°F.

Volatility of plasticizers

TABLE 15.3

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers</i>												
1	Acetin	C ₈ H ₁₀ O ₄	134	1.190 at 25° C.	1.445 at 25° C.	66.8 at 25° C.						
2	o- and p-Toluenesulfonamides (mixture)	C ₇ H ₈ O ₂ NS	171	1.353 solid		f.p. 105						
3	Diacetin	C ₇ H ₁₂ O ₆	176	1.178	1.438 at 25° C.	35.7 at 25° C.						
4	<i>N</i> -Ethyl- <i>p</i> -toluenesulfonamide	C ₉ H ₁₂ O ₂ NS	199	1.171 at 65° C.	1.522 at 65° C.	59.0 f.p.						
5	<i>N</i> -Ethyl- <i>o</i> - and <i>p</i> -toluenesulfonamides (mixture)	C ₉ H ₁₂ O ₂ NS	199	1.190 at 25° C.	1.540 at 25° C.	426				0.55	101	
6	Triacetin	C ₉ H ₁₄ O ₆	218	1.157 (av.) at 25° C.	1.429 at 25° C.	15.1 at 25° C.						
7	Dimethyl phthalate	C ₁₀ H ₁₀ O ₄	194	1.191 at 25° C.	1.513 at 25° C.	16.3 at 25° C.			0.5	0.34	>4.0	>1000
8	o-Nitrobiphenyl	C ₁₂ H ₉ O ₂ N	199	1.203 at 25° C.	1.613 at 25° C.	38 at 25° C.						
9	Diethyl phthalate	C ₁₂ H ₁₄ O ₄	222	1.118 at 25° C.	1.501 at 25° C.	12.6 at 25° C.	0.15	0.7	2.8	1.1	3.40	> 500
10	Triethyl citrate	C ₁₂ H ₂₀ O ₇	276	1.136 at 25° C.	1.441 at 25° C.							
11	Dimethyl sebacate	C ₁₂ H ₂₂ O ₄	230	0.986 at 30° C.	1.436 at 30° C.	3.5 at 38° C.						

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No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^b		
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.p.t.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Bristle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spex, 2 days, 60° C., 100% Humidity ^b	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
<i>Simple Plasticizers</i>														
1	Acetin												Kes	
2	o- and p-Toluenesulfonamides (mixture)												Mon	
3	Diacetin												Kes	
4	<i>N</i> -Ethyl- <i>p</i> -toluenesulfonamide												Mon	
5	<i>N</i> -Ethyl- <i>o</i> - and <i>p</i> -toluenesulfonamides (mixture)												Mon	
6	Triacetin												Kes, TE	
7	Dimethyl phthalate												Mon, TE	
8	o-Nitrobiphenyl												Mon	
9	Diethyl phthalate												Mon, TE, UBI	
10	Triethyl citrate												PI	
11	Dimethyl sebacate												Har	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.

DESCRIPTION OF TABLES AND FIGURES

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (131° C.)
Simple Plasticizers (Continued)												
12	Di-n-butyl tartrate	C ₁₂ H ₂₂ O ₈	262	1.094	1.444	67.7	0.5	0.7	0.3	2.4	1.50	143
13	Tri-n-butyl phosphate	C ₁₂ H ₂₇ O ₄ P	266	0.978	1.424	at 25° C. 3.4	<0.07	6.0	M	M	>4	50B
14	Monomethyl phthalate ethyl glycolate ester	C ₁₂ H ₁₄ O ₆	266	1.227	1.504	at 20° C. 185	0.1	1.2	0.5	0.1	0.65	61
15	N-Cyclohexyl-p-toluenesulfonamide	C ₁₂ H ₁₉ O ₂ NS	253	1.125	1.498	solid at 25° C. 87° C.	<0.08	1.38	1.60	0.18	0.50	93
16	Monomethyl phthalate ethyl glycolate ester	C ₁₂ H ₁₆ O ₆	280	1.186	1.501	at 25° C. 53	0.55	3.2	0.50	0.10	0.18	62
17	Di(methyl Cellosolve) phthalate	C ₁₂ H ₁₈ O ₆	282	1.160	1.439	at 25° C. 70						
18	Triethyl acetylacrylate	C ₁₂ H ₂₂ O ₆	318	1.13	1.439	at 25° C. 8.9						> 525
19	Diisobutyl adipate	C ₁₂ H ₂₄ O ₄	258	0.967	1.429	at 25° C. 8.9						
20	Tributyrin	C ₁₂ H ₂₄ O ₆	302	1.036	1.433	at 25° C. 7.5						
21	Methyl Cellosolve laurate	C ₁₂ H ₂₄ O ₆	258	0.894	1.440	at 25° C.						

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ²		
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spew, 8 days, 60° C., 100% Humidity ³	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
Simple Plasticizers (Continued)														
12	Di-n-butyl tartrate												Ke	
13	Tri-n-butyl phosphate												CS	
14	Monomethyl phthalate ethyl glycolate ester	55.4 ^c						17.5		6.1	5.9		Mon	
15	N-Cyclohexyl-p-toluenesulfonamide												Mon	
16	Monomethyl phthalate ethyl glycolate ester												Mon	
17	Di(methyl Cellosolve) phthalate	46 ^c						6.6		4.6	3.7	None	OAJTE	
18	Triethyl acetylacrylate												Pr	
19	Diisobutyl adipate												Ca	
20	Tributyrin												TE	
21	Methyl Cellosolve laurate												Ke	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.
^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁸
² For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
Simple Plasticizers (Continued)												
22	Di-n-butyl phthalate	C ₁₈ H ₂₂ O ₄	278	1.048	1.483 at 20° C.	20.3	Insol.	0.46	34	30.0	0.08	168
23	Di(Cellosolve) phthalate	C ₁₄ H ₂₂ O ₆	310	1.120	1.491 at 25° C.	f.p. 31° C.	0.2	2.0	2	0.64	0.15	30
24	n-Butyl laurate	C ₁₈ H ₃₄ O ₂	256	0.857 at 25° C.								
25	Methyl Cellosolve myristate	C ₁₇ H ₃₄ O ₂	286	0.895 at 25° C.								
26	Di(butyl CARBITOL) formal	C ₁₇ H ₃₄ O ₆	336	0.97								
27	Triphenyl phosphate	C ₁₈ H ₁₅ O ₄ P	326	1.268 at 60° C.	1.552 at 60° C.	f.p. 49.9° C.					0.02	23
28	Bis(dimethylbenzyl) ether	C ₁₃ H ₂₂ O	206	1.008	1.553 at 25° C.	29.0						
29	n-Butyl cyclohexyl phthalate	C ₁₈ H ₂₄ O ₄	304	1.078	1.508 at 25° C.	117.0						75
30	Mono-n-butyl phthalate n-butyl glycolate ester	C ₁₈ H ₂₄ O ₆	336	1.103	1.490 at 25° C.	64.9	Insol.	0.37	4.65	2.70	0.08	31
31	Tri-n-butyl citrate	C ₁₈ H ₃₂ O ₇	360	1.045	1.443 at 25° C.		Insol.	<0.2	10.0	14.5	0.16	44
32	Triethylene glycol di(2-ethylbutyrate)	C ₁₈ H ₃₄ O ₆	346	0.995	1.440 at 20° C.	11.5	0.02	1.0	0.2	24.0	0.86	

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^b		
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Shore, 2 days, 60° C., 100% Humidity ^c	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
Simple Plasticizers (Continued)														
22	Di-n-butyl phthalate	41.4°						17.6		2.7	6.3	None	Bar, CR, Har, Mon, Nov, Pitt, TK, URI, DA	
23	Di(Cellosolve) phthalate												OA	
24	n-Butyl laurate												AH	
25	Methyl Cellosolve myristate												AH	
26	Di(butyl CARBITOL) formal	55.3°						10.1		15.6	14.5	None	Thi	
27	Triphenyl phosphate												Dow, Mon, Pitt	
28	Bis(dimethylbenzyl) ether												OA	
29	n-Butyl cyclohexyl phthalate	58.8				+3	-20	6.5		0.6	8.8	None	Bar	
30	Mono-n-butyl phthalate n-butyl glycolate ester	58.2°	3000	300				3.3		17	4.3	None	Mon	
31	Tri-n-butyl citrate	55.1	2800	300		-4	-23	3.1		6.1	14.6	None	Pt	
32	Triethylene glycol di(2-ethylbutyrate)												CCCC	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁶^d For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
Simple Plasticizers (Continued)												
33	Butyl CELLOSOLVS laurate	C ₁₈ H ₃₄ O ₂	300	0.882 at 25° C. (av.)	1.439 at 25° C.	6.5 at 25° C.						
34	Di-n-hexyl adipate	C ₁₈ H ₃₄ O ₄	314	0.933 at 25° C.	1.439 at 25° C.	7.8 at 25° C.						
35	Di(1,3-dimethylbutyl) adipate	C ₁₈ H ₃₄ O ₄	314	0.926 at 25° C.	1.433 at 25° C.	5.6 at 38° C.						
36	Di-n-butyl sebacate	C ₁₈ H ₃₄ O ₄	314	0.936 at 25° C.	1.440 at 25° C.	9.3 at 25° C.	Insol.	1.6	M	M		63
37	Di(butyl CELLOSOLVS) adipate	C ₁₈ H ₃₄ O ₄	348	0.997 at 25° C.	1.442 at 25° C.	12.5						58
38	n-Butyl myristate	C ₁₈ H ₃₆ O ₂	284	0.861 at 25° C.	1.434 at 25° C.							
39	Tri(butyl CELLOSOLVS) phosphate	C ₁₈ H ₃₆ O ₇ P	398	1.023 at 25° C.	1.434 at 25° C.	12.3	0.08	6.3	Almost miscible			
40	Cresyl diphenyl phosphate	C ₁₉ H ₁₇ O ₄ P	340	1.208 at 25° C.	1.563 at 20° C.	39.8						
41	n-Butyl benzyl phthalate	C ₁₉ H ₂₀ O ₄	312	1.116 at 25° C.	1.538 at 20° C.	39.3 at 23° C.						29
42	Bis(dimethylbenzyl) carbonate	C ₁₉ H ₂₂ O ₃	298	1.063 at 25° C.	1.547 at 20° C.	155						32

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a											Supplier ^m	
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extraction, 10 days, 23° C., 4-mil Film			Spew, 2 days, 60° C., 100% Humidity ^b
									10 days, 60° C. ^c	24 hours, 70° C.	Water	Oil		
Simple Plasticizers (Continued)														
33	Butyl CELLOSOLVS laurate													Kes
34	Di-n-hexyl adipate													Kes
35	Di(1,3-dimethylbutyl) adipate													Har
36	Di-n-butyl sebacate	37.8°						7.2		0.4	16.5	None		Har, Pitt, RH
37	Di(butyl CELLOSOLVS) adipate	41.4°									18.0	None		Kes, OA
38	n-Butyl myristate													AH
39	Tri(butyl CELLOSOLVS) phosphate	43.7°						1.1		14.2	13.5	None		OA
40	Cresyl diphenyl phosphate	72	3000	292	64	475	4	-15		6.1	0.9	14.8	None	Mon
41	n-Butyl benzyl phthalate	59	2930	281	66	730	6	-11		9.5	0.6	7.8	None	Mon
42	Bis(dimethylbenzyl) carbonate	71	3000	284	63	450	2	-9		11.8	1.5	9.2		OA

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁸^m See Table 15.1 for supplier abbreviations; see page 913.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICISERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (135° C.)
<i>Simple Plasticizers (Continued)</i>												
43	Methyl pentachlorostearate	C ₁₉ H ₃₃ O ₂ Cl ₅	471	1.204	1.497 at 20° C. 1.462 at 25° C.	975.0						7
44	Methyl ricinoleate	C ₁₉ H ₃₅ O ₂	312	0.925 at 25° C.	1.462 at 25° C.							
45	Methyl <i>CALLIOCOLVA</i> palmitate	C ₁₉ H ₃₅ O ₂	314	0.891 at 25° C.								
46	Methyl hydroxystearate	C ₁₉ H ₃₅ O ₃	314									
47	Diphenyl phthalate	C ₂₀ H ₁₄ O ₄	318	1.29 solid	1.572 at 74° C.						0.006	19
48	Dicyclohexyl phthalate	C ₂₀ H ₃₀ O ₄	330	1.28								
49	2-Ethylhexyl diphenyl phosphate	C ₂₀ H ₃₄ O ₄ P	362	1.062	1.511 at 20° C. 1.487 at 20° C.	29.5	Insol.	0.34				44
50	Di-n-hexyl phthalate	C ₂₀ H ₃₈ O ₄	334	1.007	1.487 at 20° C. 1.468 at 25° C.	56	Insol.	<0.06	M	M	0.20	66
51	Di(2-ethylbutyl) phthalate	C ₂₀ H ₃₈ O ₄	334	0.965	1.480 at 25° C.	28.8 cm. at 25° C.						> 180

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-3 Specimens Plasticized to Equivalence*											Supplier ^b	
		Plasticizer p.p.t. at 25° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 25° C.	Ultimate Elongation, % at 25° C.	Shore Durorometer A at 25° C.	Stiffness at 25° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 25° C., 4-mil Film			Spew, 3 days, 60° C., 100% Humidity ^c
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Simple Plasticizers (Continued)</i>														
43	Methyl pentachlorostearate	69 ^d	2900	338	64	600	- 6	- 19	0.2		0.2	17.1	None	Hook
44	Methyl ricinoleate													Bak
45	Methyl <i>CALLIOCOLVA</i> palmitate													AH
46	Methyl hydroxystearate													Bak
47	Diphenyl phthalate													Mon
48	Dicyclohexyl phthalate	64 ^e	3000	390			+ 8	- 5	1.0		0	4.5	None	Bur, OA
49	2-Ethylhexyl diphenyl phosphate	60	2580	260	64	540	- 2	- 28	1.6		0.2	14.5	None	Mon
50	Di-n-hexyl phthalate	57	2720	293	63	570	- 8	- 31	2.8	11.0	0	12.5	None	CCCC, TE
51	Di(2-ethylbutyl) phthalate	48.5 ^f							4.4		0.2	8.4	None	TE
52	Di(1,3-dimethylbutyl) phthalate													Has

* 1000 p.s.i. tensile modulus at 75% elongation at 25° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.

^b Dash (—) indicates no data available.

^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Road 24.

^d Mixture of 1 part plasticizer to 3 parts PLAZOL plasticizer DOP.

^e Mixture of 0.6 part plasticizer to 1 part PLAZOL plasticizer DOP.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	312° F. (160° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
53	Di(butyl CELLULOSE) phthalate	C ₂₀ H ₃₀ O ₆	366	1.063	1.483 at 25° C.	42.0	Insol.	0.63	2.80	6.50	0.06	24
54	Di(CANAROL) phthalate	C ₂₀ H ₃₀ O ₆	398	1.150	1.492 at 25° C.	82.0	Insol.	2.80	0.09	0.30	0.08	18
55	Tri-n-butyl acetylacrylate	C ₂₀ H ₃₄ O ₆	402	1.046 at 25° C.	1.441 at 25° C.							61
56	n-Butyl palmitate	C ₂₀ H ₄₀ O ₂	313	0.866 at 25° C.								
57	Tricresyl phosphate	C ₂₁ H ₃₁ O ₄ P	368	1.166	1.557 at 20° C.	120.0	Insol.	0.87	5.5	2.0	0.01	7
58	n-Butyl benzyl sebacate	C ₂₁ H ₃₈ O ₄	348	1.004	1.478 at 25° C.	8.6 cm. at 35° C.						
59	Methyl acetylacrylate	C ₂₁ H ₃₀ O ₄	356	0.988	1.456 at 20° C.						0.12	29
60	Methyl CELLULOSE oleate	C ₂₁ H ₄₀ O ₂	341	0.902	1.453 at 25° C.	10.3						
61	Di(2-ethylbutyl) azelate	C ₂₁ H ₄₀ O ₄	357	0.930	1.442 at 25° C.	11.4						48
62	Propylene glycol monoricinoleate	C ₂₁ H ₄₀ O ₄	357	0.950 at 25° C.								
63	Methyl CELLULOSE stearate	C ₂₁ H ₄₂ O ₂	343	0.877 at 25° C.	1.443 at 25° C.	8.9 at 25° C.						30

No.	Name	Comparison of Properties of VinyLite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^b			
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Softness at 23° C., p.s.i. ^c	T _g , °C.	Brittle Temperature, °C.		% Loss, 4-mil Film			% Extrac-tion, 10 days, 23° C., 4-mil Film		Spew, 3 days, 60° C., 100% Humidity
								10 days, 60° C.	24 hours, 70° C.	Water	Oil		Water	Oil	
<i>Simple Plasticizers (Continued)</i>															
53	Di(butyl CELLULOSE) phthalate	47.8 ^d								2.1		4.8	11.4	None	QA
54	Di(CANAROL) phthalate	54.4 ^d								2.0		10.1	9.9	None	QA
55	Tri-n-butyl acetylacrylate														PT
56	n-Butyl palmitate														AH
57	Tricresyl phosphate	75	2930	274	58	580	+ 5	- 3		0.1		0.7	6.1	None	Cal, Mon, QA, Pitt
58	n-Butyl benzyl sebacate														Har
59	Methyl acetylacrylate	47.8 ^d								2.8		9.5	11.0		Bak
60	Methyl CELLULOSE oleate	50.5 ^d								2.6		2.0	10.3		AH, Kan, QA
61	Di(2-ethylbutyl) azelate	44.5	2800	230						5.3		1.5	12.0	None	Rm
62	Propylene glycol monoricinoleate														Bak
63	Methyl CELLULOSE stearate														AH, Kan

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.
^b Dash (—) indicates no data available.
^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.²⁰
^d Mixture of 0.63 part plasticizer to 1 part tricresyl phosphate.
^e For key to supplier abbreviations see page 902.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (135° C.)
Simple Plasticizers (Continued)												
64	n-Butyl ricinoleate	C ₃₂ H ₆₂ O ₈	355									
65	Di(1,3-dimethylbutyl) sebacate	C ₂₂ H ₄₂ O ₄	371	0.911	1.439 at 25° C.	9.3 ca. at 25° C.						
66	Di(2-ethylhexyl) adipate	C ₂₂ H ₄₂ O ₄	371	0.927	1.447 at 20° C.	12.7 at 20° C.	Insol.	0.12				41
67	Diisooctyl adipate	C ₂₂ H ₄₂ O ₄	371	0.928	1.448 at 25° C.	17.7 at 25° C.						23
68	Dicapryl adipate	C ₂₂ H ₄₂ O ₄	371	0.915	1.440 at 25° C.	8.3 ca. at 25° C.						
69	Diethylene glycol dipelargonate	C ₂₂ H ₄₂ O ₈	387	0.968	1.447 at 20° C.	19 at 20° C.						
70	Di(butyl CALLOSOOLS) sebacate	C ₂₂ H ₄₂ O ₄	408	0.970							0.04	
71	Triethylene glycol di(2-ethyl hexanoate)	C ₂₂ H ₄₂ O ₆	408	0.968	1.444 at 20° C.	16.1	Insol.	0.4	M	M	0.18	50
72	Triethylene glycol dicaprylate	C ₂₂ H ₄₂ O ₆	406 (av.)	0.972								
73	Di(butyl CARARROL) adipate	C ₂₂ H ₄₂ O ₄	426	1.03 at 18° C.		15 at 22° C.						
74	n-Butyl stearate	C ₂₂ H ₄₄ O ₂	341	0.858	1.444 at 20° C.	7.9 at 25° C.						29

M = Miscible.

PLASTICIZERS AND PLASTICIZATION

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence ^a										Supplier ^b		
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spec. 2 days, 60° C., 100% Humidity ^c	
									10 days, 80° C.	24 hours, 70° C.	Water			Oil
Simple Plasticizers (Continued)														
64	n-Butyl ricinoleate												Dee	
65	Di(1,3-dimethylbutyl) sebacate												Har	
66	Di(2-ethylhexyl) adipate	51	2730	318	67	600	-15	-53	3.5	12.2	2.2	21.0	None	CCCC, Gdr, MWI, Ken, OA, Har
67	Diisooctyl adipate													Ken, MWI, OA, Pitt, RCA, Har, Cab
68	Dicapryl adipate													Har
69	Diethylene glycol dipelargonate	51.3	2800	330			-15	-48	6.9		6.0	22.0	Yes	Em
70	Di(butyl CALLOSOOLS) sebacate													Dee
71	Triethylene glycol di(2-ethylhexanoate)	59	2680	300	66	800	-17	-50	4.7	12.4	2.0	22.8	None	CCCC
72	Triethylene glycol dicaprylate													RCA, Drew
73	Di(butyl CARARROL) adipate													Thi
74	n-Butyl stearate													CB, AH, Har, Ken, OA

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.

^b Dash (—) indicates no data available.

^c For key to supplier abbreviations see page 903.

DESCRIPTION OF TABLES AND FIGURES

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
Simple Plasticizers (Continued)												
75	n-Butyl oleate	C ₂₂ H ₄₂ O ₂	339	0.864 at 25° C.	1.449 at 25° C.	7.7 at 25° C.						
76	Tetrahydrofurfuryl oleate	C ₂₂ H ₄₂ O ₃	366	0.925 at 20° C.	1.468 at 20° C.	17.8 at 20° C.						
77	Methyl CELLULOSE acetyricinoleate	C ₂₂ H ₄₂ O ₆	399	0.965 at 20° C.	1.456 at 20° C.	42.1 at 20° C.	Insol.	0.6	M	M		12
78	o-Xenyl diphenyl phosphate	C ₂₄ H ₁₈ O ₄ P	402	1.233 at 30° C.	ca. 1.596 at 60° C.	30-55 at 30° C.	Insol.	0.26	2.22	0.3	0.006	20
79	Dibenzyl sebacate	C ₂₄ H ₃₈ O ₄	382	1.055 at 30° C.	1.520 at 30° C.	13.0 at 35° C.						
80	Di(2-ethylhexyl) phthalate	C ₂₄ H ₄₆ O ₄	390	0.980 at 20° C.	1.488 at 20° C.	81.0 at 20° C.	Insol.	0.20	M	M	0.02	24
81	Diisooctyl phthalate	C ₂₄ H ₄₆ O ₄	390	0.987 at 20° C.	1.488 at 20° C.	83.0 at 20° C.						10
82	Dicapryl phthalate	C ₂₄ H ₄₆ O ₄	390	0.966 at 20° C.	1.479 at 20° C.	66.7 at 20° C.						23
83	Di(2-ethylhexyl) tetrahydrophthalate	C ₂₄ H ₄₄ O ₄	396	0.969 at 20° C.	1.466 at 20° C.	42.0 at 20° C.	Insol.	0.26				30
84	Tetra-n-butyl thiodisuccinate	C ₂₄ H ₄₂ O ₄ S	491	1.064 at 20° C.	1.463 at 20° C.	63.6 at 20° C.	Insol.	0.23				6
85	Di(2-ethylhexyl) hexahydrophthalate	C ₂₄ H ₄₄ O ₄	397	0.966 at 20° C.	1.461 at 20° C.	35.9 at 20° C.						37
86	n-Butyl acetyricinoleate	C ₂₄ H ₄₄ O ₄	397	0.929 at 25° C.	1.456 at 25° C.	35.2 at 25° C.						20

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalence*										Supplier ^b			
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.p.t.	Tensile Strength, p.p.t. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 23° C., p.p.t.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spew, 2 days, 90° C., 100% Humidity		
									10 days, 60° C.	24 hours, 70° C.	Water			Oil	
75	n-Butyl oleate												AH, Kes		
76	Tetrahydrofurfuryl oleate	59 ^d	2590	297	67	630	-8	-34	0.8		0.3	17.2	None	AH, Em, Pitt	
77	Methyl CELLULOSE acetyricinoleate	50.8 ^d							2.4		1.2	14.5	None	Bak, Dee	
78	o-Xenyl diphenyl phosphate													Dow	
79	Dibenzyl sebacate	45.3 ^d							0.7		0.2	11.8	None	Har	
80	Di(2-ethylhexyl) phthalate	61	2360	253	67	700	-8	-31	0.7	5.1	0	15.5	None	Bar, CCCC, Har, Mon, MWI, OA, Pitt, TE, DA, Odr	
81	Diisooctyl phthalate	56	2700	300			-1	-37					None	Bar, Cab, Har, MWI, OA, Pitt, RCA	
82	Dicapryl phthalate	64	2500	270			-5	-35	0.6		0	23.8	None	RH, Har	
83	Di(2-ethylhexyl) tetrahydrophthalate	66	2590	320	68	560	-12	-37	4.1	11.5	0.3	21.8	None	CCCC	
84	Tetra-n-butyl thiodisuccinate	64	3200	318	62	760	-2	-23	0	2.5	0.5	15.2	None	CCCC	
85	Di(2-ethylhexyl) hexahydrophthalate	64	2580	300	68	750	-10	-34			9.7	0	25.4	None	CCCC
86	n-Butyl acetyricinoleate	55.4 ^d											Yes	Bak	

Simple Plasticizers (Continued)

75	n-Butyl oleate													AH, Kes	
76	Tetrahydrofurfuryl oleate	59 ^d	2590	297	67	630	-8	-34	0.8		0.3	17.2	None	AH, Em, Pitt	
77	Methyl CELLULOSE acetyricinoleate	50.8 ^d							2.4		1.2	14.5	None	Bak, Dee	
78	o-Xenyl diphenyl phosphate													Dow	
79	Dibenzyl sebacate	45.3 ^d							0.7		0.2	11.8	None	Har	
80	Di(2-ethylhexyl) phthalate	61	2360	253	67	700	-8	-31	0.7	5.1	0	15.5	None	Bar, CCCC, Har, Mon, MWI, OA, Pitt, TE, DA, Odr	
81	Diisooctyl phthalate	56	2700	300			-1	-37					None	Bar, Cab, Har, MWI, OA, Pitt, RCA	
82	Dicapryl phthalate	64	2500	270			-5	-35	0.6		0	23.8	None	RH, Har	
83	Di(2-ethylhexyl) tetrahydrophthalate	66	2590	320	68	560	-12	-37	4.1	11.5	0.3	21.8	None	CCCC	
84	Tetra-n-butyl thiodisuccinate	64	3200	318	62	760	-2	-23	0	2.5	0.5	15.2	None	CCCC	
85	Di(2-ethylhexyl) hexahydrophthalate	64	2580	300	68	750	-10	-34			9.7	0	25.4	None	CCCC
86	n-Butyl acetyricinoleate	55.4 ^d											Yes	Bak	

* 1000 p.p.t. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.p.t. per minute.

^b Dash (—) indicates no data available.^c 100% elongation, 1000 p.p.t., constant rate of loading of 810 p.p.t. per minute. See Recd.²⁵

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (130° C.)
Simple Plasticizers (Continued)												
87	Butyl CELLULOSE oleate	C ₂₄ H ₄₆ O ₂	383	0.885 at 25° C.	1.454 at 25° C.	5.5 at 25° C.						
88	Dinonyl adipate	C ₂₄ H ₄₆ O ₄	398	0.914 at 25° C.	1.445 at 25° C.	18.5 at 25° C.						
89	n-Octyl n-decyl adipate (mixture)	C ₂₄ H ₄₆ O ₄ (av.)	398	0.920 at 20° C.	1.450 at 20° C.	15.4 at 20° C.						
90	Polyethylene glycol di(2-ethylhexanoate)	C ₂₄ H ₄₆ O ₇	447	0.989 at 20° C.	1.447 at 20° C.	25.1 at 20° C.	Insol.	1.4				44
91	Butyl CELLULOSE stearate	C ₂₄ H ₄₈ O ₂	385	0.883 at 25° C.	1.446 at 25° C.	13.3 at 25° C.						
92	Tri(2-ethylhexyl) phosphate	C ₂₄ H ₄₈ O ₄ P	435	0.926 at 20° C.	1.443 at 20° C.	14.1 at 20° C.	Insol.	1.4	M	M		38
93	Di(2-ethylhexyl) sebacate	C ₂₄ H ₄₈ O ₄	413	0.918 at 20° C.	1.444 at 20° C.	20 at 20° C.						26
94	n-Octyl n-decyl phthalate (mixture)	C ₂₄ H ₄₄ O ₄ (av.)	418	0.975 at 20° C.	1.484 at 20° C.	72 at 20° C.						8
95	Isooctyl n-octyl n-decyl phthalate (mixture)	C ₂₄ H ₄₄ O ₄ (av.)	418	0.978 at 20° C.	1.485 at 20° C.	42.9 at 20° C.						
96	Di(2-ethylhexyl) sebacate	C ₂₄ H ₄₈ O ₄	427	0.913 at 20° C.	1.451 at 20° C.	19.9 at 20° C.						13
97	Dioctyl sebacate	C ₂₄ H ₄₈ O ₄	427	0.917 at 25° C.	1.447 at 25° C.	20.4 at 25° C.						
98	Dicapryl sebacate	C ₂₄ H ₄₈ O ₄	427	0.907 at 25° C.	1.444 at 25° C.	13.0 at 25° C.						

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-6 Specimens Plasticized to Equivalence ^a										Supplier ^b		
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Duroometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Long, 4-mil Film		% Extraction, 10 days, 23° C., 4-mil Film		Spew, 3 days, 60° C., 100% Humidity ^c	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
Simple Plasticizers (Continued)														
87	Butyl CELLULOSE oleate												Kas	
88	Dinonyl adipate												Pitt	
89	n-Octyl n-decyl adipate (mixture)	53	2550	295	78	1050	-7	-54	5.5	0.3	23.5	None	Here, OA	
90	Polyethylene glycol di(2-ethylhexanoate)	62	2690	308	70	750	-12	-44	3.8	8.8	3.0	19.4	None	CCCC
91	Butyl CELLULOSE stearate												AM, Kas, OA	
92	Tri(2-ethylhexyl) phosphate	59	2640	300	66	980	-16	-57	3.2	6.6	0.6	24.3	None	CCCC
93	Di(2-ethylhexyl) sebacate	57	2590	334	66	750	-17	-56	0.5		0.4	22.5	None	Em.
94	n-Octyl n-decyl phthalate (mixture)	63	2640	318	64	640	-8	-28	0.2		0.3	21.3	None	Here
95	Isooctyl n-octyl n-decyl phthalate (mixture)	67	2720	334	66	470	-12	-39		3.9	0	22.4	None	Here
96	Di(2-ethylhexyl) sebacate	60	2420	293	67	1700	-19	-56	0.08	4.6	0.3	20.2	None	Des, Har, MWI, RH
97	Dioctyl sebacate												Har, Pitt, MWI	
98	Dicapryl sebacate												Har	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.

^b Dash (—) indicates no data available.

^c For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Simple Plasticizers (Continued)</i>												
99	2-Ethylhexanoic acid diester of <i>N,N</i> -bis(2-hydroxyethyl)-2-ethylhexanamide	C ₂₈ H ₅₂ O ₆ N	484	0.956	1.458 at 20° C.	139.2	Insol.	0.5				12
100	Tri(<i>p</i> - <i>tert</i> -butylphenyl) phosphate	C ₂₀ H ₂₀ O ₄ P	495			f.p. 95-99.5° C.						0.9
101	Glyceryl triricinoleate	C ₅₇ H ₁₀₄ O ₉	933	0.959 at 25° C.	1.477 at 25° C.							0.7
102	Glyceryl tri(acetylricinoleate)	C ₆₃ H ₁₁₀ O ₁₃	1060	0.984	1.469 at 25° C.	271						
<i>Resinous Plasticizers</i>												
103	AROCLOX 1242, chlorinated biphenyl			1.378-1.388 at 25° C.	1.627-1.629 at 20° C.		Insol.	<0.1	M	M	2.50	146
104	FLAKOL plasticizer B-2H, polyester			1.055	1.469 at 20° C.	ca. 20,000	Insol.	1.0				0.5
105	G.E. 2557, polyester			1.028	1.459 at 20° C.	306						3
106	G.E. 2559, polyester			1.060	1.492 at 20° C.	4278						1
107	PARACREL C, nitrile rubber			0.984	1.470 at 20° C.	300,000						0.2
108	PARAPLAK G-25, polyester		8000	1.06								

M = Miscible.

No.	Name	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticized to Equivalance ^a											Supplier ^b	
		Plasticizer p.p.r. at 23° C., 75% Elongation, 1000 p.a.i.	Tensile Strength, p.a.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Softness at 23° C., p.a.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film			Swell, 2 days, 60° C., 100% Humidity
									10 days, 60° C.	24 hours, 70° C.	Water	Oil		
<i>Simple Plasticizers (Continued)</i>														
99	2-Ethylhexanoic acid diester of <i>N,N</i> -bis(2-hydroxyethyl)-2-ethylhexanamide	64	2720	327	63	650	-3	-23	0.4	1.7	1.1	11.0	None	CCCC
100	Tri(<i>p</i> - <i>tert</i> -butylphenyl) phosphate													Dow
101	Glyceryl triricinoleate													Bak
102	Glyceryl tri(acetylricinoleate)	68.6 ^c							1.4		2.5	9.5	None	Bak
<i>Resinous Plasticizers</i>														
103	AROCLOX 1242, chlorinated biphenyl	56.2 ^d							4.5		0.6	0.6	None	Mon
104	FLAKOL plasticizer B-2H, polyester	90	3080	360	57	440	+3	-10	0	0	0	2.4	None	CCCC
105	G.E. 2557, polyester	72	2720	810	63	580	-2	-18	0.9	2.5	2.1	17.0	Very slight	GE
106	G.E. 2559, polyester	90 ^e	2110	215	73	740	6	-6	0.6	2.2	2.7	7.1	Very slight	GE
107	PARACREL C, nitrile rubber	105	2500	410	60	480	+7	-54	0.5		0.2	0.6	None	Nau
108	PARAPLAK G-25, polyester	82	2000	310	60	480	-1	-25	0.2		1.0	0.6	None	RE

^a 1000 p.a.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.a.i. per minute.
^b Dash (—) indicates no data available.
^c 100% elongation, 1000 p.a.i., constant rate of loading of 810 p.a.i. per minute. See Reed.²⁴
^d Mixture of 0.9 part plasticizer to 1 part tricoxyl phosphate.
^e Only composition tested, does not yield standard elongation.
^f Highest plasticizer content tested. Elongation at 1000 p.a.i. = 64%.
^g For key to supplier abbreviations see page 903.

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Waxes in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	275° F. (135° C.)
<i>Resinous Plasticizers (Continued)</i>												
109	PARAFLEX G-40, polyester		ca. 6000	1.15	1.472 at 20° C.	290,000						0.6
110	PARAFLEX G-50, polyester		ca. 2200	1.084	1.470 at 20° C.	3000						1
111	PLASTOLEIN 9720, polyester		ca. 850	1.043	1.462 at 20° C.	481						2
<i>Hydrocarbon-Type Plasticizers</i>												
112	Dow 276-V2, polymolecular product derived from α -methylstyrene			1.021	1.582 at 20° C.							116
113	HB 40, partially hydrogenated mixture of isomeric terphenyls			1.007	1.570 at 25° C.	112.8						
114	KENFLEX L, hydrocarbon resin made by reaction of formaldehyde + dimethylnaphthalenes			1.01 at 15.6° C.		-1.5° C.						
115	NEVILLAC 10° (PHO), a hydrindyl phenol derivative		240 (av.)	1.075 to 1.10 at 15.6° C.	1.597 at 25° C.	23,500 at 25° C.						
116	NEVINOL			1.03 to 1.08 at 15.6° C.	1.596 (av.) at 25° C.	65 to 110						
117	PANAFLEX BN-1, alkylated aromatic hydrocarbon mixture			0.955	1.561 at 20° C.	151.6						
118	SOVALOID C, alkylated aromatic hydrocarbon mixture			1.038		106						

No.	Name	Comparison of Properties of Vinytite Resin VYNW-5 Specimens Plasticized to Equivalence ^a											Supplier ^b	
		Plasticizer p.p.t. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film			Spec., 2 days, 80° C., 100% Humidity ^c
									10 days, 80° C.	24 hours, 70° C.	Water	Oil		
<i>Resinous Plasticizers (Continued)</i>														
109	PARAFLEX G-40, polyester	67 ^d	2780	310	69	760	1	-15	0.3		1.0	5.2	Slight	RH
110	PARAFLEX G-50, polyester	75	2630	320	62	680	0	-16	0.3		1.8	6.5	None	RH
111	PLASTOLEIN 9720, polyester	69	2850	312	66	900	-7	-23	0.3	1.8	1.9	14	None	Em
<i>Hydrocarbon-Type Plasticizers</i>														
112	Dow 276-V2, polymolecular product derived from α -methylstyrene	74 ^b	2420	315	63	650	-5	-24	1.4		0.6	23.6	None	Dow
113	HB 40, partially hydrogenated mixture of isomeric terphenyls	63 ^d	2890	298	66	940	-1	-23	3.4		6.5	15.8	None	Mon
114	KENFLEX L, hydrocarbon resin made by reaction of formaldehyde + dimethylnaphthalenes													Ken
115	NEVILLAC 10° (PHO), a hydrindyl phenol derivative	56.3 ^{a,4}							13.3		0.1	0.1	None	Nev
116	NEVINOL												None	Nev
117	PANAFLEX BN-1, alkylated aromatic hydrocarbon mixture	63 ^d	2800	310	66	920	3	-21	6.1		1.3	16.8	None	Pan
118	SOVALOID C, alkylated aromatic hydrocarbon mixture	61.4 ^d	3100	280			+4	-16	11.5		1.7	9.0	None	SV

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.^b Dash (-) indicates no data available.^c 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reel 25

TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20° C.	Solubility, % at 20° C.				Volatility of Free Plasticizer in Air, mg./sq. cm./hr.	
							In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
<i>Miscellaneous Plasticizers</i>												
119	ARNEAL TOD, mixture of oleic, linoleic, and cyclic nitriles		ca. 278	0.910		30.2						
120	n-Butyl acetyl polyricinoleate, PG16		362 (av.)	0.918 at 25° C. 1.164 at 23° C.	1.460 at 25° C. 1.506 at 23° C.	3100 at 25° C.						
121	Chlorinated paraffin 40			1.164	1.506	152	0.1	8.8				8 80
122	FLEXOL plasticizer B-400, polypropylene glycol mono-n-butyl ether			0.985	1.449 at 20° C.							
123	Glycerol polyricinoleate, No. 15 oil			1.031 at 25° C.	1.483 at 25° C.							
124	HARFLEX 500, monomeric ester type			0.932	1.488 at 20° C.	17.0						
125	HERCOFLEX 600, a pentaerythritol ester		538 (av.)	1.006	1.453 at 20° C.	48.8						
126	ONOPLEX Q-10, octyl fatty-phthalic acid esters			0.952	1.475 at 25° C.	41						
127	PARAFLEX G-60, high-molecular-weight ester		ca. 1000	0.994	1.474 at 20° C.	340 to 380						0.9
128	Plasticizer DP-520, chlorinated aliphatic product			1.126 at 15.6° C. 0.97 at 25° C.	1.478 at 25° C.	66.4 ca. at 54° C.						
129	Polyethylene glycol dilaurate			0.97	1.506 at 20° C.	234						
130	SANTICIZER 180			1.071								
131	STAFLEX KA, monomeric ester type											

No.	Name	Comparison of Properties of VinyLite Resin VYNW-3 Specimens Plasticized to Equivalences ^a										Supplier ^m		
		Plasticizer p.h.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durorometer A at 23° C.	Stiffness at 23° C., p.s.i.	T _g , °C.	Brittle Temperature, °C.	% Loss, 4-mil Film		% Extrac-tion, 10 days, 23° C., 4-mil Film		Spec., 2 days, 60° C., 100% Humidity ^o	
									10 days, 60° C.	24 hours, 70° C.	Water			Oil
<i>Miscellaneous Plasticizers</i>														
119	ARNEAL TOD, mixture of oleic, linoleic, and cyclic nitriles	52.8	2400	290			-44	11.7		0.7	14.8	None	Arm	
120	n-Butyl acetyl polyricinoleate, PG16		Incompatible at 55.3 p.h.r.										Bak	
121	Chlorinated paraffin 40	68 ^o	2820	335	57	900	-5	-28	1.0	0	20	None	Halo, Dia, Hero, Hook	
122	FLEXOL plasticizer B-400, polypropylene glycol mono-n-butyl ether												CCCC	
123	Glycerol polyricinoleate, No. 15 oil												Bak	
124	HARFLEX 500, monomeric ester type	63	2400	319	67	980	-10	-61	1.1	0.4	23.0	None	Har	
125	HERCOFLEX 600, a pentaerythritol ester	58	2910	327	66	900	-3	-28	0	1.1	12.8	None	Hero	
126	ONOPLEX Q-10, octyl fatty-phthalic acid esters												OA	
127	PARAFLEX G-60, high-molecular-weight ester	66	2800	317	67	890	-2	-31	0.3	0.4	11.1	None	RH	
128	Plasticizer DP-520, chlorinated aliphatic product												Drew	
129	Polyethylene glycol dilaurate												Gly	
130	SANTICIZER 180	73	2670	308	67	430	-2	-17		2.7	1.3	12.2	None	Men
131	STAFLEX KA, monomeric ester type												Dee	

^a 1000 p.s.i. tensile modulus at 75% elongation at 23° C. on tensile machine with constant rate of loading of 15,000 p.s.i. per minute.

11. c) Will product scrap be recycled to the mills? If so, will VOC be emitted by the scrap?

No, and as such no VOC will be emitted by the scrap.

12. Are the flow rates listed in the application from points 4 and 5 correct?

Yes, each emission point has two (2) fans and the duct velocity in each is 800 fpm. Given the area of each duct (1.76 ft.²), the flow would equate to 1415 ACFM. In the original application the velocities in the ducts were incorrectly shown in feet per second. Those numbers were actually in feet per minute. Please excuse the typographical error.

13. What will be the percent of the lower explosive limit (LEL) at peak solvent evaporation rates for each exhaust stream?

The new ventilation system will be designed around peak solvent evaporation rates of 40% of the lower explosive limit and will incorporate the use of a lower explosive limit detection and control system designed to enhance safe operation.

14. What will be the percent of the threshold limit values (TLV) near the discharge points \approx 20 feet outside the plant during peak solvent evaporation rates?

The control device, which will provide at least a 90% removal efficiency can be expected to reduce emission levels 20 feet from the exhaust discharge points to well below the current (OSHA) TLV's for Toluene 200 ppm and MEK 200 ppm, however, without extensive modeling an estimated percent of the TLV at the property line cannot be established. Please advise of the necessity of performing said modeling in view of the proposed control program.

15. a) How many employees will work at this plant?

61 full-time employees.

- b) What impact will the emissions from the plant have on the ambient air quality, vegetation and visibility near the plant?

With the incorporation of a control device capable to continuous operation at or above 90% removal efficiency, adverse effects on air quality, visibility and vegetation are not anticipated since the control program reflects state of the art air pollution control technology.

16. If the solvents were recovered, could they be used or sold to a reclamation plant?

Yes, either avenue is a possibility.

17. What is the current delivered cost of the solvents used at the plant?

Toluene \$1.30/gallon

MEK \$.36/pound

18. Has the company made any studies on recovering and reusing the solvents at the plant and, if so, what conclusions were reached?

Yes, preliminary studies suggest that recovery and on-site re-use are well within the realm of feasibility, however, the necessary equipment requires a very substantial capital investment.

19. Please explain price estimate.

Enclosed you will find a copy of a budget quotation for a 25,000 SCFM solvent recovery plant (Appendix VI) which includes equipment and installation. Our original \$700,000.00 figure also included ventilation rework estimates and process control system installation. At this time it is not possible to guarantee that a solvent recovery or thermal incineration device will be of this size or cost since efforts are now underway to reduce the size of the system, however, if a system of such size is necessary due to the nature of the process (non-continuous peak evaporation type), the original \$700,000.00 figure is a very real number.

20. How many gallons of VOC per year will be shipped to Emille for disposal?

Approximately 220 (55 gallon drums) which would contain some 5,000 gallons of VOC. (By estimate.)

DISCUSSION:

In accordance with our recent conversation and the historical data provided in item one, David 'M' Company may be considered a major source of air pollution in an uncontrolled state. As such a B.A.C.T. type standard would be applicable and will likely approximate the 2.9 lbs. VOC/gallon of coating applied or 90% removal of VOC from the exhaust stream. David 'M' Company cannot meet the afore-mentioned low solvent technology standard and will be obligated to propose the control device intended for use in meeting the 90% removal standard referenced above. While I cannot guaranty at this time the exact type of control device which David 'M' Company will install, I can state that both recovery and thermal destruction techniques capable of delivering 90% removal of VOC's from exhaust streams are being evaluated.

The attached schedule estimates the time periods associated with specifying the type of control device and ultimately installing same. Would you please advise me of the proposed schedule's acceptability. We believe this schedule to be a fair appraisal of the time necessary to complete the installation of a project of this magnitude. Should you require further information and/or assistance, please do not hesitate to contact me directly.

Sincerely,



Michael A. Ware
Graphics Div. Env. Coord.
Chemicals & Coatings Group
Wheelabrator-Frye, Inc.

MAW:bjw
Attach.

October 19, 1983

ESTIMATED

OVERALL COMPLIANCE SCHEDULE

DAVID M COMPANY VOC CONTROL PROJECT

PHASE

Data Accumulation	January 1, 1984
Spreader Enclosure - (1) For Temperature Evaluation	January 15, 1984
Process Modifications and Development	April 1, 1984
Methods Comparison/Analysis - Thermal Incineration Vs. Recovery	May 15, 1984
Final Specification Development	July 1, 1984
Final Specification/Quotations and Analysis	September 1, 1984
Appropriations Request Development and Approval	December 1, 1984
Oven Enclosure and Exhaust System Installation	March 1, 1985
Solvent Recovery Delivery and Installation	August 1, 1985
Operational Shakedown, Debugging and Compliance Testing	December 1, 1985

APPENDIX VI



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~008020~~ 240189

COPY

YOUR REF:

OUR REF: E 0379
S 7212

DATE: 2 August 1983

PROPOSAL

FOR

SOLVENT RECOVERY SYSTEM

TO BE LOCATED AT:

DAVID M. COMPANY

201 VALENTINE WAY
LONGWOOD, FLORIDA



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: ~~900020~~ 240189

YOUR REF:

OUR REF: S 7212

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

SOLVENT RECOVERY PLANT

ACTIVATED CARBON ADSORPTION UNIT

AIRFLOW.....25,000 scfm.
AIR TEMPERATURE.....140° F.
SOLVENT CAPACITY.....50 gph (360 lb/hr)
SOLVENT.....Toluene

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL

David M. Company

Proposal No. S 7212

PLANT DESCRIPTION

The solvent laden air would be filtered to remove dust, cooled to approximately 95° F by the air cooler and then delivered by the fan to the three adsorbers.

Two of the three adsorbers would normally be handling the solvent laden air whilst the other would be regenerating with steam to extract the solvent from the carbon or would be held on "stand-by" following "steaming" and a short dry/cool period.

The adsorbers would cycle consecutively and automatically with override of the cycle time and steam input by gas analyser control. The proposed adsorbers are of annular bed design, the solvent air after filtering and cooling passes through an automatic inlet valve into the outer annulus of the adsorber, through the carbon bed and then exhausts to atmosphere through the inner annulus and the automatic exhaust valve.

At the end of an adsorbing period the air inlet and outlet valves close and an automatic steam valve allows steam to enter the inner annulus of the adsorber and pass through the carbon bed to the outer annulus from which point the steam together with solvent vapour extracted from the carbon passes to the condenser. During steaming some condensation occurs on the outer shell of the adsorber and is drained through filters to join the main condensate flow from the condenser to the decanter.

At the end of the steaming period the adsorber automatic steam valve shuts and the air inlet and outlet valves open to allow solvent laden air to again pass through the adsorber for a short period to dry and cool the bed (during this dry/cool period the solvent laden air is shut off from the other adsorber). The inlet valve of the dried/cooled adsorber would then close and the adsorber would be held on "stand-by" until the gas analyser, sampling the exhaust of the other adsorber, or the override timer, gives the signal for that adsorber to change to the steaming cycle, the stand-by adsorber being first brought back into service, so that there is no interruption to the adsorber cycle.

Any incondensibles from the condenser return through the condenser vent pipe to the inlet duct for recycling through the adsorbers. The condenser vent is fitted with high temperature alarm to indicate overloading from excessive steam flow or shortage or failure of the cooling water supply.

Safety features include adsorber pressure and vacuum relief valves and liquid seals.

David M. Company

Proposal No. S 7212

We would supply the following:

1. AIR FILTER - 25,000 scfm capacity

a) Pre-Filter Section

With removable and disposable fiberglass panels to protect and extend the life of the main filter.

b) Main Filter Section

With disposable filter elements of the HEPA type of fiberglass material, with an efficiency of better than 99% for particles down to 0.3 microns per standard tests.

The whole of the above would be incorporated in a galvanized steel case with access doors to facilitate servicing.

A differential resistance gauge would be fitted.

2. AIR COOLER - 25,000 scfm capacity.

Would be of finned tube design with copper tubes and aluminum fins to cool the inlet airstream from 140° F to approximately 95° F when supplied with cooling water at a temperature of maximum 86° F.

The air cooler coils would be housed in a galvanized steel case and would be drainable.

3. FAN AND MOTOR

The fan would be of the high efficiency type with airfoil impeller capable of handling 25,000 scfm of solvent laden air, when at an actual temperature of approximately 95° F, at a total pressure of approximately 18" w.g., allowing for 2" w.g. suction before the air filters.

The casing would be of substantial carbon steel construction and the unit would have spark-proof features to AMCA standards.

The impeller would be driven through a flexible coupling by a 125 HP explosion proof motor to Class 1, Group D, Division 1, 460 volt, 3 phase, 60 cycles.

A manually operated radial leaf damper would be fitted at the fan inlet for regulation of the airflow.

David M. Company

Proposal No. S 7212

4. THREE ADSORBERS - 6'8" diameter x approximately 10'0" high shell.

Each capable of handling 12,500 scfm of solvent laden air calculated at 70° F, when at an actual temperature of approximately 95° F.

The adsorbers would be of the annular carbon bed type with the carbon contained between stainless steel panels formed up into cylinders and reinforced by carbon steel stiffening bands. The activated carbon bed and screens would be supported upon a carbon steel tray to which would be connected the solvent laden air inlet valve, vapour piping to condenser and liquor drains. The air exhaust valve and the steam valve would be connected to the outlet branch of the adsorber.

The carbon bed and screens would be contained within a removable carbon steel shell, fitted with access and inspection holes with bolted cover plates. Flanged connections would be fitted to the tray so that the carbon could be easily and quickly run off into drums.

Vacuum and pressure relief valves would be provided for each adsorber.

Handrails would be fitted round the tops of the adsorbers, with a walkway over and between the adsorbers, and with an access ladder from ground level, in accordance with the requirements of OSHA.

5. ACTIVATED CARBON

We would provide the initial charge of approximately 6400 lbs of high quality carbon for each of the three adsorbers. The carbon would be manufactured by Sutcliffe Speakman and would be of the grade most suited to the required duty.

6. INLET AND OUTLET AIR VALVES - 26" diameter

We would provide an inlet and outlet valve for each adsorber 26" diameter of the mushroom type of substantial construction arranged for pneumatic operation through the automatic control mechanism.

Renewable seats would be fitted in the valve bodies and access holes with bolted cover plates provided. An indicator would be fitted to each valve so that the attitude of the valve could be easily ascertained.

David M. Company

Proposal No. S 7212

7. CONDENSER

To condense the steam and solvent vapours leaving an adsorber during the steaming period.

The unit would be of the tubular surface type with the shell in 304 stainless steel and the tubes in 316 stainless steel. The water boxes and tubeplates would be in carbon steel.

The condenser vent pipe (fitted with thermometer with high temperature alarm contacts) would be connected to the inlet duct so that any Incondensibles displaced from the adsorbers would recycle.

8. ONE DECANTER - 3'0" diameter x 4'6" high

It would be of the self-adjusting decanting type to separate automatically the condenser water and solvent. The tank would be of carbon steel welded construction fitted with a bolted cover plate and all the necessary connections. A sight glass would be fitted in the solvent outlet.

9. TRUNKING

We would provide the interconnecting trunking between the air filter, air cooler, fan and adsorbers, but do not allow for any trunking prior to the Inlet manifold of the air filter. The trunking would be of carbon steel of welded construction flanged at suitable intervals and be adequately supported.

10. THREE EXHAUSTS

An exhaust stack would be provided for each adsorber, of carbon steel welded construction, galvanized and would terminate about 6' above the adsorber platform.

11. PIPING AND VALVES

We would provide pipework to interconnect the above and manifold the service pipes to terminal flanges at the battery limits.

Vapour and liquor piping would be in 304 stainless steel, utility piping in carbon steel.

We include for all isolating valves, steam traps, pipe supports, etc.

David M. company

Proposal No. S 7212

12. PLANT CONTROL UNIT

Would consist of the following:

a) Gas Analyser (Infra-Red Type)

To measure the solvent content in the exhaust airstream of the adsorber next due for desorbing and to delay steaming until the set point was achieved. However, an adjustable override timer would be incorporated into the program to limit the period of such a delay.

A strip recorder for the exhaust gas concentration would be included and the unit would have a "high level" alarm.

b) Programmable Logic Controller

Of the solid state type which on receipt of the appropriate signals would cause the adsorber valves to sequence as appropriate. The program logic would be such that the next step would not be commenced until the previous one had been proven completed.

In the unlikely event of failure of the P.L.C. the solvent recovery plant could be operated manually by the use of pilot air solenoid valves inbuilt into the panel.

c) Annunciation Section

Of the 'Panalarm' type or similar to alarm:
"high level" - gas analyser output
"high temperature" - condenser vent

d) Control Panel

The above items would be mounted in a free-standing enclosure, suitable for indoor location in a non-hazardous area, assumed within 50' of the adsorbers.

The panel would be pre-wired and tubed to terminal points on the frame. We exclude the impulse tubing from the panel to the plant items.

David M. Company

Proposal No. S 7212

13. INSTRUMENTS

We would provide the following indicating type instruments which would be locally mounted on the plant:

- 4 Dial type thermometers
- 1 Steam pressure gauge
- 1 Differential pressure gauge (resistance through air filter)
- 1 4-Point pressure gauge (air pressure and suction within the plant)

14. STRUCTURES

We would provide structures to support items of the plant.

15. PAINTING

Where applicable, plant items would be primer coated before shipment.

16. COMPRESSED AIR

You to supply compressed air, clean, dry and oil free at a pressure of approximately 80 psig.

17. DRAWINGS AND INSTRUCTIONS

We would provide three (3) copies of each of the following:

- Foundation drawing
- Flowsheet
- Plant arrangement drawing
- Major assemblies

Plus two (2) copies of:

- Erection drawings and instructions
- Operating and Maintenance Manual

A list of recommended spares would also be provided.



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA,
1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: 900020 240189

YOUR REF:

OUR REF:

DATE: 2 August 1983

David M. Company
201 Valentine Way
Longwood, Florida

TERMS, PRICE AND CONDITIONS

PRICE OF PROPOSAL

PROPOSAL NO. S 7212

For the supply of the whole of the items as specified in this proposal, delivered only to your works in Longwood, Florida (offloading from the transport and installation by others) would be the sum of approximately.....\$ 370,000.00

(THREE HUNDRED SEVENTY THOUSAND DOLLARS)

Any Federal, State or local taxes that may be applicable are excluded.

TERMS OF PAYMENT

25% of quoted price with order.

45% of quoted price in three (3) equal payments pursuant to invoices on the following basis:

- a) upon submission of preliminary drawings comprising plant arrangement and foundation drawings and flowsheet.
- b) upon submission of confirmation that the manufacture is at least 40% complete.
- c) upon submission of confirmation that the manufacture is at least 70% complete.

25% of quoted price upon despatch or when ready for despatch if despatch date is delayed by you.

5% of quoted price on start-up and successful operation, at latest three (3) months after despatch, or when ready for despatch, whichever is sooner.

Continued.....

David M. Company

Proposal No. S 7212

TERMS, PRICE AND CONDITIONS...continuedDESPATCH

Approximately six (6) months from receipt of order assuming that there are no delays due to changes in specifications or approval of drawings.

EXCLUSIONS

We do not include for any motor controls, electrical wiring, or any special cold weather protection or insulation, or any parts, materials, work for services of any kind whatsoever except those specifically mentioned in the body of the proposal.

STANDARD OF MANUFACTURE

The goods described in this proposal would be manufactured to normal commercial standards and be suitable for the duty specified. Should you desire manufacture to any specific standards, codes or inspection schedules other than those allowed for then we will be pleased to comply with your requests, but any extra charges so incurred would be in addition to the prices quoted.

VALIDITY OF PROPOSAL

This proposal is open for acceptance for a period of thirty (30) days from the date of this quotation and after this period confirmation must be obtained that the terms and conditions contained herein are still valid.

SUTCLIFFE SPEAKMAN INCORPORATED
William C. Moses
Vice PresidentE. & O. E.

WCM/pmd

David M. Company

Proposal No. S 7212

UTILITY REQUIREMENTS

The following are the approximate utility requirements of the plant when operating at design load conditions. The figures are subject to confirmation by our final design study.

STEAM.....1190 lb/hr, dry saturated quality
at 40 psig.

COOLING WATER.....170 gpm, inlet 86^o, maximum,
return 115^o.

POWER.....100 KwH.

COMPRESSED AIR.....20 scfh.

SPACE

The equipment, we estimate, would require a floor area of approximately 45' x 20'.

The height of the equipment would be approximately 18', but if it is to be located inside, then provision should be made for lifting of the adsorber shells.

APPENDIX 'A'OUR PRELIMINARY PROPOSAL FOR THE TURNKEY INSTALLATIONOF THE PLANT QUOTED IN MAIN PROPOSAL NO. S 72121. FOUNDATIONS

We would provide concrete foundations for the appropriate items of our equipment.

Our price is conditional on the understanding that the ground is of suitable load bearing capacity, that no rock blasting or piling would be required and that all work would be above the water table.

2. MOTOR CONTROL CENTER

We include the starters for the fan motor, the water cooling tower fan and the recirculating water pumps, and which would be suitable for location in a non-hazardous area which we assume will be by the control panel.

3. WATER COOLING TOWER

We would supply and install a water cooling tower designed to cool the required amount of water from 115° F to approximately 86° F. The tower would be of induced draft type and complete with fan and motor. The system would include two pumps with motors (one normally operating, one stand-by) and the water circulating piping and valves between the water cooling tower and the air cooler and condenser of the recovery plant, the water cooling tower to be at ground level and located adjacent to the recovery plant.

Corrosion Inhibitor S? Chemically

4. INSTALLATION

We include for the necessary labour, tools, moving and lifting gear including cranes, for the installation of the items of equipment specified in this proposal onto the foundation.

4. INSTALLATION...Continued

It is understood that the work would be undertaken during our normal working hours and that we and our sub-contractors or agents would be allowed free and unrestricted access to the site.

Any overtime worked at your request would be chargeable extra at the rate in force at the time.

You would be responsible for offloading from the transport any items delivered prior to the start of the installation work and provide suitable storage and protection for all parts close to the battery limits.

All temporary site support facilities including but not limited to water, electricity, steam, drain, office and telephone would be provided by you at your expense, adjacent to the site.

Note that the quoted price assumes we will have continuity of work on site.

5. SUPERVISION

We include for the services of our own suitably qualified supervisor to supervise and give instructions to the labour provided by our sub-contractors, to secure the installation of the equipment in a competent manner.

6. IMPULSE TUBING

The pneumatic tubing from the appropriate plant items to the control panel would be supplied and installed in a workmanlike manner. It is assumed that the tubing run is a maximum of fifty feet.

7. ELECTRICAL WIRING

We would supply and install all necessary electrical wiring, cable trays, etc. to interconnect the various items of plant to the motor control center and the control panel, assuming a maximum cable run of fifty feet from the adsorbers. Where appropriate, the wiring would be to explosion proof, classification.

David M. Company

Proposal No. S 7212

8. INSULATION/WINTERIZATION

Insulation and heat tracing of appropriate equipment within our battery limits effected, assuming a minimum ambient temperature of 25° F.

9. START-UP

We include for the services of a skilled engineer for a period of two (2) consecutive weeks on site to start-up the plant and provide instructions to your personnel on its operation.

Any extra time required by you or non-continuity in attendance would be at additional cost.

10. INSURANCE

We would provide and maintain the following insurances and would require any and all sub-contractors employed by us at your site to procure and maintain the same type and amount of insurance.

- a) Workman's Compensation and employer's Liability
- \$100,000.00 each accident and aggregate disease.
- b) Comprehensive General Liability
- Bodily Injury and Property damage of up to \$1,000,000 combined single limit.
- c) Comprehensive Automobile Liability
- To cover owned, non-owned, leased and hired cars, combined single limit \$1,000,000.

Certificates of the above shall be provided to you prior to the start of our on site work.

In addition to the above and in order to insure the equipment from its delivery to your site, through installation to your acceptance of the plant, we would take out "all risks" insurance plus "builders risk" and "Installation floaters" at the appropriate time and at your expense.

David M. Company

Proposal No. S 7212

BUDGETARY PRICE

Our approximate additional price to that quoted in Main Proposal No. S 7212 for the turnkey installation of the plant as detailed in this Appendix 'A' would be the sum of.....\$ 160,000.00.

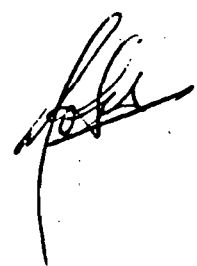
(ONE HUNDRED SIXTY THOUSAND DOLLARS)

COMPLETION

We would expect to supply and install the specified plant in approximately nine (9) months from receipt of order.

TERMS AND CONDITIONS

Otherwise as Main Proposal No. S 7212.



SOLVENT RECOVERY

WITH

ACTIVE CARBON

SUTCLIFFE SPEAKMAN, INC.

Sutcliffe Speakman have specialised for over 50 years in the design and manufacture of Solvent Recovery plant. Their unique position in the industry stems from a background of almost 80 years experience in the manufacture of active carbon for general adsorption and catalytic applications.

SOLVENT RECOVERY WITH ACTIVE CARBON

Our recovery plants are in world-wide use in a great variety of solvent-using industries, ranging from coating, impregnation and printing of materials to the manufacture of man made fibres.

Our active carbon plants for the recovery of industrial solvents, from air or gas mixtures, can give a recovery efficiency of at least 99% by volume depending on the conditions. In addition to vapour phase recovery, active carbon plants can efficiently extract and recover solvents and organic compounds from water and other liquid mixtures.

MONEY SAVING

Recovery plants save industry millions a year and the dramatic savings on solvent costs ensure more economic production.

CLEAN AIR

The adsorption efficiency of Sutcliffe Speakman plant is extremely high, effecting an important contribution to clean air.

LIQUID EFFLUENTS

Can also be purified with an active carbon plant.

WORKING CONDITIONS

A general improvement in working and safety conditions often results from the effective collection system required to achieve high overall efficiency.

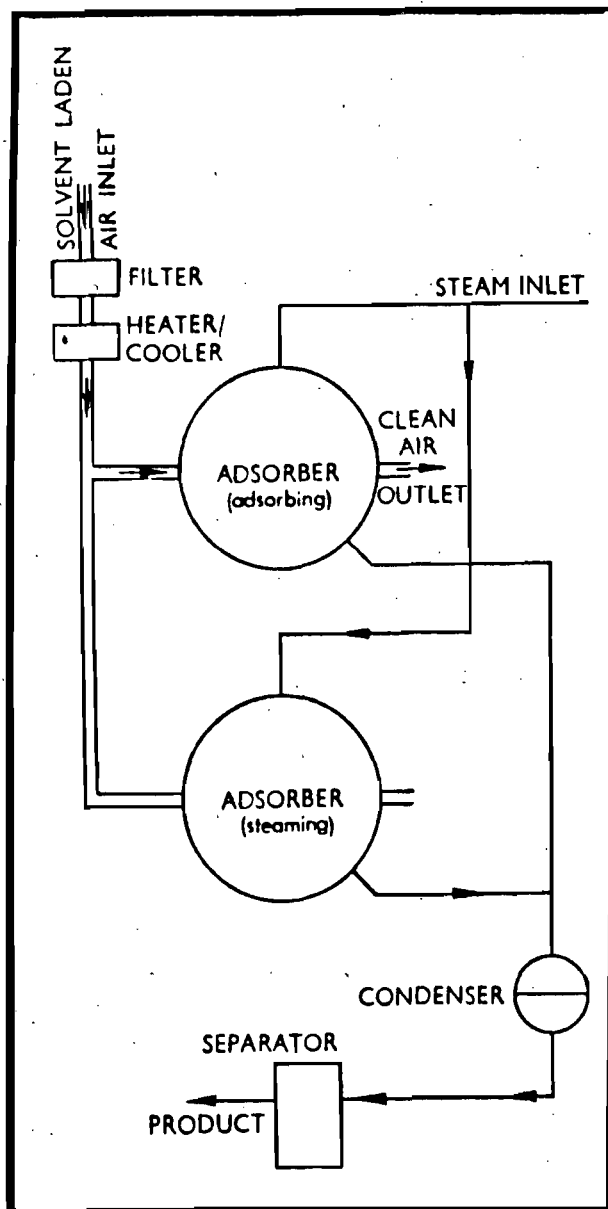
WHEN IS IT WORTHWHILE?

If you are using more than 200 gallons a week of alcohols, esters, ethers, ketones, hydrocarbons, chlorinated compounds and other organic solvents, it is certainly worth giving preliminary consideration to the economic potential of installing recovery plant. The actual economics depend upon the cost of new solvent, the concentration of solvent in air entering the plant, the number of hours per week which the process will operate and the possible ancillary equipment which may be necessary to prepare the recovered solvent for re-use. This may be required if the solvent is water miscible or where a mixture of different solvents may be recovered together and necessitate separation.

Sometimes solvent quantities as low as 1 gallon per hour can be economically recovered. WRITE TO US, WE WILL ADVISE WHETHER A DETAILED STUDY IS JUSTIFIED.

GASEOUS PHASE ADSORPTION

BASIC FLOW DIAGRAM



HOW IT WORKS

The Sutcliffe Speakman system draws air laden with solvent vapour through a ducting into the plant when, if necessary, it is first filtered to remove dust or other contaminants and then heated or cooled to a suitable temperature before passing into an adsorber which contains a bed of active carbon. The solvent vapour is adsorbed on the carbon and the clean air is discharged to atmosphere. When the carbon bed has adsorbed the designed charge of solvent, the vapour/air stream is switched to a second adsorber and steam at low pressure is introduced to the charged adsorber which causes the carbon to release, in vapour form, the solvent which it adsorbed. The steam/solvent vapour mixture is then condensed and the solvent recovered.

If the solvent is insoluble in water, a separator is provided to divide the condensate into a solvent layer and water layer which automatically leave the separator by gravity flow. If the solvent is water soluble, the condensate is taken to a distillation unit or liquid phase adsorption system for final separation.

CARBONS

The wide range of Sutcliffe Speakman carbons developed specially for solvent recovery give a high degree of adsorption efficiency at relatively high velocities. These carbons also have low retention characteristics, so that steam usage for the regeneration of beds is minimised. The most suitable carbon is selected to meet the particular conditions under which the plant will operate.

PLANT

Sutcliffe Speakman design their adsorbers to give maximum surface area of carbon, thereby ensuring the minimum resistance to air-flow, and a relatively lower power consumption. Adsorbers are designed with static or rotary beds.

Automatic operation of the adsorber valves with variable time cycle, is effected by instrumentation or by the robust Sutcliffe Speakman standard control unit.

In normal applications the adsorbers are not subjected to pressures of more than 0.5 lbs/square inch. Where necessary, plants can be designed for operating at higher pressures.

If required, plant can be designed for further extension to suit clients needs.

The rotary bed plants and a range of static units can be supplied as fully "packaged" plants. Wherever practicable, all plants are pre-assembled in our works before despatch. Full erection and maintenance services are available including contract visits by Technical Service Engineers.

SOLVENT CONCENTRATION IN AIR

The adjacent diagram shows the quantities of solvent in relation to air flow at given volume/volume concentrations, corrected to the normal temperature and pressure conditions at the inlet to the recovery plant, with indication of solvent value per annum.

It is important that the concentration of solvent in the air stream from a process is, where applicable, well below the lower explosive limit of that particular solvent vapour in air, to ensure a satisfactory operational safety margin. It is normally considered that the quantity of air flow from a process should be sufficient to ensure that the concentration of solvent vapour in air is no more than a maximum of 50% of the lower explosive limit.

Conversely it is usually necessary to ensure that the concentration is not too low, or the size of plant required to handle air flow may be larger than can be justified economically in relation to the quantity of solvent to be recovered. The ideal concentration range is usually between 20%-40% of the lower explosive limit.

Sutcliffe Speakman technical staff are available to advise on the collection of solvent from your process to ensure that maximum overall recovery efficiency is achieved, and that the desirable air flow from an economic point of view is sufficient to collect the solvent vapours effectively.

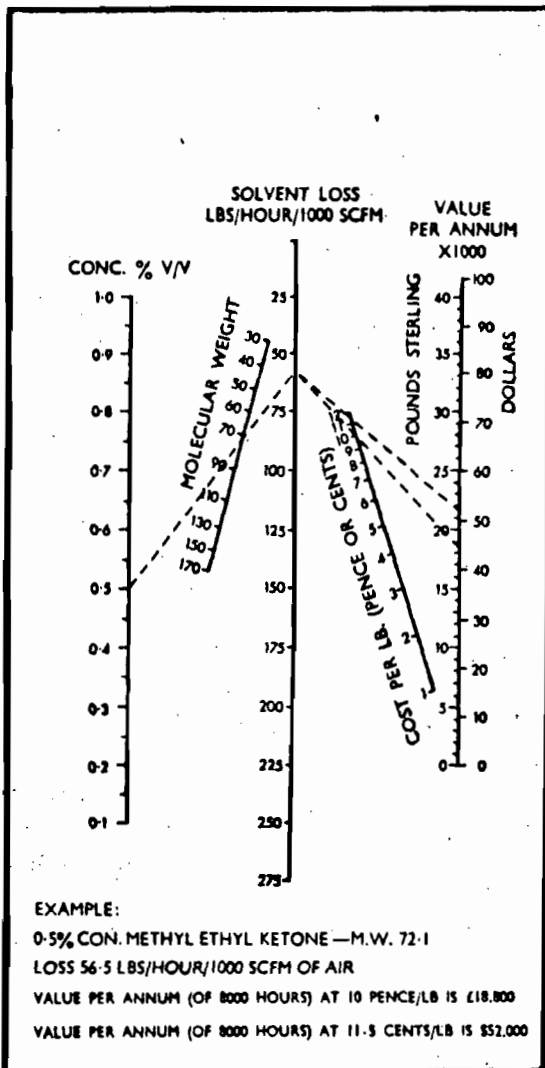


TABLE OF MOLECULAR WEIGHTS AND LOWER EXPLOSIVE LIMITS FOR SOME TYPICAL INDUSTRIAL SOLVENTS

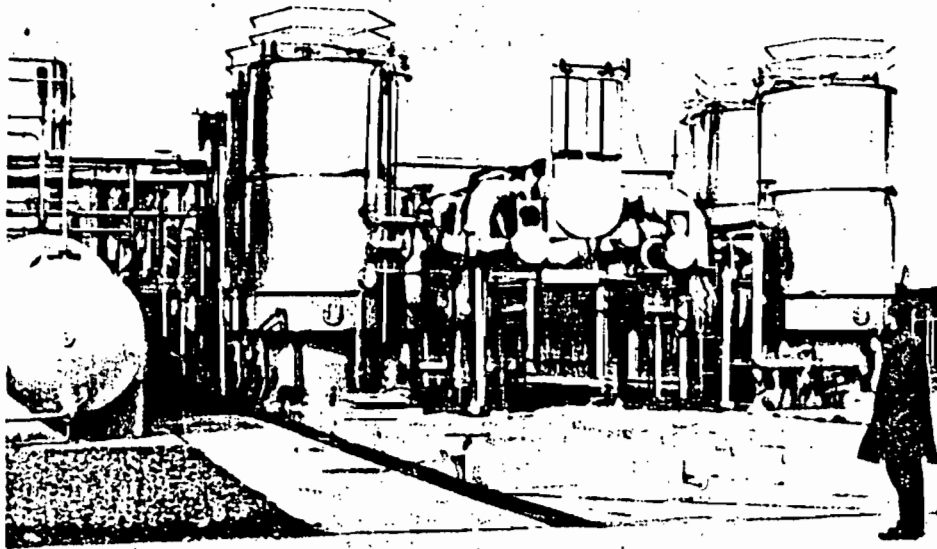
	A	B		A	B
Acetone	58.08	2.15	Methyl Acetate	74.08	4.1
Benzene	78.11	1.4	Methyl Alcohol	32.04	8.72
Butyl Acetate (Normal)	116.16	1.7	Methyl Ethyl Ketone	72.10	1.81
Butyl Acetate (Iso)	116.16	2.4	Methylene Chloride	84.93	Non-flam.
Butyl Alcohol (Normal)	74.12	1.45	Nonane	128.25	0.74
Butyl Alcohol (Iso)	74.12	1.68	Octene	114.23	0.95
Carbon Disulphide	78.13	1.0	Pentane (Normal)	72.15	1.3
Carbon Tetrachloride	153.84	Non-flam.	Pentane (Iso)	72.09	1.3
Chloroform	119.39	Non-flam.	Perchloroethylene	165.85	Non-flam.
Cyclohexane	84.16	1.35	Propyl Acetate (Normal)	102.3	2.0
Decane (Normal)	142.28	0.7	Propyl Acetate (Iso)	102.3	2.0
Dichloroethylene	97	9.7	Propyl Alcohol (Normal)	60.09	2.15
Ether (Diethyl)	74.12	1.85	Propyl Alcohol (Iso)	60.09	2.02
Ethyl Acetate	88.10	2.25	Tetrachlorethane	167.86	Non-flam.
Ethyl Alcohol	46.07	3.3	Tetrahydrofuran	72.10	1.84
Ethylene Dichloride	99.97	8.2	Toluene	92.13	1.3
Furfural	96.08	2.1	Trichloroethylene	131.4	Non-flam.
Heptane (Normal)	100.2	1.0	Xylene	106.16	1.0
Hexane (Normal)	86.17	1.25			

A Mol. Wt.
 B Lower Explosive Limit in Air % V/V at 20°C and approx. atmospheric Pressure

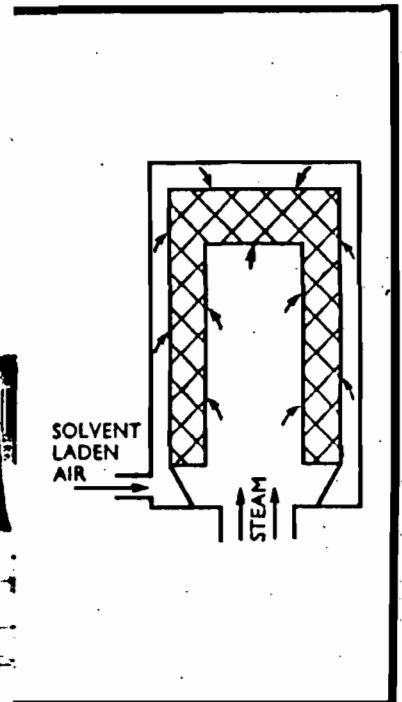
Sutcliffe Speakman manufacture two principal types of Gaseous Phase Recovery Plant— Both types employ active carbon.

STATIC BED PLANT

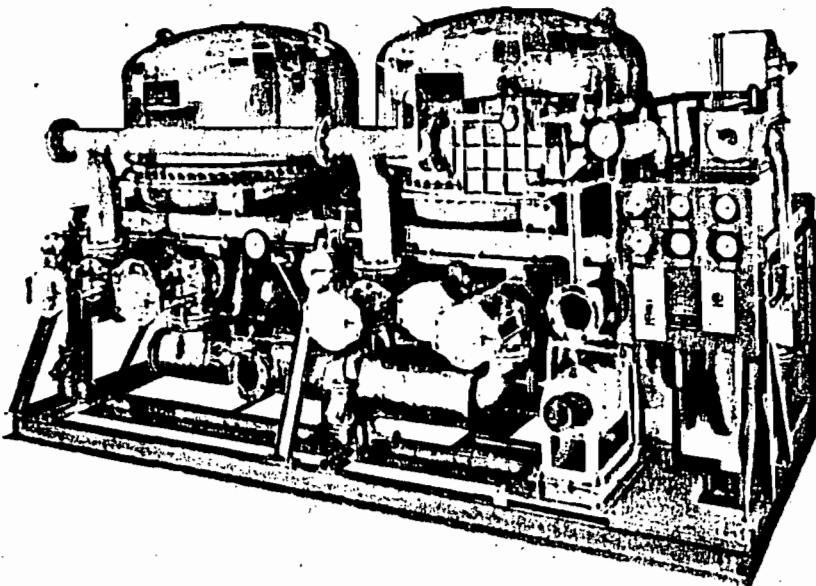
A static plant usually consists of two or more adsorbers each containing a vertical annular or flat bed of active carbon. One or more of the adsorbers receives the solvent laden air whilst one or more are being steamed out to recover the solvent. The adsorbers are suitably valved and cycle automatically so that the process is continuous. The smaller plants can be of packaged design, skid mounted. Plants are individually designed to meet customers specifications.



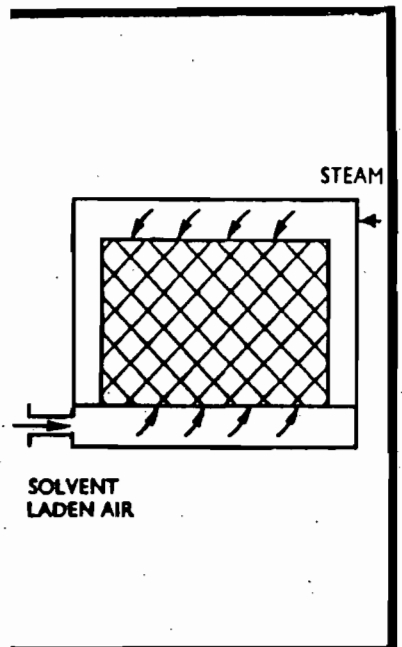
Adsorption section of a large recovery complex at an Acetate Fibre Factory in Belgium. The plant is arranged for future extension.



ANNULAR BED



Packaged type fully automatic plant of cylindrical flat bed design



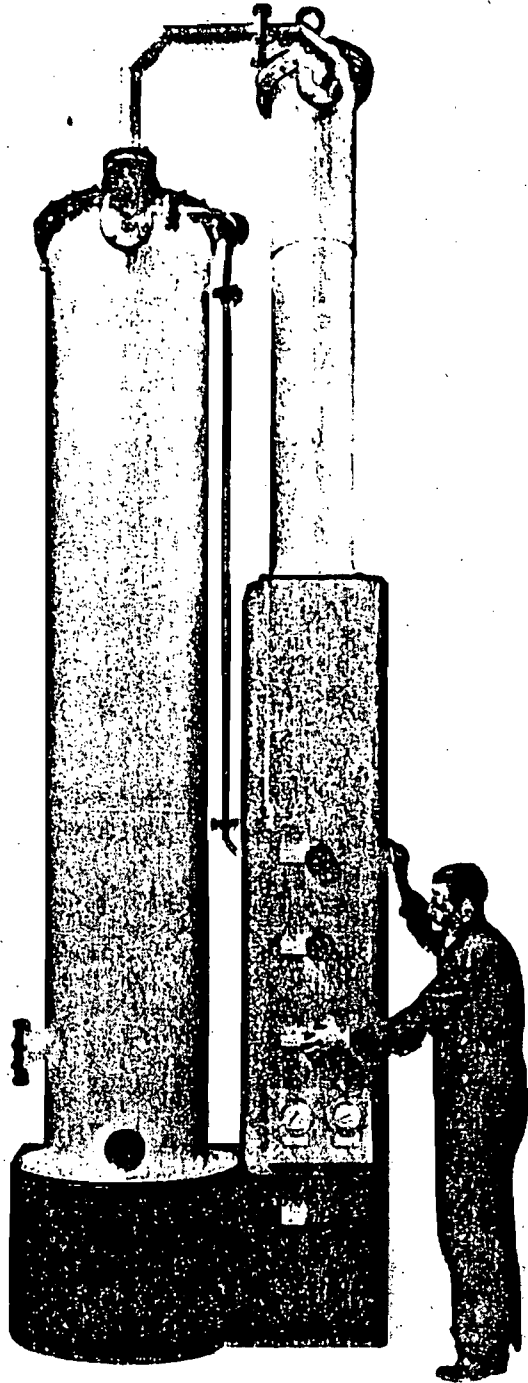
FLAT BED

LIQUID PHASE ADSORPTION

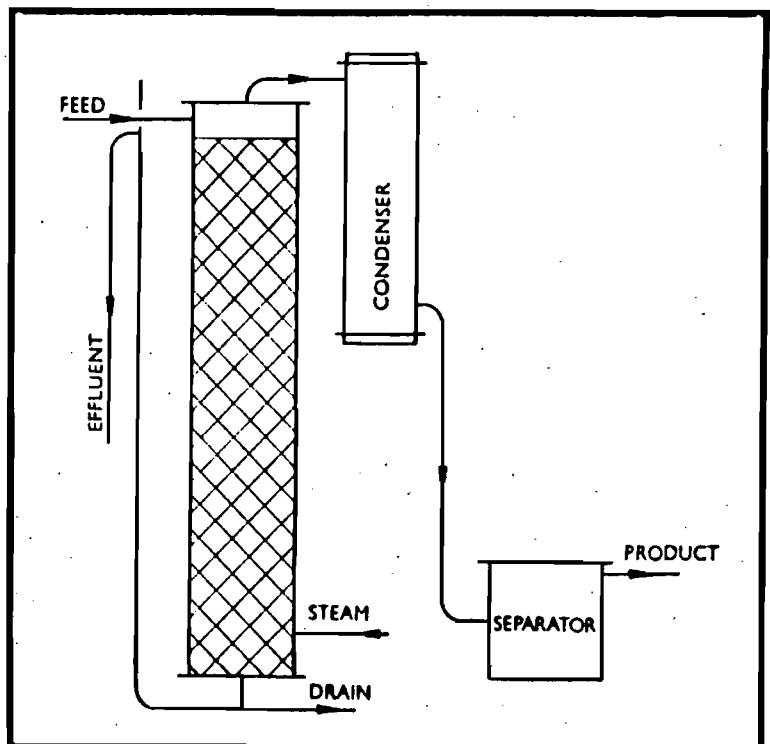
This system uses active carbon for the adsorption of small amounts of certain water-miscible solvents from solutions with water. The weak solvent/water liquor is passed through an active carbon bed and, when the carbon has adsorbed the designed charge of solvent, the solvent is distilled off by our patented system. This process effects substantial increase in concentration of a weak solvent/water solution, the water in the inlet feed passing through the bed and to effluent drain. During steaming of the carbon a small amount of water is introduced and the products of steaming taken to a separating tank from which the water layer recycles through the system.

In certain cases the carbon can be used to adsorb selectively a particular solvent component from a mixture of solvents with water.

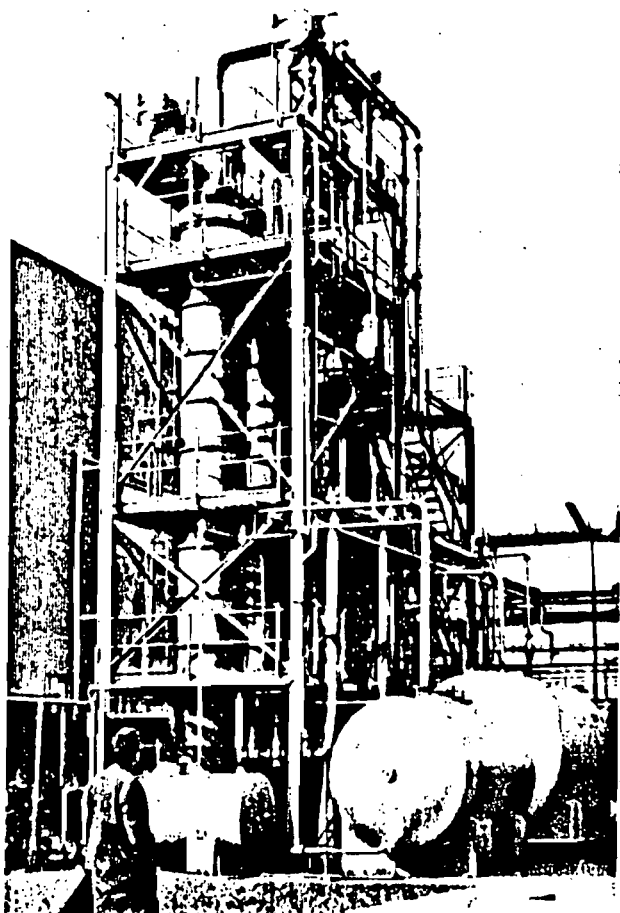
A particular feature of this system is the extreme simplicity of operation and control which can be manual or automatic.



Liquid phase adsorption unit.



DISTILLATION



Section of a large distillation complex.

Sutcliffe Speakman supply a complete range of distillation equipment, which may be required in the separation of water-miscible solvents from the products of recovery of the main adsorption plant, or for other purposes.

The range of distillation equipment includes both perforated plate and bubble cap designs, and the operations can be automatically controlled in relation to steam input and reflux ratio.

Stills can be supplied either for continuous or batch operation.

There is also a range of small units for the recovery of solvents from contaminated mixtures, i.e. by extraction of residues from waste lacquers or in cases where solvents are used for washing down printing and paint machines, etc., the solvent can be cleaned for further use.

ENQUIRIES

In submitting enquiries for evaluation, it is helpful if the following preliminary details can be submitted.

Gas or liquid flow rate

Operating pressures

Temperature and relative humidity of air or gas stream

Quantity, nature and value of compounds to be extracted

Other substances in air, gas or liquid stream of a contaminant or corrosive nature

Normal operating time of process

Details of steam, cooling water, power and compressed air services available.

TYPICAL SOLVENT RECOVERY APPLICATIONS

Acetate Fibre · Insulated Cables · Balata Belting · Leather Cloth · Hydrocarbon Extraction (Natural Gas) · Metallic Foil Printing and Lacquering · Compressed Asbestos Fibre Products
Oil Extraction · Degreasing · Paper Impregnating and Lacquering · De-sulphurising of Gas
Paper Printing · Dry Cleaning · Pressure-Sensitive Tapes · Explosives · Rotogravure Printing
Film Casting · Rubber-Dipped Goods · Rubber-Spread Goods · Surgical Bandages
Shoes, etc. · Solvent Manufacture · Tail Gas from Petroleum Wells · Viscose Fibre
Viscose and Acetate Film · Weatherproof Sheeting · Process Off-Gas



SUTCLIFFE SPEAKMAN INC

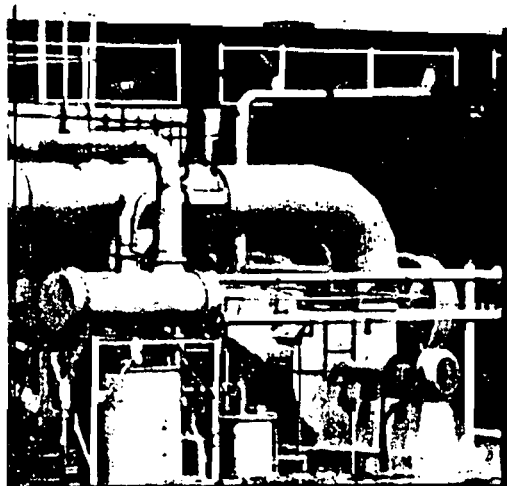
SUITE 200, HEAVER PLAZA,
1301 YORK RD. LUTHERVILLE, MARYLAND 21093
TELEPHONE: 301-337-2800
TELEX: 909020

The many facets of activated carbon

**Reprinted from
Chemical Age 26 June 1981**

activated carbon

and business advantages of solvent recovery



toluene and SBP2.

annular bed is the most efficient when it comes to steaming the solvent off the carbon bed (desorption). This is because the steam enters the inner annulus of the adsorber working its way outwards in contrast to other

designs where the adsorber vessels have to be efficiently lagged to prevent waste of heat energy. With the annular bed design by the time the steam has worked its way through the carbon bed to the outer annulus it has performed its useful work, and the cooler outer shell of the adsorber vessel serves as an air condenser and relieves the load on the main condenser with regard to the amount of cooling water required.

The basic components of a typical solvent recovery plant are:

- Air filter (to remove any airborne particulate matter in the solvent-laden air stream).
- Air cooler/heater (to temperature adjust air stream for safe and good adsorption on the carbon bed).
- Main fan(s) (to provide correct rate of airflow through the adsorbers).
- Activated carbon adsorbers (where the solvent(s) are adsorbed/desorbed).
- Condenser (to condense solvent vapour and steam from the desorption cycle).

- Separator (decanter) — (where non-miscible solvent/water mixture separates).

- Product tank (for collection of solvent from the separator).

In operation the solvent-laden air is passed to two of the adsorbers while the solvent is desorbed (by steaming) in the third adsorber.

The process operates automatically and continuously, usually with infrared gas analyser control to ensure optimum efficiency, and each adsorber consecutively undergoes an adsorption and desorption cycle.

The simplest type of solvent recovery is where there is a single solvent which is non-miscible with water, eg, xylene. This solvent will collect as the upper layer in the decanter and, being virtually insoluble in water, can simply be decanted for re-use whilst the lower (water) layer is passed to drain.

If, however, a mixture of solvents is being recovered and some of these are miscible with water then additional plant is necessary to separate the solvents. This usually takes the form of distillation and dehydration equipment depending on the solvents involved and the final specification required by the customer for the recovered solvents.

Each solvent recovery system has its own particular technique for recovery, but the basic adsorption/desorption on activated carbon is common to all. When designing a solvent recovery plant for a particular set of conditions one must have detailed information on such parameters as airflow, air temperature, humidity, solvent concentrations, maximum/minimum process conditions, and details of any impurities in the solvent-laden air stream. The last factor can be particularly important since it can have an important bearing on materials of construction and expected carbon life.

There are several aspects which require careful attention to ensure that the best efficiency of solvent recovery is achieved and that at the same time minimum energy is consumed in the running of the plant. The first practical step which can be taken to maximise recovery efficiency is to present as high a solvent concentration as is safely possible with minimum airflow.

This will be largely governed by the type of process upstream of the

Table 1: Some typical results of solvent adsorption by carbons

Solvent	Concentration (%V/V)	*Initial adsorption (%W/W)	§Cyclic adsorption (%W/W)	**Steam ratio	Boiling point range (°C)
Methylene chloride	1.0	28.3	17.3	1.4	40.5
Arklone P	0.5	44.9	20.8	1.4	47.6
Acetone	1.0	20.3	12.5	2.3	56
Tetrahydrofuran	0.5	22.0	9.0	2.0	66
Hexane	0.48	21.3	8.2	3.5	68.7
Ethyl acetate	0.5	27.6	13.6	2.1	77.2
Trichlorethylene	0.5	44.6	19.9	1.8	86.7
n-Heptane	0.12	22.4	5.9	4.3	98
Toluene	0.4	23.3	9.6	3.5	110
Methyl isobutyl ketone	0.2	22.0	9.0	3.5	115.9
Mixed rubber solvent	0.3	25.7	10.3	3.8	120-160
Shellsol E	0.3	35.6	11.9	3.7	153-193
3, 3, 5-trimethyl cyclohexyl acetate	0.1	36.4	7.9	4.5	206

*Initial adsorption refers to the solvent adsorbed by new carbon to 'breakpoint', i.e. when solvent can be detected in the effluent air stream. The 'slip' level at this point is normally of the order of 20 ppm.

§Cyclic adsorption refers to the solvent which can be adsorbed and desorbed consistently at an economical steam consumption.

**Steam ratio refers to the total steam used to desorb the solvent from the adsorbent end is an overall figure inclusive of condensate and moisture condensed on the carbon bed.

SUTCLIFFE SPEAKMAN INC

S E N S O R C O N T R O L S

for SOLVENT RECOVERY

**WITH PARTICULAR REFERENCE
TO THE USE OF INFRA-RED GAS ANALYSERS**

Extract from paper prepared by
Dr. Robert D. Hill
Previously of Chemical Plant Division
Sutcliffe Speakman & Co. Ltd.

and

Presented at the 30th Annual Meeting of the
Gravure Technical Association
Chicago, April, 1979

There are many different types of sensors one can use on a solvent recovery plant but all function either to report information and/or to act on that information to control the solvent recovery plant's operation in some way. For example, proximity sensors can be used to sense whether a valve is open or shut, and flow metering and LEL meters can be used in conjunction with each other to measure the total solvent presented to an adsorber. My remarks will concentrate on using sensor control to optimize the conservation of energy, and maximize the efficiency of solvent recovery, and the role of the infra-red gas analyser in helping to achieve these aims.

The first practical step we can take in maximizing solvent recovery plant efficiency is to ensure that we are presenting, safely, as high a solvent load as possible in the minimum airflow passed to the solvent recovery plant. To a large extent this will be governed by the type of process upstream of the solvent recovery plant. Wherever possible, and economically achievable, recirculation of a portion of the solvent laden air should be considered. This will significantly increase the efficiency of solvent recovery across the carbon bed. The examples given below illustrate the typical improvement in efficiency of recovery of solvent which can be achieved:

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- (a) Inlet concentration 1000 ppm
Outlet concentration 35 ppm

$$\eta = \frac{1000-35}{1000} \times 100\% = 96.5\%$$

- (b) Inlet concentration 2000 ppm
Outlet concentration 35 ppm

$$\eta = \frac{2000-35}{2000} \times 100\% = 98.25\%$$

The side benefits from ensuring maximum concentration of solvent vapor that is presented to the solvent recovery plant are that it will reduce the size of the solvent recovery plant required and with it the capital outlay, space needed, and services required to run it. The sensors applicable to this area of plant operation are typically LEL meters coupled to variable damper controls which will vary proportion of solvent laden air from the process which is recirculated and diluted with fresh air.

Let us now consider in some detail the adsorption/desorption cycles which are at the heart of the solvent recovery plant's operation.

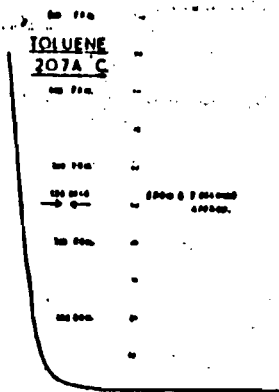


Figure 1

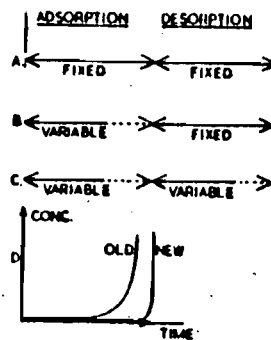


Figure 2

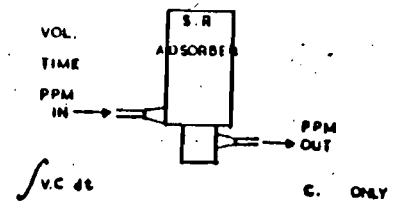


Figure 3

In Figure 1 we see the typical rise in solvent concentration as solvent (in this example - Toluene) starts to "slip" through the bed of activated carbon as an adsorption cycle nears its

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end. It can be seen that there is a rapid rise in solvent concentration - about 100 ppm in around 135 sec. With the particular sensor favored by Sutcliffe Speakman a rise of 5 ppm can readily be detected, and when a pre-selected concentration is reached (usually around 50 - 75 ppm) the sensor produces a signal which is used to initiate an automatic cycle controller which changes over the particular adsorber vessel being monitored from "Adsorb" to "Desorb" mode by opening/closing appropriate valves.

Let us now examine this process more closely. In Figure 2; on Line A, we have an "ideal" situation where a constant concentration of solvent in a constant airflow is presented to the solvent recovery plant. Provided constant conditions are maintained and the activated carbon's adsorptive properties also remain constant, then it is merely a matter of timing the length of the adsorption and steaming cycles to achieve optimum operating efficiency. This ideal situation is rare. What is frequently encountered in practice is that a variable concentration of solvent is presented to the solvent recovery plant. Now, if the adsorption cycle has been time controlled for the full designed load of solvent, and if, for example, half that load has been adsorbed on to the carbon bed in the preset time, not only will the carbon not be utilized fully, but an excessive amount of steam also will be consumed in the desorption cycle. As much as one third of the steam used for desorption could, for example, be wasted.

In Line B of Figure 2 we have the situation where we must be able to sense when the active carbon has adsorbed the optimum amount of solvent, and solvent "slip" is starting to occur through the carbon bed. In this case the length of the adsorbing cycle will be a variable and this is represented by the dotted part of the line in the figure.

However, what will be the situation if, in addition to the solvent concentration varying, we have a process in which there are airborne contaminants with the solvent which cause an unavoidable and progressive deterioration with time of the activated carbon (irrespective of the design or manufacture of the plant or source of carbon)? In some large, multi-adsorber plants associated with such processes it is often the practice to overhaul one adsorber at a time and to replace the poisoned carbon with a charge of new carbon. In this situation one can have an adsorber containing new carbon working in conjunction with one containing old carbon.

On Line D of Figure 2 we see a graphical representation of how the "slip" curves for old and new carbon compare.

In Figure 3 we see the two basic approaches used to measure when an adsorption cycle should be terminated. On the left side

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as solvent laden air enters the adsorber we can measure solvent concentration and flow rate, and compute, by integration with respect to time, the total quantity of solvent passed to the adsorber. When a predetermined amount of solvent has been passed a signal is generated which is used to change over the adsorber from Adsorb to Desorb mode. This approach, as used by some companies, seems to be a fundamentally less reliable approach in that it assumes a given carbon performance at all times. Furthermore, it can never give any useful information about what is happening inside the adsorber.

For example, an unusually high "slip" of solvent could indicate that valve seatings require renewal. It is considered that the fundamentally correct approach is to sense what is happening on the exhaust side of the adsorber and use this information for control purposes. It should be noted that this entails measuring only one parameter (solvent concentration in the air stream) against time. Now, whereas the inlet concentration is typically in the 1000 - 3000 ppm range we are now looking at a 0 - 100 ppm range on the outlet of the adsorber. For optimum efficiency we need a sensor which will be capable of measuring very low ppm's, have a fast response, be stable, be accurate, have high reproducibility, not be affected by water vapor and by other extraneous factors which are frequently encountered in the industrial situation. This tends to exclude the use of LEL meters (too insensitive), gas chromatography (slow response and not continuous), and flame ionization detectors, and brings us to the infra-red gas analyser.

Many gases absorb energy in the infra-red region because of the resonant molecular vibrations within the molecules of gas or solvent vapor. These resonances are highly specific and give rise to I-R spectra which are characteristic of the substances in question. The I-R spectrum of a molecule has been likened to the human finger print by means of which one specific wavelength depends on the concentration of absorbing molecules placed between the infra-red source and the detector, and this fact is utilized to determine the concentration of a selected component in a gas or vapor mixture.

Because of the complexity of typical gaseous infra-red spectra, some overlapping of absorption bands frequently occurs, and it is usually necessary to select a very narrow wavelength band to avoid cross sensitivity. Figure 4 shows the 1-8 microns region of the I-R spectrum of toluene and shows several characteristic absorption bands for this solvent.

There are several types of I-R gas analysers potentially available for use, and broadly they can be divided into single wavelength/single beam; single wavelength/double beam; and double wavelength/single beam instruments. In my experience all two-beam energy absorbing systems suffer from the inherent defect of

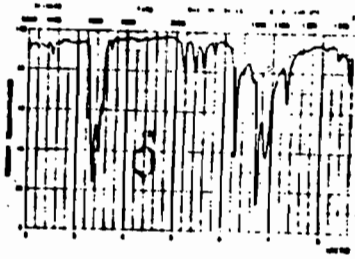


Figure 4

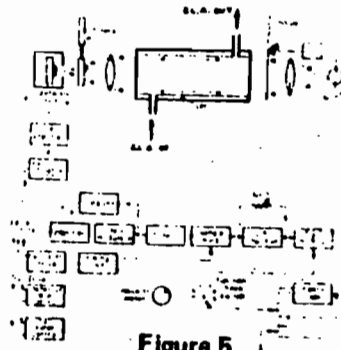


Figure 5

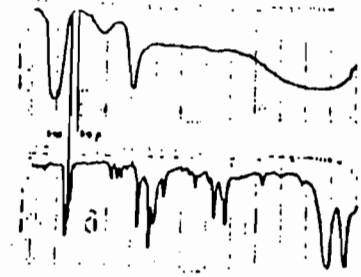


Figure 6

being sensitive to background absorptions caused by fouling of the sample cell windows, broad band absorptions in the sample gas (such as that caused by water vapor), and temperature differences between sample and reference beams. However, most of the disadvantages of the two beam system can be eliminated without losing the advantage of a continuous reference measurement by examining, alternately in time, the I-R energy transmission at the peak absorption wavelength of the measured component and that at a nearby wavelength which represents the "background" energy, but which is itself unaffected by the measured component. This can be achieved with a two wavelength/single beam instrument.

Such a system is shown in Figure 5. By having the source and detector units in optical alignment with each other, and arranging for a pair of very narrow band pass interference filters to be switched alternately in time into the beam (actually at 6 Hz) it is possible to produce a sensitive, accurate, reliable, I-R gas analyser which is immune from many of the extraneous factors which can plague other types of I-R gas analysers.

Let us now turn to the critical area of deciding what wavelengths to select for the solvent(s) in question. This is an area to which, I believe, too little consideration is often given - probably because it calls for expert knowledge in the field of infra-red spectroscopy.

If the solvent is toluene there are, as we can see from the I-R spectrum for this solvent (lower spectrum in Figure 6) several bands to choose from. Or are there? If we look at the spectrum of water (upper spectrum in Figure 6) we can see that there are several regions where water is absorbing energy strongly and has the potential ability to interfere. We must remember also that we are wanting to detect a 50 ppm level (say) which is only a 0.005% v/v concentration whereas if there is a 40% R.H. situation with respect to water vapor that corresponds to a 2.9% v/v concentration of water vapor - i.e. 580 times greater! The possibility of interference from water is obvious. However, if we select wavelengths 3.45 microns for toluene and 3.9 microns for the reference, then we can see that we shall meet the best criteria for having the solvent and reference wavelengths near to each other, but with the latter at a point of least interference from water. In this way we can ensure that even 100% RH will not affect this sensor's performance



Figure 7



Figure 8

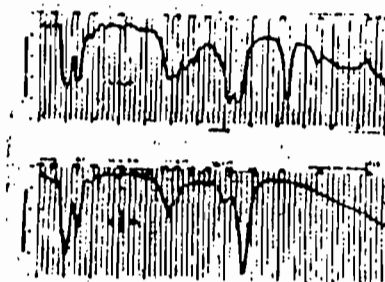


Figure 9

To illustrate just how complex the I-R spectrum of water vapor really is and how the effect of raised temperature can cause the molecules to become more agitated and resonate more strongly, Figure 7 shows the I-R spectrum for water vapor at 100° C, using an expanded scale between 5 and 8 microns. It will readily be appreciated that it is most important that the designer of a solvent recovery plant must have access to such specialized knowledge if the best sensor control design is to be achieved.

Let us now consider some other commonly used solvents: Figure 8 shows the I-R spectra for acetone and methyl ethyl ketone. Here the CH stretching absorption band at around 3.5 microns is relatively weak and it is usual to select the C=O (carbonyl) band at about 5.8 microns instead.

Figure 9 shows the I-R spectra for methyl alcohol and ethyl alcohol. Here one can use the CH band at around 3.5 microns as both alcohols have strong enough absorption bands.



Figure 10

Figure 10 shows the I-R spectra for ethyl acetate and hexane. Here we have a good illustration of the relative weakness of the CH band at 3.5 microns for ethyl acetate and the corresponding strength of the CH band for hexane at the same point in the spectrum.

In conclusion, in order to get the best out of a solvent recovery plant the following objectives should be achieved:

- (1) Maximize the solvent concentration and minimize the airflow within safe/economic limits.

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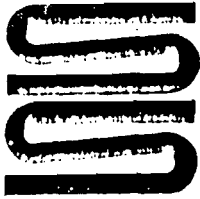
(2) Use I-R Gas Analyser control of the adsorption cycle to nullify the effects of varying solvent load and possibly also varying air flow in order to achieve the fullest use of the charge of active carbon in any particular adsorber.

(3) Select a 2 wavelength/single beam instrument which can guarantee:

- (a) immunity from water vapor
- (b) Virtual immunity from all normal extraneous factors such as voltage variation, temperature change, etc.
- (c) fast and accurate response
- (d) Is fully automatic and requires virtually no attention.

(4) If the carbon is likely to have its activity modified by the process gas/vapor stream, then control the steaming cycle also by sensing a falling concentration of solvent in the steam/solvent mixture coming from an individual adsorber.

As manufacturers of their own activated carbon for the past 75 years plus solvent recovery plant which embraces the sensor technology, I have described, I believe, that Sutcliffe Speakman have one of the best systems for optimizing efficiency and minimizing the use of energy.



SUTCLIFFE SPEAKMAN INC

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YOUR REF:

OUR REF:

DATE: 4 August 1983

DESIGN FOR EXPANSION/RECYCLING

Mr. Michael A. Ware
Environmental Coordinator
Wheelabrator-Frye Incorporated
Graphic Supplies Division
Chemicals and Coatings group
2010 Indiana Street
Racine, Wisconsin 53405

Dear Mr. Ware:

We have pleasure in referring to the discussions with your colleagues and yourself during the visit to David M. Company, Longwood, last Friday, and we appreciated the opportunity to discuss the project for the recovery of toluene from the three rubber spreading machines.

For your preliminary information we have pleasure in enclosing our Preliminary Proposal No. S 7212, in triplicate, for a plant to recover up to 50 gallons per hour of toluene from 25,000 scfm airflow, as you requested, for your initial studies. We understand that consideration is being given to the possibility of making changes to the process operations to reduce the peak evaporation rates of solvent so that the capacity of the recovery plant might be reduced.

The proposed plant is quoted on the basis of supply only, with the addition of a preliminary proposal, Appendix 'A', for the turnkey installation of the plant which we would be pleased to firm up after further study of site conditions.

The proposed plant is very similar to a considerable number of installations that we have supplied for operation with rubber spreading machines in which particular industry we have over 40 years experience.

We will be pleased to quote also for the supply of suitable hoods for the spreading machines and a collection system when you so require.

David M. Company

4 August 1983

We trust that the enclosed information is sufficient for your immediate requirements but we will be very pleased to discuss any aspects further with you at your convenience.

Yours sincerely,



William C. Moses
Vice President

WCM/pmd