STATE OF FLORIDA

DEPARTMENT OF ENVIRONMENTAL RECTUATION
ST. JOHNS RIVER DISTRICT BOB GRAHA GOVERN VICTORIA J. TSCHINK SECRETA SUITE 232 ORLANDO, FLORIDA 32803 SAINT JOHNS RIVER DISTRICT RIVER DISTRICT
APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCES G 24 1983
SOURCE TYPE: Blanket Spreading Operation [x] New [] Existing RAOM
APPLICATION TYPE: [x] Construction [] Operation [] Modification
COMPANY NAME: DAVID "M" COMPANY COUNTY: SEMINOLE
Identify the specific emission point source(s) addressed in this application (i.e. Lime
Kiln No. 4 with Venturi Scrubber; Peaking Unit No. 2, Gas Fired) Festooning and Mixing
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. Furt I agree to maintain and operate the pollution control source and pollution con facilities in such a manner as to comply with the provision of Chapter 403, Flo Statutes, and all the rules and regulations of the department and revisions thereof also understand that a permit, if granted by the department, will be non-transfer and I will promptly notify the department upon sale or legal transfer of the permit establishment. *Attach letter of authorization Signed:
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 been designed/examined by me and found to be in conformity with modern enginee principles applicable to the treatment and disposal of pollutants characterized in permit application. There is reasonable assurance, in my professional judgment,
1 See Florida Administrative Code Rule 17-2.100(57) and (104)
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Nº 74416

RECEIPT FOR APPLICATION FEES AND MISCELLANEOUS REVENUE
Received from Senclair & Valentine Co. Date Cler 10, 1983
Address 201 Valentine Way Longwood Dollars \$ 1,000.00
Source of Revenue David "m" lo Blanket Spunder
Revenue Code 001001 Ck # 1588 Application Number AC59 - 73966
By K Tulloch

PROFESSIONAL 8719	John W. Seabury Neme (Please Type) Seabury-Bottorf Associates, Inc. Company Neme (Please Type) 3702 Silver Star Rd., Orlando, Florida 32808 Mailing Address (Please Type) Oate: August 11,1983 Telephone No. 305/298-0846
,	II: GENERAL PROJECT INFORMATION
end expected improvements in whether the project will resumence.	nt of the project. Refer to pollution control equipment, equipment performence as a result of installation. State ult in full compliance. Attach additional sheet if installation of a lithographic printing blanker manufacture.
	e following sources which are part of a common exhuast eyeteng area, and Mixing area. Please refer to emission points l
n the following process lavo	ut diagrams to ascertain particular emission points and exh
entilation configurations	in this application (Construction Permit Application Only)
Start of Construction upon :	
for individual components/uni	yetem(a): (Note: Show breakdown of estimated costs only its of the project serving pollution control purposes. chall be furnished with the application for operation
Indicate any previous DER per point, including permit issue	rmits, orders and notices associated with the emission

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	1 ° p	ower plant, hrs/yr N.A.; if sessanel, describe:	
			· · · · · · · · · · · · · · · · · · ·
		nis is a new source or major modification, enewer the following quest: or No)	lone.
	1. 1	la this source in a non-attminment area for a particular pollutant?	No
		. If yes, hes "offset" bean applied?	No.
	t	. If yes, has "Lowest Achievable Emission Rate" been applied?	No
	c	. If yes, list non-ettainment poilutants.	No_
		one best eveilable control technology (BACT) apply to this source? If yes, see Section VI.	Yes
		ose the State "Prevention of Significent Deterioristion" (PSD) equirement apply to this source? If yes, see Sections VI and VII.	No
		on "Standards of Performence for New Stationary Sources" (NSPS) opply to this source?	No
:		o "National Emission Standards for Hazardous Air Pollutants" NESHAP) apply to this source?	No
		leasonably Available Control Technology" (RACT) requirements apply is source?	No
		. If yes, for what pollutente? 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 enswer of "Yes". Attach any justification for any enswer 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 Incimprators)

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

Contam	Inants	_ Utilization	·
Type	x wt	Rate - 1be/hr	Relate to Flow Diagram
None Known	-	Variable	Used to formulate cements Mixing (2), Spreading
v.o.c.	30-70%	Variable	Mixing (2), Spreading (3,4,5), Festooning (6
MEK Toluene			(At step #6 99%+ of
		•	off)
1	None Known V.O.C. MEK	None Known - 30-70% MEK	Type X Wt Rete - 1be/hr None Known - Variable V.O.C. 30-70% Variable MEK Variable

*The process utilizes rubber cements which are mixtures of various rubber compounds i.e.(nitrile, e 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.

- 1. Total Process Input Rate (lba/hr): Variable
- 2. Product Weight (lbe/hr): Variable
- C. Airborne Contaminents Emitted: (Information in this teble must be submitted for each emission point, use additional sheets as necessary)

Name of	Emies	ion ¹	Allowed ² Emission Rate per	Alloweble ³	Potent		Relate to Flow
Conteminent	Meximum lbe/hr	Actuel T/yr	Rulo 17-2	lba/hr	lbe/yr	T/yr	Dingram
VOC - MEK/	103	84	BACT	To be determined	420,000	210	1
Toluene	5	4	BACT	To be determined	20,000	10	3
.11	94.5	75.6	BACT	To be determined	378-000	189_	4
11	94.5	75.6	ВАСТ	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.

Paference applicable emission stendards and units (e.g. Rule 17-2.600(5)(b)2. Table II,
E. (1) - 0.1 pounds per million BTU hest Input)

³Calculated from operating rate and applicable standard.

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

D. Control Devices: (See Section V, Item
--

Name and Type (Model & Serial No.)	Conteminant	Efficiency	Range of Particles Size Collected (in microns) (If applicable)	Basis for Efficiency (Section V Item 3)
	1			
		,		
				· · · · · · · · · · · · · · · · · · ·

£. Fuels

	Consumpt	ion*	
Type (Be Specific)	evg/hr	mex./hr	Meximum Heat Input (MMBTU/hr)
	·		

*Units: Netural Gas--MMCF/hr; Fuel Oils--gallons/hr; Cosl, wood, rafuss, other--lbe/hr.

Fuel	Ana:	Lye	i # ;
------	------	-----	--------------

Færgent Sulfur:	·	N.A.			Percent	Ash:	N.A.		
Density:	N.A.			lbs/gel	Typical	Percent	Nitrogen:	N.A.	·
Meat Capacity:		N.A.		BTU/16		N.A.			BTU/gel
Other Fuel Cont	aminen	ts (which	may ca	use eir p	ollution)	ı	N.A.	J	
		(•

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

Annuel Averege N.A. Meximum N.A.

8. 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.

			ft.	Stack Diumete	r:	page)
a Flow Rate:	ACFH_		_DSCFH	Gae Exit Temp	erature:	• F
	1					1 . !
	SECT	TION IV:	INCINERA	TOR INFORMATI	ON	
		Not	Applic	able		, , , , , , , , , , , , , , , , , , ,
ype of Type 0 (Pleatic		Type II (Refuse)		If Type IV (Patholog-ical)		Type VI (Solid By-prod.)
ctual b/hr ciner- ated						
ncon- rolled be/hr)						
		•		•		hr)
al Weight Incine	rsted (1be/h ef Houre of	Operation	per day	Deeign Cep	ecity (lbe/	
proximate Number	rated (lbe/h	Operation	per day	Deeign Cep	ecity (lbe/	hr)
tel Weight Incine proximate Number nufacturer	rated (lbe/h	Operation Heat Re	per dey Mode	Deeign Cep	ecity (lbe/	hr)
tel Weight Incine proximate Number nufacturer	rated (1be/h ef Houre of Volume	Operation Heat Re	per dey Mode	Deeign Cap day/ 1 No	ecity (lbe/	hr)wke/yr
el Weight Incine raximate Number ufecturer e Canetructed	rated (lbe/h ef Houre of Yolume (ft) ³	Operation Heat Re	per dey Mode	Deeign Cap day/ 1 No	ecity (lbe/	hr)wke/yr
el Weight Incine raximate Number ufecturer e Canetructed imary Chamber condary Chamber	rated (lbe/h ef Houre of Volume (ft) ³	Operation Heat Re (BTU)	per day Mode	Deeign Cep day/ 1 No Fuel	acity (lbe/wk	Temperature (*F)
el Weight Incine raximate Number ufecturer e Canetructed imary Chamber condary Chamber	volume (ft)	Heat Re (BTU)	per dey Mode alease /hr)	Deeign Cep day/ 1 No Fuel Type	BTU/hz Stack T	Temperature (*F)
rimery Chamber secondary Chamber seck Height:	Yalume (ft)3	Heat Re (BTU) Stack Dies ACFM	per day Hode slease /hr)	Deeign Cep dey/ l No Fuel Type DSCFM*	BTU/hr Stack T	Temperature (*F)
el Weight Incine raximate Number ufecturer e Canetructed imary Chamber condary Chamber ck Height: Flow Rate:	Volume (ft)3 ft.	Heat Re (BTU) Stack Dies ACFM ign capacied to 50%	Mode Alease Ater: Aty, sub- axcese	Deeign Cep dey/ 1 No Fuel Type DSCFM* mit the emiss	BTU/hr Stack T Velocity:	Temperature (*F) sapFP

H. Emission stack geometry and flow characteristics: From estimates

Emission Point (SCE process floor diagram):

Stack Height: 18 ft.

Stack Dimensions: 4.5×4.5 square (ft.)

Gas Flow Rate: 12,600 ACFM 9960 DSCFM

Cas Exit Temp.: 150 (*F)

Water Vapor Content Ambient (variable)%

Velocity: 600 FPS

Emission Point <3> (See process flow diagram):

Stack Dimensions: 1.5 x 1.5 square (ft.)

Stack Height: 12 ft.
Gas Flow Rate: 1575 ACFM 1410 DSCFM

Gas Exit Temp.: 95 (°F)

Water Vapor Content Ambient (variable)%

Velocity: 700 FPS

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 Water Vapor Content: Ambient (variable)% Velocity: 800 FPS

Gas Exit Temp.: 150 (°F)

Emission Point < (See process flow diagram):

Stack Height: 12.5 ft.

Stack Dimensions: $2(1.33 \times 1.33 \text{ outlets})(ft.)$

Gas Flow Rate: 1415 ACFM 1120 DSCFM

Gas Exit Temp.: 150 (F)

Water Vapor Content: Ambient (variable)% Velocity: 800 FPS

Brief description	or obelectud	, characta		COULTER	0041694	NOE	<u>Appliable</u>	
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	,							
								
								
Sitimate disposal					fran hh		(ac rubbar	4000
	or any arriv	ient other	than that	ewitted	rrom th			*****
ah, etc.):	Not Applicab		than that	981668	TEDM CR			******
ah, etc.):	,		then thet		Trom Cn			
ssh, stc.):	,		then thet	owicced	Trom Cn			

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

 To a construction application, attach basis of amission satisfies (e.g., design calculations, design drawings, participant manufacturer's test date, atc.) 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 parmit from a construction permit shall be indicative of the time at which the test was made.

SEE APPENDIX 1 ATTACHED

- J. 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 begnouse include cloth to air ratio; for acrubber include cross-section sketch, design pressure drop, etc.)
- NOT APPLICABLE

 5. With construction permit epplication, attach derivation of control device(e) afficiency. Include test or design data. Items 2, 3 and 5 should be consistent: actual emissions = potential (1-afficiency).

NOT APPLICABLE

- 6. An 8 1/2" x 11" flow diagram which will, without revealing trade secrete, identify the individual operations and/or proceess. Indicate where raw materials enter, where adiid and liquid waste exit, where gaseous emissions and/or eirborne particles are avaived and where finished products are obtained.
- SEE APPENDIX 2

 7. An 8 1/2" x 11" plot plen showing the location of the establishment, and points of sirborns 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 plen of facility showing the location of menufacturing processes and outlets for mirborns emissions. Relate all flows to the flow diagram.

SEE APPENDIX 3

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- 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

applicable to the source?	
[] Yes [] No	
Contaminant	Rate or Concentration
N . A .	N.A.
	The second secon
	- constant of the constant of
	and the second s
• •	trol technology for this class of sources (If
[] Yes [X] No	
Contaminant	Rate or Concentration
BACT has not been declared for this	class of sources.
200	
. What emission levels do you propose as b	
Contaminant	Rate or Concentration
VOLATILE ORGANIC COMPOUNDS	AVERAGE COATING VOC CONTENT OF 5.4 1bs./
	VOC/gal. OF COATING DELIVERED TO COATING
	APPLICATOR.
. Describe the existing control and treatm	
1. Control Device/System:	2. Operating Principles:
3. Efficiency:*	4. Capital Costs:

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Brier description of					
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	f any efflue	nt other than t	that emitted fr	om the stack (s	scrubber water
	f any efflue	nt other than t	that emitted fr	om the stack (s	scrubber water
Ultimate disposal of ash, etc.):	f any efflue	nt other than t	that emitted fr	om the stack (s	scrubber water

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 stach 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 air-borne 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.

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Useful Life: Operating Costs: Energy Maintenance Costs Emissioner Conteminent Rate or Concentration 10. Stack Perameters Halaht: ACFH flow Rate: Temperatures Velocity: FPS Describe the control and treetment technology eveileble (As many types as applicable use additional pages if necessary). LOW SOLVENT TECHNOLOGY CONTROL VOC EMISSION Control Device: FORMULATE COATING TO b. Operating Principles: BY FORMULATION GIVEN SPECIFICATION Efficiency, 1 NOT APPLICABLE Cepitel Cost: ESTIMATE, \$10,000.00 d. Useful Life: CONTINUING Operating Coat: \$20,000.00 Energy: 2 NOT APPLICABLE Meintenence Cost: NOT AVAILABLE Aveilability of construction materials and process chamicals: AVATLABLE Applicability to manufacturing processes: APPLICABLE Ability to construct with control device, install in available space, and operate within proposed levels: ABLE TO INITIATE CATALYTIC INCINERATION Operating Principles: CATALYZED INCINCERATION Control Device: CATALYTIC INCINERATOR b. Efficiency: 1 90% + Capital Coat: \$500,000.00 d. Useful Life: 10-15 YEARS Operating Coat: NOT AVAILABLE AT THIS WRITIN Energy: 2 NOT AVAILABLE FROM BUDGET h. Maintenance Cost: NOT AVAILABLE Availability of construction materials and process chemicals: AVAILABLE Explain method of determining efficiency. ²Energy to be reported in units of electrical power - KWH design rate.

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- J. Applicability to manufacturing processes: APPLICABLE WITH SOME PROCESS MODIFICATIONS
- k. Ability to construct with control device, install in evaluable epace, and operate within proposed levels: NO FORSEEABLE DIFFICULTIES
- SOLVENT RECOVERY
- . Control Device: CARBON ADSORBER
- b. Operating Principles: PHYSICAL ADSORPTION

c. Efficiency: 1 85-90%

d. Capital Coat:\$600,000.00-\$700,000.00

- . U. oful Life: 10-15 YEARS
- f. Operating Coat: NOT AVAILABLE AT THIS WRITING
- g. Energy: 2 NOT AVAILABLE FROM BUDGET QUOTATION
 - h. Meintenence Cost: NOT AVAILABLE
- i. Aveilability 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
- a. Control Device:

b. Operating Principles:

c. Efficiency: 1

d. Capital Costs:

e. Umeful Life:

. Operating Coet:

a. Energy: 2

- h. Maintenance Coet:
- i. Availability of construction materials and process chemicals:
- Applicability to manufacturing processes:
- k. Ability to construct with control device, install in evailable 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. Cepitel Cost: \$10,000.00

- 4. Ueeful Life: CONTINUING
- 5. Operating Cost: \$20,000.00
- 6. Energy: 2 NOT APPLICABLE
- 7. Maintenance Coat: NOT APPLICABLE
- 8. Menufacturer: NOT APPLICABLE
- 9. Other locations where employed on similar processes: NONE KNOWN
- a. (1) Company: NOT APPLICABLE
- (2) Meiling Address:
- (3) City:

(4) States

¹Explain method of determining efficiency.

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

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Rate or Concentration
(4) State:
Rate or Concentration
on of systems:
nen available. Should this information not
OF SIGNIFICANT DETERIORATION
ABLE
() SO ² Wind epd/dir
,
day year aonth day year
to this application.
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. Wa	e instrumentation EPA referenced or it	e equivalent? [] Yes [] No
b. Was	e instrumentation calibrated in accord	ence with Department procedures?
[]] Yee [] No [] Unknown	
. Heteoro	ological Date Used for Air Quality Hod	eling
	Year(s) of data from / / month day y	
2, Sur	rface data obtained from (location)	
3. Upp	per air (mixing height) data obtained	from (location)
4. Ste	ability wind rose (STAR) data obtained	from (location)
	er Modele Used	
1.		Modified? If yes, attach description.
1	•	Modified? If yes, attach description.
		Modified? If yes, attach description.
3		
4		Hodified? If yes, attach description.
	copies of all final model runs showing the stables.	g input data, receptor locations, and prin-
. Applice	nto Haximum Allowabl's Emission Data	
Polluta		
TSP		grama/aag
50 ²		
	•	grame/sac
	n Data Used in Modeling	
point a		sta required is source name, description of ordinates, stack data, allowable emissions,
. Attach	ell other information supportive to the	ne PSD review.
ble tec		s selected technology versus other applica- oduction, taxes, snergy, etc.). Include a sources.
		al material, reports, publications, jour- on describing the theory and application of

2. Instrumentation, Field and Laboratory

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

24 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 🗘 :

Emission point \bigcirc 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.

Emission Point 1 Total = (spreader (# 1: 77.4 T/yr.) + (spreaders (2 and 3) 504 T/yr. x .25) + (festooning 5 tons/yr.) + (mixing general 2.0 T/yr.) = 210 T/yr.

Emission Point 🔇 :

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 $\stackrel{4}{\checkmark}$ captures 75% of spreader #2's exhaust. Therefore maximum potential emission from emission point $\stackrel{4}{\checkmark}$ would be

252 T/yr. x .75 = 189 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 T/yr. x .75 = 189 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 min./hr. x 4.3 lbs./min x 1.0 = 38.7 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 min./hr. x 14.0 x .25 = 31.5 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 min./hr. x 14.0 x .25 = 31.5 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.

Emission Point 🐠 :

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%

Emission Point 🜖 :

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%

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.

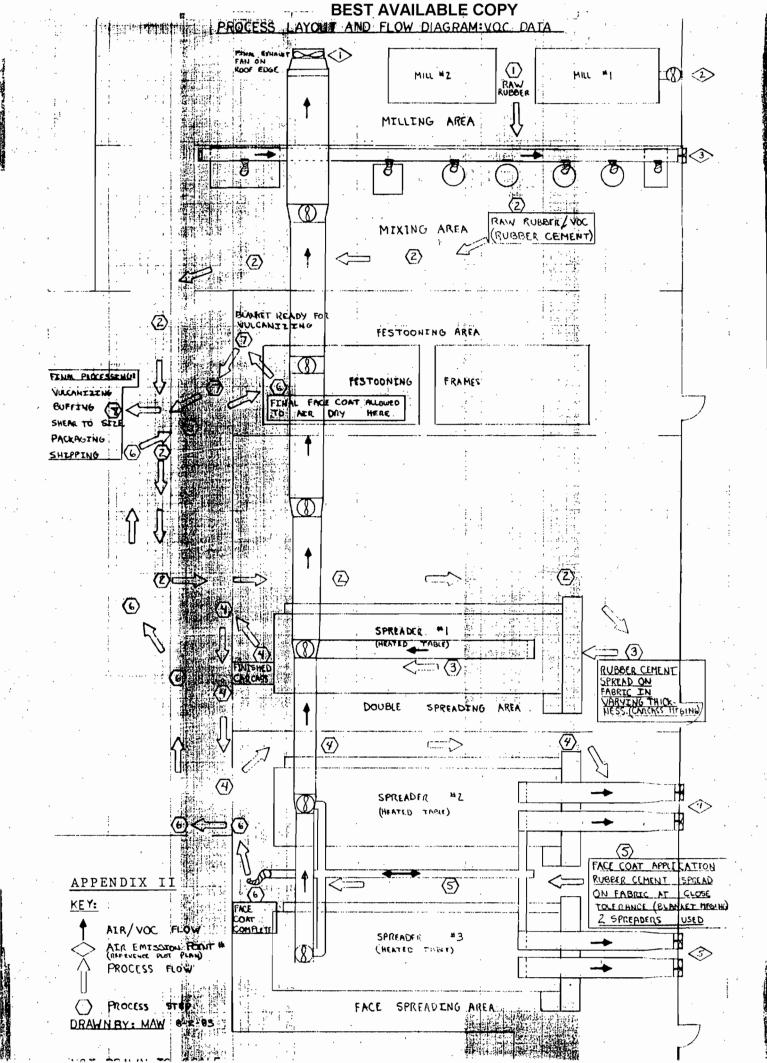
Emission Point	Potential Emissions x	Actual Emissions	
\Diamond	210 T/yr.	84 T/yr.	
\Diamond	10 T/yr.	4 T/yr.	
4	189 T/yr.	75.6 T/yr.	
	189 T/yr.	75.6 T/yr.	TOTAL ACTUAL TONS/YR. = 239.2

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

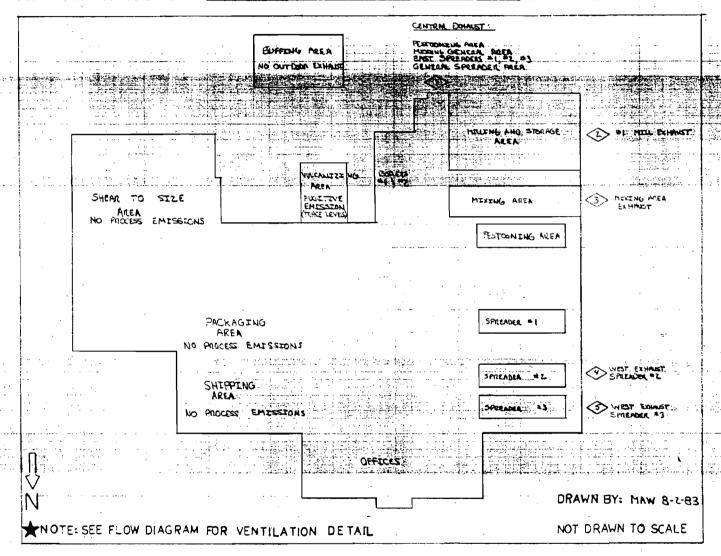
Step 🛈 Rubber is milled into sheets for use in cement making. No calculation of emissions is included in this application since only particulate emissions would be regulated and this source is a minor one which will be dealt with separately from VOC emissions. Rubber is combined with VOC (MEK/Toluene) and mixed to the de-Step sired viscosity for spreading. Rubber cement is spread on cotton fabric in repeated passes of Step (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 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. Following final cure the blanket may be buffed or sheared and Step

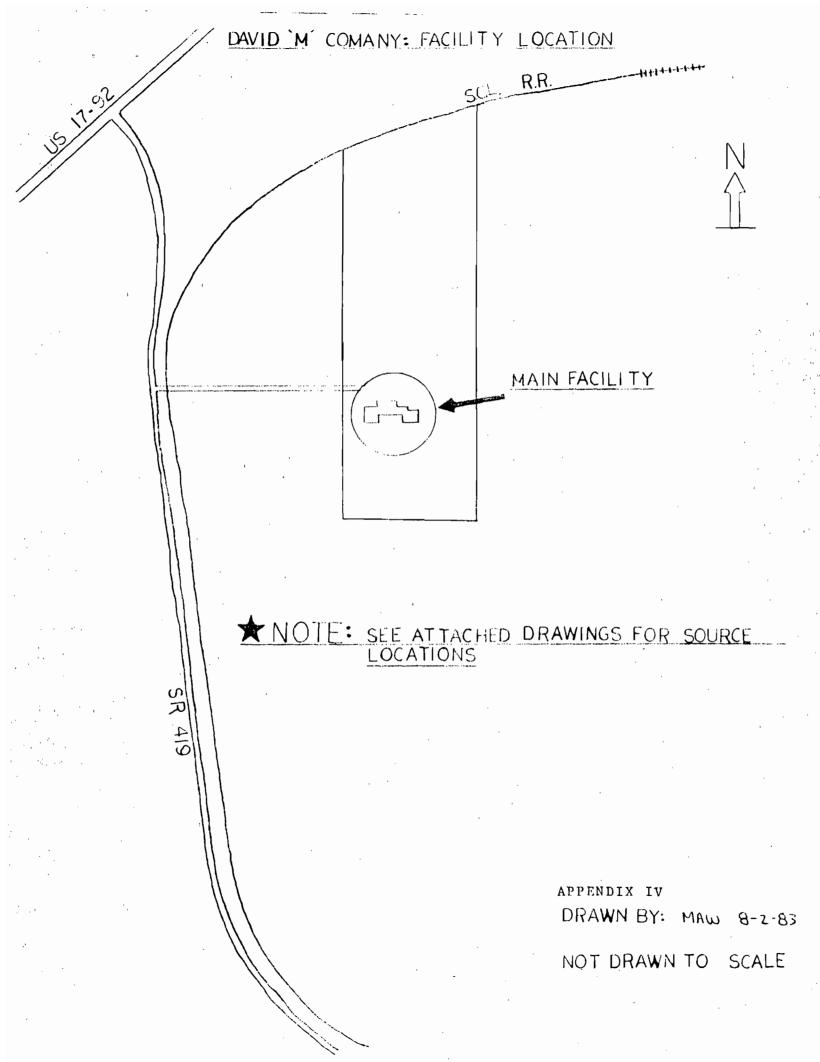
then packaged for shipping.

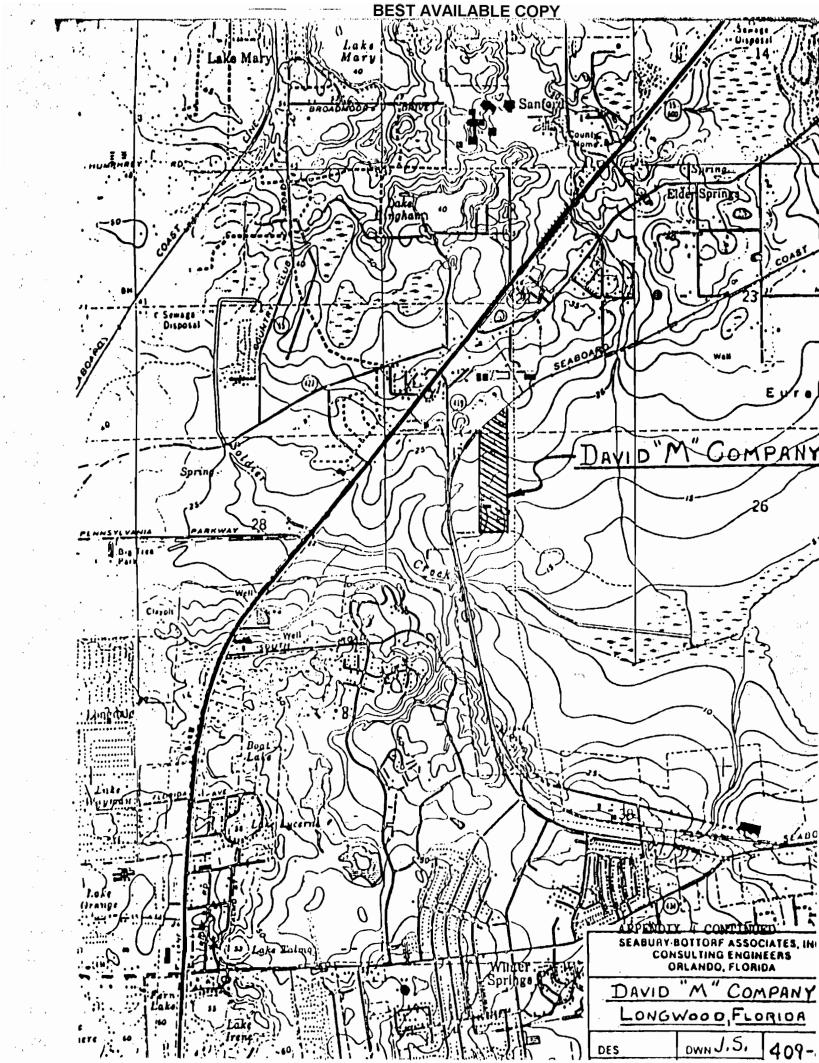


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APPENDIX III PROCESS LAYOUT AND EMISSION POINTS:







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CASE HICHORY SHEET

አ ጀርር አለ ለብ •	David "M" Co/Spreaders 1,2,3 AC59-73966	
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CASE HISTORY SHEET

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DER DEC 21 1982

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

December 17, 1984

1-2-84 Copies to Carol Furbleson & Charles Cellis

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

Page 2. 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

ESTIMATED

OVERALL COMPLIANCE SCHEDULE

DAVID M COMPANY VOC CONTROL PROJECT

PHASE	ORIGINAL DATE	REVISED DATE
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

То	be	fil	ed: <u>\</u>	OIVAC	W	Co
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06	C	Case	Ν,.	84-0	04	†

MEETING DOCUMENTATION

,	Attendees: (Attach list) X
1	Location: TALLAHASSEE Date: 12/12/84 Time: 3.00 PM
9	Subject: Permitting Enforcement Other
l	Meeting requested by: John Gamble, Signal vop Group, (312) 371-2440
-	Meeting Objectives: <u>Submit REVISED APPLICATION FOR PERMIT TO</u> <u>Construct Air Pollution Control Equipment For A lithographic</u> <u>Printing blanket manufacturing facility and new process equipment</u>
1 *	Notes: Notes: Notes: Nobert H. Boddeley, Proj. Eng., (312) 391-3552 (Chief Spekeiman) John H. Gamble, Director, (Contact for questions) HERB MYCROFT, Tech. Dir., David M. Co. Ian Parry, Sutcliffe-Croffshow 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 boilding and pour cement slab for carbon absorption unit—at their rish. Told Co. Preliminary Determination will be out in January, 1985. Told Co. they could not increase production and emissions prior to Told Co. they could not increase production and emissions prior to receiving permit to construct. Agreed that new mixing equipment woult increase production or
	emissions. Explained permitting procedure/schedule. Explained permitting procedure/schedule. Total Co. Final DATE (12/1/85) Reviewed Consent order schedule. Total Consent order. 15 DATE THEY MUST MEET OR REVISE Consent order.

(over)



DAVID M GOMPARY

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

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 mutally 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.

Sin**¢**e**t**e

ayne L. Brady eheral Manager

& Vice President

0157029 $\mathbb{N}_{\mathbb{O}}$.

RECEIPT FOR CERTIFIED MAIL

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

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S Form	SENDER: Complete items. 1, 2, and 3. Add your address in the "RETURN TO" space on reverse.
3611, Jan. 1979	I. The following service is requested (check one.) Show to whom and date delivered
۵.	(CONSULT POSTMASTER FOR FEES)
RETURN RECEIPT, REGISTERED, INSURED AND CERTIFIED M	Mr. Michael A. Ware 201 Valentine Way Longwood, Florida 32750 3. ARTICLE DESCRIPTION: REGISTERED NO. CERTIFIED NO. 0157029 (Always obtain signature of acidressee or agent) 1 have received the article described above. 5:GNATURE DAddressee Daubentzed agent 4. DATE OF DELIVERY 5. ADDRESS (Complete only a processed) 6. UNABLE TO DELIVER BECAUSE: CLERK'S INITIALS
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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:

- 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. Faney, P.E.

Deputy Chief

Bureau of Air Quality
Management

CHF/WH/s

cc: William Darling
Nancy Wright

9-18-84 Cepaies to Charles Collins an Manay Wright

MICHAEL A. WARE Environmental Coordinator



Wheelabrator-Frye Inc. Graphic Supplies Division Chemicals and Coatings Group 2010 Indiana Street



BEST AVAILABLE COPY



DAVID M GOMPANY

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

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 surmisal 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,

Muchael O. Ware

Michael A. Ware

cc: W. Brady

J. Spangler

H. Mycroft

B. Baddeley

T. Westman

J. Gamble

Incompletences
response District
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return for file

Patty









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

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,

DAYID 'M' COMPANY

Wayne/L) Brady

Vace President and General Mgr.

WLB:bjw Attach.

cc: M. A. Ware

R. H. Baddeley

J. Spangler

H. Mycroft

T. G. Westman

J. H. Gamble

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.



JUL 9 1984

CONSENT ORDER

Dept. of Environmental Regulation Office of General Counsel

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 emmission 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 Exibit 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).

FOR THE RESPONDENT:

President and General

Manager David "M" Company 201 Valentine Way Longwood, Florida 32750

DONE AND ORDERED this

1984, in Orlando, Florida.

FILING AND ACKNOWLEDGEMENT

FILED, on this date, pursuant to \$120.52 (9), Florida Statutes, with the designated Department Clerk, receipt of which is hereby acknow-

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

11481463

BOOR PACE SEMINOLE COUNTY FLORIDA

This Warranty Deed Shale and contained the 29th day of September A D 1977 by

HOOVER BALL AND BEARING COMPANY homes at Post Office Box 10033, Ann Arbor, Michigan 48106 and having its principal place of bereinalise culted the granter to

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

becommittee called the grantee

because word horrow his respect groups of conference colleges all the conserves to this engineers and the horrowing and representatives and proposed of conferences, and the conserves and conferences.

. Witnesselh: That the arouter for and in consideration of the sum of \$ 10.00 olupble considerations, receipt inherent is hereby acknowledged by these presents does grant, bargain, sell, alien, remove release concer and confirm unto the granter, all that certain land situate in

Lot 6, SHUMAN'S ADDITION TO EUREKA HAMMOCK, according to the plat thereof as recorded in Plat Book 2, Page 53, of the Public Records of Seminole County, Florida.

Subject to reservations in that certain Deed from the State of Plorida 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: County, Florida;

FOGETHER—with all the tenements, hereditaments and appurtenances thereto belonging or in any-

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

And the grantur hereby covenants with said granter that it is lowfully spixed of said land in for simple: that it has good right and lawful authority to sell and convey said land; that it hereby fully warrants the title ita said land and will defend the same against the lawful claims of all persons whomsoever; and that and land a 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 fayor of Plorida Power Corporation dated May 8, 1973 and recorded June 4, 197] , in Official Records Book 982, Page 695, of the Public Records of Seminole County, Florida.

In Witness Whereof the granter has coused these presents to be executed in its name and its corporate seal to be because allized, by its proper officers thereunto duly authorized, the day and year first above written.

HOOVER BALL AND BEARING COMPANY

ATATE OF Michigan COUNTY OF Washrenaw

I HEREBY CERTIFY that on this day before me, an officer dule so Robert E. Ressler and Douglas E. Peck, respect

Wice Francis and Secretary,

had and allered and in the Courty and Story inc almost the 29th to, or September

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The Instrument prepared by:

October 13, 1933

PHASE

Data Accusulation

Spreader Enclosure - (1) For Temperature Evaluation

Process Modifications and Development

Methods Compartson/Analysis - Thermal Incineration

Final Specification Development

> Final Specification/Quotations and Analysis

Appropriations Request Development and Approval

Oven-Enclosure and Exhaust System Installation

Solvenc Recovery Delivery and Installation

Operational Shakedown, Debugging and Compliance Tescing

January 1, 1984

January 15, 1934

April 1, 1934

May 15, 1984

July 1, 1934

September 1. 1984

December 1, 1984

March 1, 1985

August 1. 1985

D.E.R. - ORLANDO EXHIBIT NO. 2

Best Available Copy Nº 81922 DEPARTMENT OF ENVIRONMENTAL REGULATION



DAVID M GOMPANY

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 fours,

DAVIZ M COMBANY

Wayne J. Brady Vice President & General Manager

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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 spreaders, 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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TATE	SELEN	. <i>"Э</i> Ч1 Д 7Н		SCRIF	PTROM	, WOLL	LBS	#INS	SO YDS	CEN	TOLUENE WSED	SYD	# !LYD	TOL	FEET Adgrawin	F/FEET	ি হৈছে একিল ভা
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1/13/83	120	80.00	771	PASS	., .	70.5	14.0	4.7	266.66	.052	9.870	.037	.082	2.100	76.59	,,,,	
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1/13/83		TOTALS	TALC P	ASS		70.5	23.0	16.9	266.66	-006	743-571	9 60	-135		21.30	21.30	
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DTAL PR	000CT	ION TIME	06:16:2	0	Est of behavior	enskamien vien	MA	13.4	was a survivor		WAX	164	365	9.270	# 63.63		e e e e e e e e e e e e e e e e e e e
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-UN DN 0	1/23/	84 AT 14	:11:55			DAVID	TOLUE	NE AMALY	212					****	PAGE 22	* : · ·
DATE	LEN	₩ Æ DTH	RE	SCRIPTION	*. Tu	LBS	MINS	SO YDS	CE M	TOLUENE USED	TOL SYD	TOL	TOL	FEET	F/FEET	
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		-					."						100 mg	(A) ***		
					>>>				CEM	TOLUENE	TOL	TOL	TOL	FEET	F/FEET	•
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	83 : 湖 次		"3RD .!	PASS 💥	_ ₹256.8	3 2 3 . (D	222.22	-058	7.384	- D33	~ 5073	1.893	30.92	हरू 6.92	
11/02/			RUESER	70072 1517	PASS 62.7	13.0	5.6	222.22	-058	8.151	-036	-081	1.455	53.57	~~~~53.57	H. H. Fasts
	8 3 <i>``@'</i> J/00		ZND .t	PASS 16	4 14 14 MARIND Z • 1	233.I	3.5	- FIZZZ - ZZ	:: D58	720 D. 151	₩ 50	1806	THE STORY	90.90	1230.90	State of the state
11/02/	83 <i>X</i> 00	80.00	3901	PASS	62.7	32.0	D 4.2	222.22	-144	20.064	-090	-200	4.777	71.42	71.42	
31/02/	8 3 47 /2 0 0) ::::::::::::::::::::::::::::::::::::	L NGH ★TH N	PASSa 🦠 🔧	○ (源) 62.7	/	0 #75-1	₹222•22	-112	T-15-675	· 070	- E- 156		源 58 - 82	32256 ⋅ 82	100
11/02/	83 100	, 80.00	9 51H 1	PA 5:3	62.7	7. (5.1	222.22	0.51	9.389	-019	- 043	-860	58.82	58.82	
*1/02/	83季₹100	170 D.00		PA 55, Lie 1787	1 62. T	# 35D. (-225	学经51.350	受影 141	75-513	5-598	汉元 3.57	表 3.57	The W
11/02/					62.7		4.6	555.55	-180	25.080	-112	-250		65.21		
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1/02/	83 100	80.00	13TH F	PASS	62.7	28.0	2.7	225.55	-126	17-556	-0/9	.175	6.502	111.11	. 11111	
	B3 🐃 00		STOTE T	PA :ES	2.7	425.	D 1151, 2- B	··· 222-22	-112		TO	350	500	WW7-19	- AD7-19	
1/02/				T IST PASS				222.22				-156		78.94		
		3380.00			PASS 764-2		3-1	225.22			-063	- W-1-1	5556			1775
1/02/				PASS	64.2	23.0	2.6	223.55	-103	14.786	-066	-147	5.273	707-14	707-14	
1/02/		50.00		PASS .	\$5° TARRA-2			. ZZZ •ZZ	-157	22.470		- Se 22-	-1129,321	57.69		
1/02/				P 4:55	64.2	19.0	3.5					-121	3.485	85.71	85.71	
1/02/				PASS	- A-2		, , , , , , , , , , , , , , , , , , ,	222-22			- CAA-	~211	3.745			
1/02/			\		64.2					17-976						
1/02/	83 300	33300.00	TIN .	P433	THE WITH \$4 - 2	33.7.2 (·		222-22	-970	T. C. D. 1			365 4.708	220-00	120.00	
1/02/	82 300	80.00	TO CO	2455	D	22.	3.0	222.22	- 000	14.124	.063		7.708	100.00	100.00	
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	grand the major	U U I A L S	reas increases and a second	THE THIRD BY THE STORM	THE STREET, SHOWING THE STREET		103.0	~~// /-/ 2	-114	414.441		CACAGO IN BACTORISM		101 T. 36		
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DIAL	PASSES	. 20				(IIII	N 2.5			# I M	-049	- 109	3.755			return a service
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RUN IN 01/23/84 AT 14	:11:24	DIAID W LOFAEN	NE ANALYSIS	And the second s	PAGE 17
Beredika (1997) Budak Jakisa (1997)	and a little and a companies of the	Parties	CEM TOLUENE TO	TOL TOL FEET	F/FEET
POATE , WLEN DEWEIDTH	DESCRIPTION YOU	LBS MINS	SO YDS SYD USED ST	D MALAD GANIN SOUTHIN	of MIN Charges
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1/01/83 195 65.00 1/01/83 295 765.00	RUBBER 106 1ST PASS 70.		352.07 .181 45.055 -12 352.07 .207 51.392 -14	7 .231 2.489 32.32	26.26
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DATE	LEN.		Total De	SCRI	PTION					CE W	IULUENE	106	TOL	T (DL FEET		t materials Later
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FOTAL PAS	SES	25	et gattav engre OSIAAIR	ال ان فمشكات الاس	WAY T	er ere dage		18 -W2-1	TOWN WE	超50%-15	MIN HAX AV6	178	- 397	7 - 5	06 - 34.4	0 100-00	
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RUN ON	01/23/	'Æ4 AT 14	:11:23	·	D	MAYID. W	70LUE	NE ANALY	SIS			~			PAGE 15	
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" DATE	LD	! . ₩IDTH	DESCRI	PTION	TIPL	LB5	MINS	SC VDS	SYD	. USED	SYD	LYD	Promomin	-MIN	MIW.	
				•						•						
10/27/8	3 : [3] 00	**************************************	TOPHER 3 49	2TIST PASS	37.2:	25.0	#Q5.3	*222.22	SIES	**************************************	- D64	125 SET 43	# FP 4508	**************************************		The setting and and
10/27/8	3 100	80.00	2ND PASS	5	57.2	18.0	4.2	222.22	-061	10-296	-046	-102	2.451	71.42	71-47	ACC. 11 12 12 12 12 12 12 12 12 12 12 12 12
10/27/6	3 3/2 00	50.00	" 3RD PASS	the second of	57-2	13.0	# 13 O		-058	7.436	-033	074	1750YB -806		76 . 02	
10/27/8	3 100	80.00	RUPBER 700	2 IST PASS	58.5	15.0	4.3	222.22	- 067	8.775	-030	-087				
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10/27/8			JOD DASS		58.5	35-0	12.3	222.22				204	1.464	24 30	24 30	The state of the s
10/27/8				ingering suprement in a great	58.5	27.0	: . 4. 3	222.22	121	415.795	-071	× = 157	· ************************************	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	-amag - 76	1:
10/27/8			5TH PASS	TROPHICAL AL	58.5	24.0		222.22		14.040	- 063	140	4.011	85 71	P.S. 71	
10/27/8		7 0 . OO	STH PASS	から 東京 ない でき は 100 100 100 100 100 100 100 100 100 1	58.5	20.0	7472-5	********	2000	1.700	052	7072 117	.680	550.00	æ 30 - 0 0	• • • • • • • • • • • • • • • • • • • •
10/27/8		80.00	7TH PASS	New Spirit Con Str. Spirit	58-5	19.0	3.0	222.22	085	11.115	.050	.111	2 - 850	76.92	76.92	1
10/27/8	3 7928 00	38.02 DQ	MINNATH DASS	A 45 FERRIS THE STEWNS HOSE	~a.~	1 3 2 - D	:23 5	100000	- 144		70.000	was 1 8 7		45 - 71		
10/27/8	3 7 00	80-00	OTH PASS		58.5	26.0	5 2	222.22	117	15.210	068	152	2 025	57-50	57-50	The state of the s
10/27/8	3 48882 80	45.00 D 3.00	THE PASS		50.35 1	TORREST OF THE	TERRE - 10	WW 25 . 23	*****	0.04	THE PARTY OF	THE CANADA	OOO A STREET	70.00 FA . 02	**********************	LINE COURS - No. 1907 - LINE -
10/27/8	3 100	80.00	11TH PASS	A MANAGEMENT AND A STATE OF THE AND	58.5	25.0	3.1	222.22	1112	14.625	· 065	.146	4.717	06.77	96.77	Maria and Maria
						7777 3 - 0	145 MD L A	14 19 2 2 2 2 2 2	- 05.0	AD ACTIONS	- C - C - C - C - C - C - C - C - C - C					AND THE PROPERTY OF THE PROPER
10/27/8	3 100	80.00	13TH DACC	THE PERSON OF TH	58.5	32.0	1 . 0	222.22	- 144	18-720	- 084	187	10.400	166-64	166.66	A A CHARLES AND ASSESSMENT OF THE PARTY OF T
10/27/8			THAT HE DASS	The state of the s	58.5	- 34 - N	~. 302	3222-22	- 100	THE PLAN	₩ . 063	T0%_ 1 A D	- 10.400 - 10.400	CA 15	*142-85	
10/27/8			15TH PASS	w. remark and a series of the series of	58. S	22.0	2.0	222.22	- 000	12.87C	.057	.128	6.435			
B/27/8			STATH BASS	,	58.5	10.0	∵.3. 6			₹% 5.850	.3-026 ∂\$0-€	0 58			70 - QA	TRANSPORT OF THE
10/27/8			DUBBED AAC	O IST DASS	65.6	10.0	3.4			6.560	.029	.065	1.929	88.23	88.23	र विकास कर के किया है। जा स
20/27/8				9 IST PASS	45.	7.17.0	TEN 1	222.22					**************************************		73.17	
0/27/8			3RD PASS		6 5.6	16.0	3.2			10.496	.C47	.104	3.280	93.75	93.75	
3 27/8			SATE DAGE	المان المان المان ال	5 5.6	18.0		222.22		711-808	-053		723	. 420 - 00	-420-00	The second section is the second seco
-0/27/e				Jim 152. 124. 144.	65.6	15.0	2.5 2.1	222.22		5.240	.044	3006	4.685	142.85	142 B5	
:0/27/8			WATER BASE	ing a community of the	45.6			222 - 22	*- 300		39 P4		SHOOTING WOOD	71-42	FW: 71-42	
10/27/8			7TH PASS	a hara a sanada kamada kalenda da d	65.6	25.C	3.1	255.55	113	16.400	073	154	5.290	96.77	71-42 96.77	'
:0/27/8			THE DASS	manyar manya mpanyaran	85.8	23.0		222.22			-067	150		-414-11		TO THE PROPERTY OF A STATE OF THE PARTY OF T
0/27/6			DIN PASS	orgo reguesta. Orgo reguesta	45.4	18.0	4.1	222.22		11.608	.053	118	2.880	73.17	73.17	
-0/27/8			MANTH BASS	the second of the second	55.4			. 525 - 55			056		.T793		2115.36	
0/27/8			TITH PASS		<u> </u>	16.0	1.9	222.22		15.496	.047	-104	5.524		157.69	
B/27/B			12TH DISC		55.6	15.5	~2. i			10-168	-045	-101	15 7214 - 841		442.05	
0/27/8			TATH DASS		65.6	14.5	1.7	222.22		9.512	.042	.095	5.595			
0/27/8			TATH BASS		45.4	28.0	3.1	222.22			-082	-183	15.925	96.77	96.77	
0/27/8			ISTH DASS	• •	65.6	27.5	2.7	222.22		18.043	. 381	.180	6.661		111.11	<u>.</u>
	3 3100		WASTN DASS		45.6	20.5	2.5	222.22		#13.44B	-060	134				
0/27/8			17TH DACE		45.6	9.5	1.9			6.232	.028	-062	3.280		157.89	
0/27/8			TRIM PASS		55.6	9.5		222.22		36-232	-028	-062	2-047	142-85		
0/27/8			101H PASS		45.4	8.5	2.5	222.22		5.576	.025	.055	2.230	77120.00	142.85	مسائع دي .
	3 mg 200		WEZOTH PASS	A granten of the contract of the	45.4 -		:-2.5	.555.55		7-544	033	- 075	1377792-901		115.38	* ************************************
	3 100		WEDIN PASS	, S	65.6	12.5	1.8	222.22		8.200	• 035	.082	4.555		166.66	
	3 1100				55.4 ·	12.5					.033				166.66	
	3 100		23PD PASS	e	65.6	14.0	3.1	222.22		9.184	.041	-091	्राक्ष्य • 191 2 • 962	96.77		
	3 .781 00					77.13.0	2.5			8.528	.036	085		20.00		
3/27/8			DETH. DACE		45 4	14.5	1 4	222 22		9.184	.041	. 291	5.740			
0/27/8			26TH DACE		45 -6		72 / 4	222.22		6.856	.039	088	:5.535			
0/27/8			2714 0455		45.4	11 6	2 0	222.22		7.544	.033	.075	3.772			
0/27/8			SATH DASS		65.4	. 11.0	1 - 6	222.22		7.216	-032	-072	- 4 . 00 B			
3/27/8			2010 0455		45.4	11.0	2.3	222.22			.032	.072	3.137			
	3 100		TAIC DASS		45.4	7 70 - 5	15.3	222.22		6.232	.028	. 062			19.60	
U/ 2 1/ 0	3 100	707416	TALL PASS	e e e e e e e e e e e e e e e e e e e	UJ•U	007.0	150-7	666.55			.020	, 0002	5401			
	5.1	- OTALS	·	. 10	Same of the party	201.0	150. /	. 000.00	* .	302.733				erication in	1. 2.2	- T
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DYAL		ION TIME	00 241 240			= 1	12.3			-A Y	-105	. 234	10.400		187.50	
VEDACE	1 85 6	FMENT DE	D DACS IA .	195		70/	3.1		*	AV.	.052	.116	4.148		112.29	
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IUN DN	01/23/	84 ÅT 14	: 11:22			DAYLD P	TOLUE	NE ANALY	S 1 S	•		- 7		!	PAGE 14	T
2. DATE	LEN	WIDTH	_ 10	ESCRIPTION	7 TOL	LBS	WINS	SO YDS	CEM SYD	TOLUENE USED	TOL	TOL	TOL	FEET	FIFEET	₩. ^
0/26/8 10/26/8	100	80.00 80.00	240 380	1492 451 Pass Pass	PASS 57.	20.0	2.5	222.22	.090	11.440	.051	.114	5.576	120.00	120.00	
10/26/6	3 100 3 1100	80.00 #813.00 80.00	RUEBE)	₹ 7002 IST	PASS 58.	5	10.4	222.22	.288 .211	37.440 27.495 45.045	.168 	.374	3.600	28.84 28.84	28.84	
0/26/8	3 100 100	ક્કું . 00 . 0 0 20 . 0 53	• 4TH 5TH	PASS	56. 58.	33.0	5.3	222.22	-045 -148	5.850 19.305	•026 •086	.193	103. Parisa 1.290	756.60 66.66	65.56	
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0/26/8	100	00.00	RUBB E	PASS 1151	PASS 69.	53.0	6.2	222.22 222.22	.238	36.868	.165	.368	5.949	48.38	48.38	
0/26/8 0/26/8 0/26/8	3 III 00 1	80.00 28 80.00	i kirir • TH	PASS 🚉 🏗	69.6 69.6	27.0	202. 7	222.22	-121	-848.792	## 084	**************************************	6.960 6.286	we41.11	-411-11	- Line of the second
0/26/8:	100	60.00	67H 7⊺H	PASS	69.6 69.6	15.0	∞ 2.3 3.3	222.22	.081 .085	13.224	• 956 •059	.132	4.007	90.90	70.90	Contraction of the Contraction
0/26/83	1 00	80.00	ÇTH	PASS	69.6	15.0	2.7	222.22	- 067	15.440	.046	.104	3.866	111.11	111.11	
DTAL PA	SSES	22 TIME	05:44:2 05:44:2	**************************************	a allandaria Taraharia		# xg.1 x 10.4	All the Con-) - / . AT	WAX	- 2026 - 202	958 150	8.190	28.64 142.85	142.85	T. TOP
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RUN DN	01/23/	84 AT 14	:11122	4	٠	DAVID M	70LUE	NE ANALY	515	•		, 190 9	(grande i minde jagen. Grande i samer i	PAGE 13	7
DATE	LEN		I DE	SCUIPTION	TOL	LBS	MINS	50 YDS	CE M SYD	TOLUENE USED	TOL SYD	TOL	TOL	FEET	F/FEET	
30/25/ 10/25/	83 100	8 (I ~ C ()	E ZND I		57.2	14.0	4.3	222.22	.063	8.008	-036	-080	1.862	69.76	69.76	a the same of the same of the same of
10/25/ 10/25/	E3 100	60°00) 00°00	RUBBER	7002 1ST PASS	58.2		7.2	222.22	.198	25.608	.115	• 256	3.556	41.66	1.56	
10/25/		150.00			58.2 58.2	49.0	· · · · · · · · · · · · · · · · · · ·		-216	27.936 28.518	125 128	· 285	5.820	61.22	61.22	
10/25/01	3 7 1 00 8 3 1 00	80.00	'작성 4TH (5TH (PASS HAMM SUSSIFIED	58. 2	33.0 51.0	3.3	222.22	.148				で表表 - 820 6・745	92290.90		
	B3 🚉 100		ME BTH	PASS		39.0		222.22	÷175	22.698	102		-829	÷3.82		7200
				PASS NEW TENERS	୍ୟ 5 8∙2	-∷737.0	38 3 S	, 222.22	×- 166	31.21.534		- 215	152			
10/25/	100	्यू म 80.00	FRUDSER	PASS	78 69. 0	24.0	· 7	~222.22.	# 108		:::::::::::::::::::::::::::::::::::::	1835 165	29.33 ·523		. 63.62	
10/25/	83 🗯 🗷 🚥	<i>孫</i> 称780.00	. : : : : : : : : : : : : : : : : : : :	PASS 作。不是这种对此的	<i>∷</i> ₹ 369. 0	:::::::::::::::::::::::::::::::::::::	- Sept		≫-081	1024-20	ব্যক্তা চেত্ৰ	top- 4 24	ापुरुक ः 17 5		≈#25 . 99	The Control of the Co
10/25/	3 100	60.00	5TH 1	PASS Yester	269.0	19.0	~2.1 ⊘2.0	222.22	35085	12.420	- 958	· //李131	5.914 ************************************	150.00	142.85 	- Alleria
30/25/0 4 0/25/ 0	100	. BO.00	। साठ । सार िहु≝्।	PASS Light The	0.9.0 0.986;;;;	16.0 77% 30. 0	1.8	222.22	•072	11.040 20.700	049	-110 207⊈e	0.133	73-17	166.66	
±0/25/0 ₹ 6/25/ 0		80.00		PASS PASS	69.0	28.0	2.0	222-22	-170	19.320	- 050	. 193	7.430	175.38	115.38	,
-0/25/		80.00	' 101 - 1	PASS TO THE PASS	69.0	15.0	2.2	222.22	.067	10.350	- 046	103	4.7D4	136.36	136.36	
10/25/1	83 1 0 C		1274 (PASS PASS	69.0	14.0	3.6	222.22	• 06 3	9.560	. 94.3	-070	2.003	63.33	63.33	A STATE OF THE STA
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DTAL	PASSES	ICH TIME	06:37:39	5 28.115 VARD 3.289		MA	x 13.1			WAX	204	.453	11.072	166.66	165.66	<u>-</u>
VERAGI	E L es c E Les C	EWENT PE EMENT PE	F PASS ? F SOULRE	28.115 TARD 3.289	: •	AV	6 3.9			AVG		. Pà 1 PA	~~-669	74.14	~ 94.14	Apple of Comments
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 I dn o	1/23/6	34 AT 14	: 11 722				DAVID P	TOLUE	NE ANALY	515	. TO HENE			And the Samuel States of the Sales	इत्या क्यांच्याम् १५	PAGE 12	t garantari
						, TÔL						100	105	TOL	FEE	FIFEE	
24/83		800	2ND	PA55		57.2	29.0	7.2	222.27	133	16.588	-074	- 165	2.303	41.60	41.56	
24/83 24/83	1 120	(£(J) = 00 (€(J) = 00		P 7002		55 59.6	95.0	7.2	222.22	• 202	26.910	- 121	- 269	3.737	41.00	41.00)
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		80.00		PASS A	- 10	<u>;;;;;::35</u> 6.2		3.3.	7-222-22	25 198	25.116 25.608 20.952	115	₹6.256	· 27.760	2.5EP0 - 90	100 PO	150
24/83	77 00 EV7	OO . OO	THE BEST H	PA55	F	30.2	27.0	3.4	* 222 - 22	35121	7135.714		∵:::::•157	. MEET - 5021	300 . 2 :		4.4
24/83		80.00	0 I H	PASS		5 6 4 2	25.0	2.6	222.22		14.550 12.222 12.222	-065	- 145	5.506	115.36	115-38	
24/83 24/83	100	80.00	RUBBEI RUBBEI	PASS 188	IST PA	\$\$ 58.2	21.0	3.5	222.22	.094	12.222	· 054	-122	3.492	85.71	85.71	11 12 13
24/83	100	80.00	350	PASS		56.2 SS.∰85.4	44.0	2.9	222.22	198	25.608 25.608	-115	.256	8.830	103.44	103.44	ENGRAPTION OF
24/83	100	80.00	240	PASS		65.4	29.0	4.0	222-22	-130	18.766	.085	-189	4.741	75.00	75.00	
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PAGE I	Les CE	MENT PE	R SQUAF	E YARD	3.199	. fr	ра ферена — 1961—	-/		- Pit-V	TO SERVICE TO THE	B 022	THE STREET THE RESE	angang separat se	. , ,		· As an annual · ·
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				1.4	11.5	700				•••							: ::::::::::::::::::::::::::::::::::::
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						\$.,		• • •		• • •				A Supplement	
39/27/83	₹ 00 E	- T71. 016	TRANSF	P/SPATE	R TEST	PASS TOT.	5 36.0	- 133 A	5213.88	-16B	1300 × 300	4"WE 113	·***********	4,37,5147	****** 23	2000 - Q2	
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		77.000	THE PARTY	TW/SS	40	**************************************	5 46.5		213.88		41.137	- 052	- 111	11 mm2 = 855	- A6.92	#### 03	127 Table 1
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19/21/83	100	77.00	1711			~ M ~	/ /A - ()	2.7	213.66	• 112	19.000	- 4000					
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	1 00 T	77.00			- 114 J	58.	7 ::32.0		213.88		388.784		267	3.080 5.680	96.77	93.14	man and
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	100	77.00		PASS	43.	58.	7 21.0	2.6	213.88	- 050	11.153	.052	111	3.28¢	115.38	711.05	Section 1985
9/27/83		77.00	- 101 20 Th	20 SS	なかずが デー・**	Maria de la companya	7 33.0		213.88		74819.371	707-090	10 T	-739\$ -457			PROTESTINATED .
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9/27/83		77.00	26 T M	PASS	্ ক্রিক্ট	TOTAL THE PROPERTY AND A SECOND	~326.0	3.1	213.88				725151		96.77		
9/27/83	100	77.00	271 H	CASS	and display	75.00 (156.05) 58.00	29.0	2.5	213.68		16.935	.079	169	5.774	120.00		
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9/27/83		77.00	. TOTATH	PASS		68.	2	5.4	213-88		- 21.142		211	7.356	· 88.23		
	100	77.00	5TH	PASS		68.	26.0	2.4	213.88		17.732	- 882	.177	7.386	125.00	120.31	
	100	77.00	6TH	PASS			2 a 41.0		213.88		27.962	-130	279	3.531	55.55	53-47	काई इत्यक्षिक प्रभावित ।
	100	77.00	7T H	PASS				~~~5.6	~213.68	-135	19.778	.092	• 1 7 /	3.531	53.57	51.50	
9/27/83	~ ≈1 00 :	ge: 77.00	aryrapp o I H	PASS	33 BVF.	elibries en	0 - 89mm S	··· 3.3	- 213-68	-130	** <u>19.99</u> 6	· · · • • • • • • • • • • • • • • • • •	490 Perior	·····································	~ 90 .90	:: <u>∵</u> 67-60	and the sales attached
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9/27/83		77.00		PASS	214 - 48.	66.	2 700.0	3.9	213.88		9 :32.73 6	° a~153	327	-393	76.92		
9/27/83	100	77.00		FASS		66.	2 45.0	3.6	213.89		27.280	-127	-272	7.577	63.33		
9/27/83	3.00	77.00		PASS	7.75	65.	2 - 31-0		213-88		21.142	098	-:L211		-90.90	767.50	- अपूर्ण स्थिति ।
	100	77.00		DASS		65.	28.0	3.2	213.85		19.096	-089	-190	5.967	93.75		art t
9/27/83		77.00		PASS		68.	2 38.0	3.8	213.88		25.916		259	<u>6.820</u>			•
	165	77.00	IETH	PASS		68.	2 37.C	3.1	213.68		25.234	• 11 /	-252	8.140			
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	1 00	77.65	1718	F455		66.	10.5	3.3	213.65			-033	.071	2.170			
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DATE	LEN.	"WIDTH	27 € 2.89	SCRAP	TIDN	¥DŁ	LBS	MINS	SO YDS	CEM	TOLUENE USED		TOL			F/FEET	
7/26/83	**** ****	78.50			1ST PAS						24.300			3.626		45€9 3 . 05	Control of Section
7/26/83 7/26/83	98 98	78.50		PASS	,~ • •	67.				-098	14.175	.066	-144	3.081	63.9	62.71	
7/26/83	98	78.50		7002	IST PAS	D /•	5 14. 0 2 28.0		213.68		-9.450 16.856		• 172	3.180	55.47	7 54.43	
/26/83					AST PAS				~213.68		19.866		::-202	7 75 BAD	22.47	₩ 84 · 84	
1/26/83	98	78.50		PASS	5 1 M (1 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2 1 1 2				213.68		27.090	.126	.276	A - 752	21.2	, 50.61	
/26/83		3478.50			"X\$"	13850.	2 37.0		213.68				-227	5.180	≈0 2750 - 3 7	7.00	The state of the s
/26/83	98	78.50	5 T H	PASS		60.	2 33.0	3.3	213.68		19.866	.092	.202	6.020	89.09	, 6	
		· 50			الأمامي المعاطفة الذي المراشعة المعادلة المامية المعادلة المامية المعادلة المامية المستقلمات المامية المستقلمة المعادلة المامية المام				213-68	-229	:::::::29.49B	130	. 301	7. P. 218		7 -90-15	7.72
/26/83	98	78.50		PASS		60.	2 24.0	4.5	213.68	-112	14.448	.067	.147	3.210	65.33	5 5 - 10	
/26/83		测解8.50	SERBIH	PASS -		77 760 .	2 .7855.5	店熟2 ●●	. 3213.6 8	259	333.411	.#; ≈156		深州13.921	93822 - 50	.420.50	THE STREET
/26/83	98	78.50	OTH	PASS	& THE THE WAY HAVE THE WORLD	60.	2 31-5	5.6	213.65	.147	18.963	-086	-193	3.386	52.50	51.51	
726/83	- 60% 96 - 60%	78.50	1114	PASS	St. The Local Activities	.00±;©∪•.	2 33.530.0	30 Sept. 4	::::::::::::::::::::::::::::::::::::	-100		AND TOI	100 th 201	変な形で・2/4	3.55	3 04.04	CHECK VALUE
					CONTRACTOR TO BE NO COLUM				213.68	m 133	17.157		.175	3.812	65 - 33		Art Birchister of the Art
/26/83	OR	78.5C	TTH.	PASS	CONTRACTOR DAY	- 0 0	2 10.0	2.4	213.68	-088	11.438	- 053	-116	4.765	122.50	120.20	
/26/83		78.50	TOTH	PASS	· · · · · · · · · · · · · · · · · · ·	350.	2		213.68	-102	11.438	##=061			-917-60	115.59	
/26/83	96	78.50	RUBBER	7002	1ST PAS!	5 58.	7 21.5	2.5	213.65	-100	12.520	.059	.128	5.04E	117.60	115.39	
/26/83	98	78.50						** 2.7			EE0.033		156 122				mamilia a galladak sud
/26/83	95	78.50	300	DASS		5A.	7 10-0	2.7	213.65		11.153	.052	-113	4.130	105.88	106.84	
/26/83	98	78.50	ATH	PASS	": " . "	58.	7. \$\text{\$1.7.0}	2.3	213.68	.079	7. 9.979	*** • D4 6	£101	**************************************			
/25/83	96	78.50	5 T P	PASS		5e.	7 17.C	2.3	213.68		9.079	.046	-101	4.338	127-02		
/26/83	. 98		STH.	PASS .	<u> </u>	56.			213.69		45.262	+071	· 155	359			
/26/63	98	78.53			IST PASS				213.68		12.198	- 057	-124	4.691	113.07		
/26/83	- 98	778.5C		PASS							***********	- 854	25117	-200	76.00		
/26/93	98	78.40		PASS		64.			213.58		17.976	-084	.1E3	5.992			
/26/83 /26/83	/: 798 98	78.5C	STH.	PASS	· · · · · · · · · · · · · · · · · · ·	64.			213.68 33.68		13.488	- 969 -063		4.953	100.00		
/26/83	. 98	T78.50		PASS :	ر سودند پ	64.			∌213.68		731-556	.:: 054	-117	1393 984			
/26/83	98	76.50		PASS	:	64.			213.68		12.940	.060	.131	4.612	91.87		. • •
V26/83	~^ 9e	78.50		PASS					213.68			-057	-124	5.062			1.777
/25/83	99	78.50		PASS	• •	64.			213.65		11.556	- 054	.117	4.615	122.50		
V26/83	98	78.51	LOTH	PASS	n 4.51	5.4			213.68		10.272	.048	~ 104	"" - 466	127.62	125.42	• • • •
/26/83	98	78.50	11TH	PASS		64.	2 16.0	2.4	213.69	.076	10.272	-048	- I 04	4.280			
/26/83	:: 98	78. 5∦				64.					10.272	-048	-164	:::#3 - 8 ○ ◆	108-86		and the second of the second
/26/83	39	78.50	13TH	PASS		64.			213.68		16.914	.051	-111	*4.745	127.82		
/26/83	~ 9 8	78.50	1479	PASS ,	ادار المعادية الأدارية. مراوعيا الأمطاعة أدارية المار	64 . i	2 . 236.0	2.5	213.68		30-272		104	-106	117-60		
/26/83	98	78.50	TALE P	A 5 5		64.	2 18.0	15.2	213.69	-084	11.556	. 054	• 1 1 /	. 100	17.5		
											-571.110		21491452	Martin and Albertail Co.	7 M 2 - 7		ए क्स १ भेड़ तायह
TAL PAS	SES	37	- make 2000	ener we	Section 1	- 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ent i mana na i	N .02.3	100	+ 2 3 Feb.	NI Milwight Control	-044	165 096	2.250	-43.B		
7 BBS	DUCT	ON TIME	A7.67.4		1" (7,)			. * = = =		64. A	· · · · · · · · · · · · · · · · · · ·	.156	.340	13.921	127.62	125.42	

SON ON O	1/23/	84 AT 14	:11:22			DAVID P	TOLUE	NE ANALY	S 1 S				يونه ي طونشيند - ي	And Age	PAGE 8	
DATE		. "WEDTM			*	· LBS	MINS	SO YDS		TOLUENE USED	TOL 570	TOL	TOL		F/FEET	
39/23/83	-100		FAMESER 3149	2 15T #PASS		77A1 34 25 1	1 TOTAL - 129 -		7	~7 19.9 41				67.40		·
19/23/83		811.75	IND PASS		57.E					12.427		124	3.030	73-17	73.85	ACTION AND ADDRESS OF THE PARTY.
19/23/83		"02.75				16.0	. 23.4	224.30		9.248		092	~~ ~ 2.720	mr48.23	10000 A Q . 06	
39/23/83	100	FB5.7'5	RUBBER 700			26.0	5.8	224.30	.115	15.652		- 156	2.0070	J E	26161	
39/23/83		10.75	2ND PASS	₹ <u>.</u>	60.2		7144 6 - B	224.30		31.906		26319	9-692	27744 - 11	~~~ -44. 53	
19/23/83		80.75	3RD PASS		60.2	38.0	4.5	224.30		22.876	-101	-228	2-041	00-00	h/~/V	
		では あり、75 りの、75	TH PASS				3.3	224-30		19.866		198		705.00	77 P. 10	THE SECTION
19/23/83		##80.75	TESTHEPASS		6 D. 2	30.0	2.9	224.30		18.060		180	0.221	103.44	102	
9/23/83		80.75	7TH PASS	1 THE	60.2	33.0	5 - 1	224.30	-147	19.866	.088	108	3.895	38.82	50.37	
		3880.75	BTH PASS							**************************************	ાલ ≂ 13 5	7000mm304	~~~10.133	- 00 a 00	#78400 - 94	The state of the s
9/23/83	1 00	80.75	9TH PASS		60.2	25.0	4.0	224.30	-111	15.050	-067	150	3.762	75.00	75.70	A No. of the last
9/23/83	33 DO		THE OTH PASS		(2) 0.2	题 有51.0 .	。第2. 7	温224-30	-136	3548.662	083	186	AND 06-911	Bat 11-11	· 3412-15	
9/23/83		80.75	11TH PASS		60.2	32.0	2.2	224.30	-142	19.264	.085	-192	6.642	103.44	104.42	
9/23/83	. 767 DO	80.75	13TH PASS	一种的一种,在一种	**********	- Care Care Care Care Care Care Care Care	MANAS - 2	#224 - 30	- 176	23.779	∞e-106	237	- 10 - 338	(100 mg)	94.63	Special section of the second section of the section of the second section of the section of
9/23/83 9/23/83		3380.75	JESTATH PASS		60.2	20.0	3.2	224.30		12.040	.053 %=045	.120	3.702 924	93.75		
9/23/83	100	80.75	15TH PASS	researcher in regulation in	60.2	11.0	- 3.5 3.7	224.30		6.522	-029	.066	1.789			
9/23/83		# 80.7 5	TESTH PASS	¥***	60.2	10.5	2.0	224.30		7576.321	026		14:912 -257			CARLETTER & AR.
9/23/83		80.75	17TH PASS		6C.2	11.5	2.2	224.30		6.723	.030	-069	3.146	136.36		en en Bestelle de
9/23/83	1 00	80.75	AUSBER 660	9 457 PASS		*34.0	5.7	224.30		23.290	-103	232	-085	-52.63	53.12	
9/23/83	100	80.75	2ND PASS		66.5	55.0	5.9	224.35	. 245	37.575	-167	~~°. 376	~~~B.385	50.84	51.32	• •
9/23/83		60.75	3RD PASS	Her. 45	, :68-5	35.0	2.9	224-30		23.675					104.42	
9/23/83	1 00	80.75	ATH PASS		68.5	30.0	2.6	224 -30		20.550	-091	- Z05	7.903		116.40	
9 /23/83 9/23/83		80.75 60.75	TO STH PASS	· · · · · · · · · · · · · · · · · · ·	68.5	29.C	72.9	224.30		79.160 IS.165	- 085	198	5 :850		704.42	4.62
9/23/83		********	TTH PASS		7.68.5	54.0		224.36		36.996			525 -8 -941		65-83	18 19 18 DI
9/23/E3	100	80.75	BTH PASS		68.5	37.0	3.1	224.35		25.345	.112	.253	8.175	96.77	97.68	The second second
9/23/83		80.75	OTH PASS		66.5	29.0		224.30		19.865			2006-019			; ss*
9/23/83	1 00	80.75	TOTH PASS		66.5	55.0	4.9	224.35		37.675	.167	. 37€	7.688	61.22	61.80	•
9/23/83		88.75	TAITH PASS		68.5	◆3.0	3.6	224.36		29.455	-131		wgg:-8 - 181			* *** *
9/23/83	1 00	80.75	"12TH PASS		68.5	20.0	3.0	224.30		13.700	. De 1	137	• .566			• •
9/23/83		80.75	TITH PASS		68.5	26.0	5.5	224-30		19.186	-985		ger ~ 8 · 718			
9/23/83	1 00	60.75	14TH PASS		68.5	28.0	2.4	224.30		19.180 9.590	.085 -042	.191	7.991 5.047	125.00		والمواسيوس معولوسوسي
9/23/83 9/23/83		80.75 80.75	15TH PASS		∵ ∙68.5 68.5	15.C	1.9	224.30		10.950		.109	5.768			To the same of the
9/23/83		80.75	17TH PASS				2.3	224.30		e.905			.xee 3.671			
		10 mm		ger y tare you and a single	68.5	19.0			·	· · · · · · · · · · · · · · · · · · ·	.058	.130	3.827	88.23		***
9/23/83	च <u>.</u> 00	93 ~ 60.75	TALC PASS	and the second second	₩68.5		11.6	m224.30	-09 B	15.070			299 benerala			Section 1
		TOTALS	TALC PASS OB:10:04 R PASS 30.3 R SOUARE YAR		- 0.0	1155.0	136-2	2523.4D	.135	740.543	•	•	ind the .	CONTRACT CONTRACT	4 * **	,
					in a second	10 Mg - 1 (122 mg - 1		د مد مست دیک پیوا		SAC MVX Alk	_1_	** ** ***	7000	:		
DTAL PA	5 SE 5	38		•		MIN	1.5		•	WIN	-028	.063	1.789	94 - II		
JIAL PR	DOUCT	ION WINE	08:10:04			MAX	6.8	•		TAX.	-167	- 376	866.04. 5.663		159.37 94.53	
TERAGE	L 25 ()	CHENT PE	H MADD JÜ.J	99 D.B. 140		▲ V G	, 3.5			# A P	-086	• 1 4	3.663	93.03	*~.33	,
/ERAUE	LOS C	CHENI PE		D 30179								•				

RUN ON	01/23/	84 AT 14	:11:22		DAVID .	TOLUE	NE ANALY	S I S			i.s.			PAGE 7	
				2				CEW	TOLUENE	TOL	TOL	TOL	FEET	F/FEET	. **
DATE	LEX	MIDTH	DESCRIPTION	TOL	LBS	MINS	SO YDS	SYD	USED	SYD	:LYD	MIN	MIN	#1W	
99/22/0									20.808						Total Appen.
39/22/0		81.75 B1.7 5		57.6 57.8	26.0 18.0	**************************************	227.08 2 27.0 8		15.028	• 066	• 150	3.130	62.50	63.86	
09/22/8					27.0	3. 8	227.08		15.498	.068	. 154	4.078	78.94	80.67	C- C/88%
09/22/6				57-4	30.0	₹ - 5	227.08			33-075	172	3.826			Table 1
39/22/6			3RD PASS	57.4	41.5	4.3	227.08		23.821	-104	.238	5.539	69.76	71.29	Killian Market Profess 1.
19/22/6					. 36.0	3.4	227.08			7 096	218			3200-16	- E
)9/22/0				57.4	33.5	3.2	227.08		19.229	.084	192	6.009	93.75	75.80	
19/22/6		61.75		57.4	**************************************	3.3	227.08		22.960	· 101	.229	6.957	90.90	92.90	THE REST
19/22/						ം ഷ 0	227.08					-59m -696			TARREST AND
19/22/8		81.75	9TH PASS	57.4	45.5	3.0	227.08		26.117		.261	6.696	76.92	78.60	ALLAN CONTRACTOR OF
39/22/6				3657-4	W40.5	72.7			323.247	WEE 102	232	1088 - 610	3111-11	-213-54	THE SERVER OF
19/22/6			IITH PASS	57.4	36.C	3.7			20.664	.090	-206	5.584	81.08	82.85	**
19/22/8							2227.08		ு ் இ. 368			₩₩5.925	₩ 996 - 7 7	**************************************	
19/22/6					58.5	6.1	227.08		31.982	-140	. 318	5.226	49.18	50.25	
3 9/22/6 39/22/6		81.75	يريا «ZND PASS» الميران SRD PASS	54.5			-227.0B		25.070	-099	<u>₩#-250</u> • 226	6.112	73.17 81.08	*74.77 82.85	
39/22/6			ATH PASS		41.5 29.0	3.7	227.08		22.617	. 6.00 at 19.		***** E 1 &	T -	. 87.59	THE STATE OF THE S
39/22/6					44.6	5.1	227.08		29.612	.130	296	5.806	58.82	60.11	
39/22/8				67.3		· 4- 8	227.08				TA 235	5.888	75.00	76.64	Tributa Stra
19/22/8	3 100	81.75	3PD PASS	67.3	36.0	4.3	227.08		20.190	. 586	.201	4.695	69.76	71.29	, · ·
19/22/8		81.75			21.6	3-0	227.08		13.818	18-060	-t-was 1 36		200.00	-145-10	
9/22/8		81.75	24D PASS	65.8	26.5	3.9	227.05		18.753	.082	.187	4.935	78.94	E0.67	
39/22/6		3.75	SRD PASS	65-8	14-0	2.7	227.08		9-212	X,# 04 0	<i>ે</i> લ≁ 9 92	727F2-411	141-11	*113·S	
19/22/8		81.75	ATH PASS	65.8	51.5	5.5	227.05		33.887	-149		6.161	54.54	55.74 74.77	*
3 9/22/ 0		81.75	6TH PASS	8.65.0 mgg/55 5.50	35.C	3.8	227.08		28.952 23.030	- 127 • 101	289 230	6.060	78.94	80.67	Company of the second of the s
19/22/8		8:.75		्र (१९४ - १९ ७५ - ६	17.0	4.3	227.08		* 8711-186	33-049	776211	TER2-601	69.76	71-29	***
19/22/6		e 75	TALE PASS	65.5	24.0	17.2			15.792	-069	.157	.928	17.64	16.03	
. 77 227 0		TOTALS		: 100 J			6565.32		\$13.257						e i je u upagim in kranjange en 🗒
OTAL F	ASSES	29	and analysis	The state of the s	· WIN	2.7	a comp		THE PRINCIPAL IN	* ***	15 09 2	-2.040	48.36	49.44	
DTAL	RODUCT	ICH TIME	C7:42:55	2	MAX				神食 X	-149	. 33e	8.610	111.11	113.54	-
VERAGE	LES C	EMENT PE	R PASS - 35.413 🛷	、地 小田上楼	- A ¥6	-4.1	: .		-AVG	~ - 09Z	115		76.94	~76.62	the first papers of the same and the
VERAGE	Les C	EMENT PE	F SOUARE YARD 4.52	22			, .		•	•			**		

STOTOL CHANGED

FROM 57.4% TO

54.5% CN THIS

PASS

RUN ON C	11231	84 AT 14	11:22				DAVID M	TOLUE	NE ANALY	SIS	٠.			The street street	, , , , , , , , , , , , , , , , , , , ,	PAGE 6	3. 73 °
				÷	' .	•	*			CEN	TOLUENE	LAS TOL	TOL	ŤOL	FEET	F/FEET	
DATE	LEN	WIDT	H DE	SCRIPTI	DTMI	TÖL	LBS	MINS	. 50 YDS	SYD	DESED	SYD	*LYD	1995 SM 1 M	Tomb		្រែក្រោយ
99/21/83	- TE 00	7746.D	"RUBBER	1402 13	PPASS	557.B	20.0	*:55 8.25	F3 27.77	·***	221.560	56.090		.778	_XE 46 -15	· 26299	S Name of the last
09/21/83	100	46.0	D ZND	PASS		57.8	11.5		127.77		6.647	.052	.066	1.621	73.17	42.07	4 1 10 3 3 1 1 1 1
09/21/83						57.0	** B . O		9127.77			~ ~036	∵00 0 4 6	2.296	77 60 -10	39.20	TO THE REAL PROPERTY.
09/21/83				7002 15			18.0	4.5	127.77		10.332	-080	.103	2.296			
09/21/83					.s	57.4					33.12.628	098	126	· 建建物 - 354		59.48	二两里更是二
09/21/83					~~;		18.0 13.0	3.0 4.2 ∴	127.77		10.332	-058	-103	2.444 004 Essentar	100.00		
09/21/83				7002 15			42.0	5.0	127.77		24.108	.188	.241	821	60.00	34.50	
09/21/83					. گانگاد است به						47.462		₩ D74				
09/21/83	100	46.00	3 PD	PASS		57.4	9.0	4.0	127.77	.070	5.166	. 04D	.051	1.291	75.00	43.12	C. S. All AND MANAGEMENT
09/21/83				PASS 🗯 🚉	1	r; `57 . 4		₹.2	4127.77	-086	## #6.314		~~063	10 - 670	FE 36 . 36	77078 - 40	**************************************
09/21/83						57.4	9.0		127.77			.040	.051	1.519		20012	
99/21783			H GERRELL C	PASS 金田光		57.4					314.					90.78	
09/21/83 09/21/8 3		46.00	D. SEETH.	PASS MINI	consider to	3/44 4 35	9.0	7.0784 3 .2	127.77	155	5.166	.040 	.051	2.718 499983-567			-
09/21/83		46.00	914	DASC MANN	[4] [1][2][2][2][2][2][2][2][2][2][2][2][2][2]	57.4	12.0	2.8	127.77	-003	6.888	.053	.068	2.460	107.14	61.60	. man indian
99/21/83				PASS: "T"	يو دود دود دود درستا الدا الداملية المحمد	57.4					- 25 BBB	# <u>%</u> ÷ 053		∰###5.444		·*************************************	- the transfer of the contract
99/21/63	100	46.00	1114	PASS		57.4	11.0	2.3	127.77	.096	6.314	.049	. 063	2.745	130.43	75.00	· · · · · · · · · · · · · · · · · · ·
79/21/83				PASS 🥂		~57.4	9.0		9127.77			·- 040			150-00	26.25	The state of the s
96/51/83				D A < C		57.4	10.0	2.0	127.77		5.740	. D44	. 057	2.870	150.00	86.25	
19/21/83				PASS		₹57. 4	7.0	- 1.9	127.77			~~:-031	D+D	Z2-119	157.09		
)9/21/83		46.00		DA 55	,	57.4	8.0	1.8	127.77		4.592	.035	. 045	2.551	156.66	95.83	
)9/21/83		46. 00		PA 75		57.4	10.0 14.0	2.9	127.77		332	-062	103	20 6. 63.63	135.43	75.00	TREE
39/21/83				D4 675			13.0	2.0	127.77		7-462	-056		3.731	150-00	- 66 . 25	
19/21/63		46.00		7:00 2 10	DACC	57.4	12.0	2.0	127.77		6.889	.053	. D68	3.444	150.00	~ 85.25	
39/21/83						57.0	12.0	~ 2.0	127.77		6.888	-053		3.444	150.00		STEEN PROVIDE
39/21/83		46.C		PA 55		57.4	9.0	2.6	127.77		5.166	.540	-051	1.985	115.38	66.34	
79/21/83				6e07 19	T PASS			%-1	127.77			094	121	2.954	73-17	42-07	
)9/21/E3		46.00				67.3	12.C	2.7	127.77		e.076	.063	. sec	2.991	111-11	63.88	
39/21/83						67.3	13.0	2.4	127.77		8.749	.066	-967	:3-645	125-00	71.67 49.28	- Table
19/21/83 39/21/83		46.00				67.3	.26.0	3.5	127.77		17.498	.136 -152	174	4.000	是。71 185。71	49.28	
19/21/83		46.00				67.3		2.9	127.77		9.422	.073		3.248	103.44	59.48	
9/21/83						67.3	6.0	1.9	127.77			-031	75 94 9	2.125		90.78	-2. <u>4.</u>
9/21/83		46.00				0/.3	10.0	2.4	127.77		6.730	.052	-067	2.804	125.00	71.87	
39/21/03		46.DI		PASS "	•	67.3	10.0	2.2	127.77		6.730	per - 052	- D67	3.059 2.557	136.36	78.40	- en 3
9/21/83	100	46.00	D 10TH	PASS		67.3	9.5	2.5	127.77		6.393	.050	• 55 5			69.00	
9/21/83			D ** * 11784 *	PASS 11/7 11	en cy	~~67.3	9.0	3.1	127.77			-047	- 960			55.64	e can em data da e
·6/21/83			1214	PASS		67.3	4.5	1.0	127.77			-023	-030	3.028	300.00	172.50	
-9/21/83	100		TALC P	PASS MAIN PASS ASS M		467.3	8.0	14.9	127.77			:- D42	· • 95 3	-361	20.13	11.21	
		TOTALS					542.5	112.6	5110.60	.106	327.491						in the section of
STAL PA	SSEC	4 C					VIN	1.0			WIN	.023	.530	1.050	46.15	26.53	, " 1"
TITAL DE	DOLL	ION TIME	87:42:0	•				6.5			MAX	-108	- 291	5.576			. 2
VERAGE	LESC	EMENT PE	PASS	13.562			AVG	2.6			AVG	.063	. 56 :	2.892		67.54	
VERAGE	LBS C	EMENT PE	E 87:42:0 ER PASS ER SQUARE	YARD 4	245								· -			ś	

PUN ON 01/23/84 AT 14:11:22	DAVID	M TOLUENE ANALYSIS		WORLD TO THE	PAGE 5
·	IPTION TOL LBS	CEM	TOLUENE TOL ""	TOL FEET LAD STREET	F/FEET
19/20/83 97 77.75 RUBBER 14 19/20/83 97 77.75 2ND PAS 19/20/83 97 377.75 3RD PAS	92 (1ST PASS 7.8 29.5 S 57.6 21.0	5 209.49 140 C 4.4 209.49 .100	12.138 .057	175 2.795 66.13	64.27
9/20/83 97 77.75 RUBBER 70 19/20/83 97 77.75 2ND PAS	02 157 PASS 57.4 51.6 S 7 200 57.4 2049.6	0 編4.4 (209.49 %233	9.537	096	58.91 58.91
9/20/83 997 977.75 94TH PAS	5 1-1321451 157.4 1726.6 5 57.4 44.6	3.1 209.49 124 0 2.6 209.49 210	25.256 .120 .	153	108:77
9/20/83 97 77.75 7TH PAS	5 57.4 32.5	209.49 2057	18.655 .089	108	117.83 - 12 300 V.
9/20/83 ~@97 ~@977.75 @@901H PAS	\$ 18.7 PASS 18.7 32.5	5 4.1 209.49 112	6.077 .029	2.870 97.00 189::::::::::::::::::::::::::::::::::::	68.97
9/20/83 97 77.75 2ND PAS 9/20/83 97 77.75 3RD PAS 9/20/83 97 77.75 ATH PAS	S 67.4 2€.0 S 1. 2000 / 1267.4 安治1.0	0 4.0 209.49 .133 0 %43.6 %299.49 -100	18.872 .090 .	194 4.718 72.75 195	70.70
9/20/83 197 177.75 11/5TH PAS 9/20/83 97 77.75 6TH PAS 9/20/83 197 77.75 7TH PAS	\$ 15.77 167.4 1733.5 \$ 67.4 26.0	5 */ 3.1 209.49 % 159 0 4.3 209.49 % 124	17.524 .083	232	
9/20/82 97 77-75 8TH PAS 9/20/83 97 97-75 9TH PAS	67.4 15.5 67.4 10.7 7.6	2.3 200.47 .673 5 .2.5 200.49 683	10.447 .049 44.1.795 .056	107 4.542 126.52	122.96
TOTALS		5 E7.6 5027.76 .130	386.521	121 46-93	3945-64
DTAL PASSES 24 TIAL PRODUCTION THE 06:53:11 VERAGE LES CEVENT PER PASS 27. VERAGE LES CEMENT PER SQUARE VA	#1 	IN 2.3	2 461 - 139	062 1.882 87.70 301 9.713 126.52	422-96
VERAGE LES CEMENT PER SQUARE VA				State Control of the	
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RUN ON OI	/23/	84 AT 14	:11:55		ı	DAVID P	TOHYE	NE ANALY	S 1 S	V			en grunnag Program Wr	عبيبار د د	PAGÉ 4	
DATE	"LEN				TOL	LBS	V	SO VDS	CE M	TOLUENE		TOL	TOL	FEET	F/FEET	ki manan
89/19/83	, °75	~~80.50	"RUBBER 1492	1ST PASS		₹35.0	5.7	212.4 2	5164	7739-355	25091		- 3-200	₩ 98.3 0	18:50	
09/19/83	95	80 5C	2ND PASS		55.3	22.0	4.3	212.42	•103	12.166	.057	-128	2.829	66.27	66.69	
09/19/83 09/19/83	95			167 0466	355 • 3			212-42			_::-039	35087	2-2-1	77-02	77450	artistical des
709/19/83	95 95	80.50 30.50		TOL PASS		34.5 351.0		212.42			.096	.215		6.72		
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09/19/83	95	80.50	5TH PASS		59.3	33.5	4.1				.093	-209	4.845	69.51		事法 シャン
D9/19/83	··95			5 100 to						27.574						**********
09/19/83	95	80.50	7TH PASS		59.3	4 C . C	4.5	212.42	-188	23.720	-111	-249	5.271	63.33	63.72	
309/19/83	95			ETTER TRE	:59.3	9.0	Chr. 5.9	£ 212.42	:-230	海29.057	्राह्म ः १ ३६	305	-450	13.07 mg/d	∙হল্লে⊉3.63	38.32
09/19/83	95	80.50	AIH PASS		54.3	4 / • U	W Zel	, 212.92	• 2 2 1	2/08/1	• 1 3 1	• 293	W 0 . /Y/	.\ 0A.21	09.94	471
999/19/83	795	表表80.50	音法 10TH PASS 頭	्ट अस्ति सहस्रा ध	359·3	3391.03	16-2-10	15212-42	193	AUX24-313	- Table 1	255	20141-577	×4.35.71	· 3436 · 55	
09/19/83	95	80.50			59.3			212.42			-097		6.695			
3 89/19/83 09/19/83	्राक्ष ्ट्र 95			TOT DACE	83Y • J	. 用源之 / · D · I	*** ** • • • •	## 212- 12	-129	- SEE - 20 /			, 100 mm	109.61		- Suchan S.
*89/19/83	• "95		2ND PASS	131 PASS	C.V.J	19.5		212.42			.054	.121	770,539 -269		110.29	
09/19/83	95	80.50	3FD PASS		59.3	16.5	2 5	212.42	A77	9.784	.046	-102	3.913	114.00		
109/19/83	::195			PAGE TOE							117		v:::5.657		e 65.17	
09/19/83	95	80.50			69.2			212.42		16.027	.075	.168		X 114.00	114.70	ati I
09/19/83	-95	80.50	"BRD PASS	TO JOHN HE	68-2	27.5		212.42		28.755	-088	₩ 58 97		× 135.71	136.55	
09/19/83	95	80.50	ATH PASS		66.2	23.0		212.42		15.556	.073	. 165	6.274	X 114.00		10
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09/19/83		80.50			66 · S	21.0	2.5	212.42		14.322	· 067	-150	4-93E	/ 98.27	96.88	
.09/19/83	* 95	- 80-50		Parina and Charles Control			3-0	212.42		21.142	-099		7-047		-95.59	CONTRACTOR
09/19/83	95	80.50			66.2	18.5		212.42		12.617	.059	-132	5.040	7 114.00	114.70	
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AVERAGE L	BS C	EMENT PE	R PASS 29.758			AVE	3.5		3 2 3	AVE		- 294	5.582	<u>.:</u> . 68 . 62	69.37	10 Te 10 19
AVERAGE L	.85 C	EMENT PE	06:47:30 R PASS 29.756 R SOUARE YARD	4.342	4 **** 4 * * * * * * * * * * * * * * *	Samuel 134			metarant - a (154)	And the second of the second	perconnection of	Charles and the way of the	MATERIA .e			
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		4.4		the best of the same of the sa		********	200.00		TOLUENE	70.		Sec. 12. 1933	· č		·
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39/16/83	75	晋 5.75	THEBBER 1	492 25T PASS	356.4 W	25.9775.1	35.31	375T	BO	886	2 30 112	22.484	17757-47	322.57	
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09/16/83 D9/16/83	75	45.75	ZWD P	CO2 15T PASS		16.5 6.1 24.5 (% 5. 7	95.21	.173	9.702 4.406	.101	.129	1.590	30 . RR	21.09	
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39/16/83	75		SRD PA		64.8	15.0 4.0 14.6 3.9		-157	9.720		129	:- 2.430 2.387		33.86	
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AVERAGE	LBS C	EMENT PE	F PASS 13	4557	2	AV6 3.9	. 1967/17	~~ ~!! ~	AVG	.083	.156	2.119	60.96		
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BUN DW	01/23	/84	AT 14	:11:55			~ •	DA	VID P	TOLUE	NE AN	ALYSIS				, (4;			PAGE 2	?
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9/15/8			61.7		PASS	er, Aaron a. Franklik trii		. B 4	25.5	3.6	. 171	.52 .51	48 🤫	3514.994	₩ 087					
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9/15/6			61.7			33 7			دَدْهُ ،			.52 .1		35.627						
5/:5/8			61.79		DASS	4,17		. 5	17.C			52 .0		11.305		.113		50.00		
9/15/6			61.7			7 2ST 1	PASS					.52 -1		#20.366						
6/15/5	3 100		61.7	: 24:	; PASS		64	• 8	11:.5	3.6		EZ .7		13.652		.106	2.970	83.33	64.32	
9/15/6			61.75			A STATE OF THE PERSON.	्राक्ष्यम् <mark>य</mark>	-8 ··	743-B			.52 .0		······ 8-424			1987: Z -005			
5/15/E			61.79		PASS	•		. 6	12.5			.52 .C		8.100		-081				
9/15/6			61.79			14 2 x			43.5					· 8. 748			· · · · · · · · · · · · · · · · · · ·			
9/15/8	2 100		61.75	TALC	PA 55				16.0	15.2	171	.52 .D	93	10.368		.103	_			
		····•	TALS				** :	*, **	410-2	134.5	49 74	-bo - 1	4 2	427.612		*	-	- The Control of the		
OTAL P	4 C CF C	. >0		. ,		110 200	. ** ***		~ -1		and start	- /~		- min	0	076	m: 1.572		26-92	•
			TIME	07:30	0.1			, a ,	· 51	X 8.6		****		WAX	-241	414				
				R PASS										~~~~~~			·			
				F SOUAT			2			_ , ~ , ~ ,	~~	. 1500			. 3000		Harrison.	,	4	

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

DAVID P TOLUENE ANALYSIS

CEN TOLUENE TOL TOL FEET F/FEET

WHATE LEN WINTH DESCRIPTION STOL SLES WINS SO YDS SYD USED SYD LYD WIN WIN

GRAND TOTAL AVERAGE LOS CEMENT PER PASS 27.233

FERAND TOTAL AVERAGE LOS CEMENT PER SOURE VARD .152

1/4 7 PAH 195.660

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1. 2

TOTAL CONTROL OF THE CONTROL OF THE

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 Product Code: 4-Black Special Magnehelic Setting: .07-.08 Roll Length: 50 yards (resurface blanket) Operator: Mack Duct Temp: 135°F

Roll Width: 79"
Plys: 3-ply

Yards (resurface blanket) Operator: Mack Duct Temp: 135°F Flow ACFM: 5698

Y Flow SCFM: 5076

Pass	Rubber	% VOC	Cement Lbs/Pass	Toluene Lbs/Pass	Min/ Pass	Lbs Tol/ Min	Ft/Min Tach	Ft/Min Calc	ppm Meter	% LEL Meter	% LEL Calc
1	В 1280	58.5%		Mixture of coluene, MEK nochloroben-	&	N.A.	40-42	45.5	N.A.	N.A.	N.A.
			ze	ene							
2	В 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 N	ot 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:
Plys:

62" One Product Code: One Ply Red

Operator:

Mack

Magnehelic Setting:

Duct Temp:

.07-.08 135°F

Flow ACFM (a 135°F):

: 5698

Flow SCFM (a 70°F):

5076

Avg

	·					Lbs					
			Cement	Toluene	Min/	Tol/	Ft/Min	Ft/Min	ppm	% LEL	% LEL
Pass	Rubber	% VOC	Lbs/Pass	Lbs/Pass	Pass	Min_	Tach	Calc	Meter	Meter	<u>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 - 5 7	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	3	27.0	18.8	13.6	1.38	Not on		1313	10.3	7.81
							Tach				

Readout

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:

Avg % LEL:

14.9%

2.5

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:

Duct Temp:

.07-.08 135°F

Flow ACFM (a 135°F): 5698

Flow SCFM (a 70°F): 5076

						Avg					
						Lbs					
			Cement	Toluene	Min/	Tol/	Ft/Min	Ft/Min	ppm	% LEL	% LEL
Pass	Rubber	% VOC	Lbs/Pass	Lbs/Pass	Pass	Min_	Tach	Calc	Meter	Meter	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 Rea	ading)	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

Tase No. 84-0.044





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 BAOM

P 408 530 377

RECEIPT FOR CERTIFIED MAIL

NO INSURANCE COVERAGE PROVIDED— NOT FOR INTERNATIONAL MAIL

(See Reverse)

Ware
\$
;
\$
:

200	SENDER: Conspicts Hems 1, 2, and 3. Add your address in the "RETURN TO" space on severse.
m 3811, Jan, 1979	1. The following service is requested (check one.) Show to whom and date delivered
	(CONSULT POSTMASTER FOR FEES)
_	2 ARTICLE ADDRESSED TO:
RETURN	Mr. Michael A. Ware 2010 Indiana Street Racine, Wisconsin 53405
AECEIP)	3. ARTICLE DESCRIPTION: REGISTERED NO. CERTIFIED NO. INSURED NO.
, F	P408530377
G	(Always obtain signature of addressee or agent)
RECEIPT, REGISTERED, Machine	I have received the article described above. SIGNATURE DADDRESS DAUthorized agent DATE OF DESIVERY DATE OF DESIVERY S. ADDRESS (Complete only if requestion) 2 33
	6. UNABLE TO DELIVER BECAUSE: COERL'S
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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:

- 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 <u>Overall Compliance Schedule</u> 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 State of Florida
DEPARTMENT OF ENVIRONMENTAL REGULATION

INTEROFFICE MEMORANDUM

And/Or To	outing To District Offices o Other Than The Address	
To:	Loctn.:	·
To:	Loctn.:	
То:	Loctn.:	
From:	Date:	
Reply Optional []	Reply Required []	Info. Only []
Date Due:	Date Due:	

TO: Nancy Wright

THRU: Bill Thomas

FROM: Willard Hanks wmk

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 Aunt copy 11/22/83

ESTIMATED

OVERALL COMPLIANCE SCHEDULE

DAVID M COMPANY VOC CONTROL PROJECT

PHASE

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



Graphic Supplies Division 7520 University Des Moines, Iowa 50311 Tet. (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 0 2 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

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

Michael a. Wave

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:

VEAD	OPERATING HOURS/DAY	ESTIMATED % NORMAL BUSINESS VOLUME	ANNUAL EMISSIONS
YEAR	OPERATING HOURS/ DAT	VOLUME	ANNUAL EFESSIONS
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?

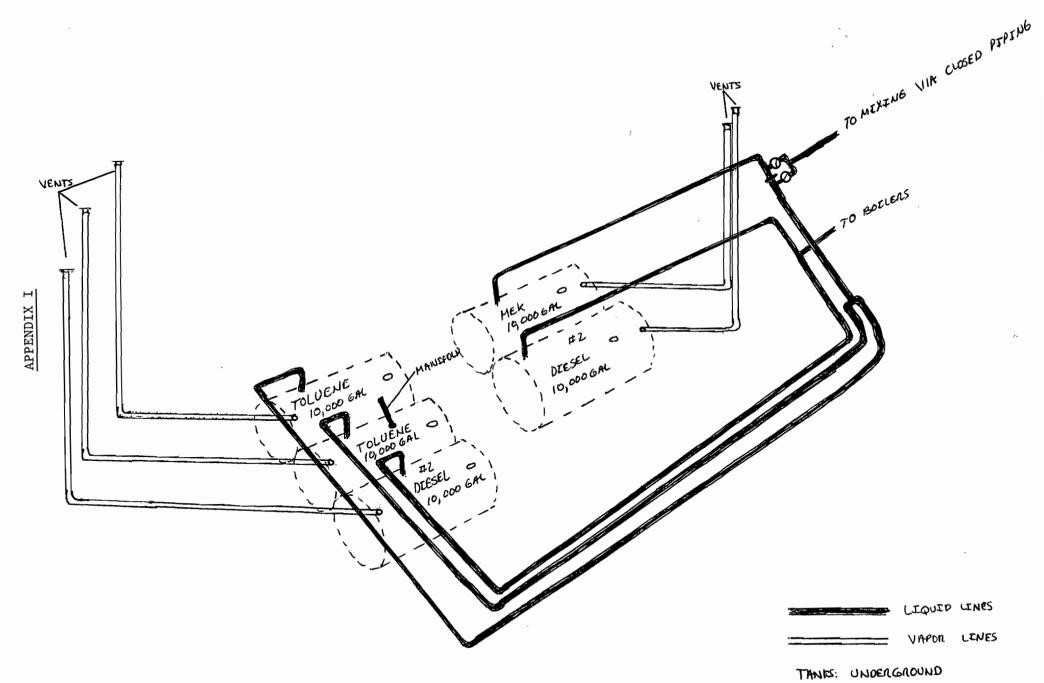
A coaxial vapor recovery system of the type 1 or 2 nature will be installed to minimize VOC emissions during off loading. In addition, an OPW 523, or 523 - S pressure vacuum vent will be installed on each tank vent line to minimize VOC emissions during storage. Appendix II contains example information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

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



PLUMBING: UNDERGROUND TO PUMPS

COAXIAL

SECTION B

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

System C

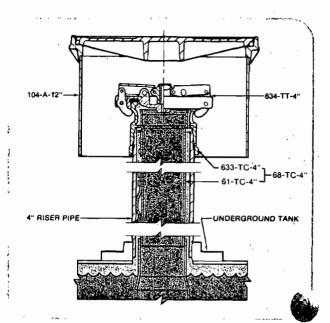
TYPE 1 = 68-TC - 3" & 4" Series Fittings
(Non Poppeted)

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

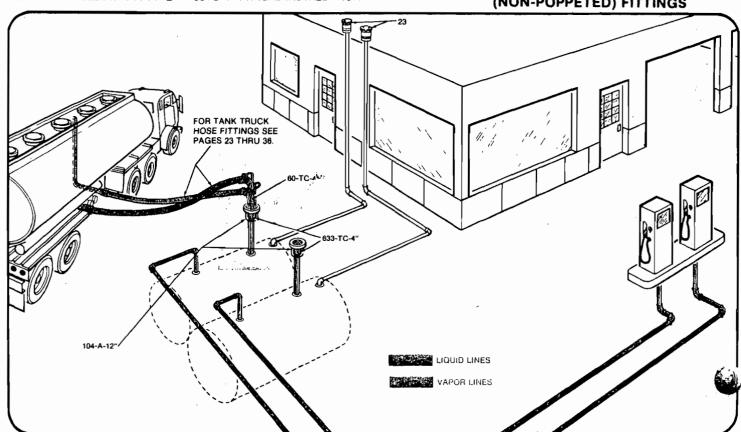
An easy, inexpensive means to convert existing conventional 3" & 4" size underground storage tank fill pipes to coaxial installations. Normally little to no digging or tearing up of concrete is necessary. Simply remove existing tight fill adaptor and fill pipe (if installation is so equipped), inserting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

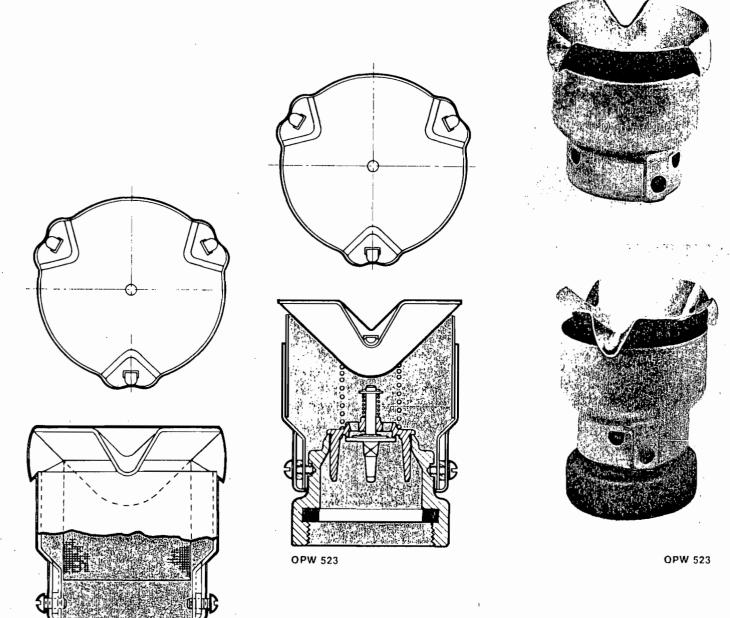
NOTE: Type 2 poppeted fittings are certified by California Air Resources Board (CARB)

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



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





OPW 23, 523, 523-S VENTS

OPW 23

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 $1\frac{1}{2}$ ", 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 $\frac{1}{2}$ 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.

COMPOUND	# SAMPLES ANALYZED	AVG LBS. TOLUENE/100 LBS.	AVG LBS RUBBER/
7002 5503	48 22	58.9 63.7	41.1 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.

COMPOUND	HI	LO	AVERAGE	NUMBER TESTED
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 > .3 microns in diameter.

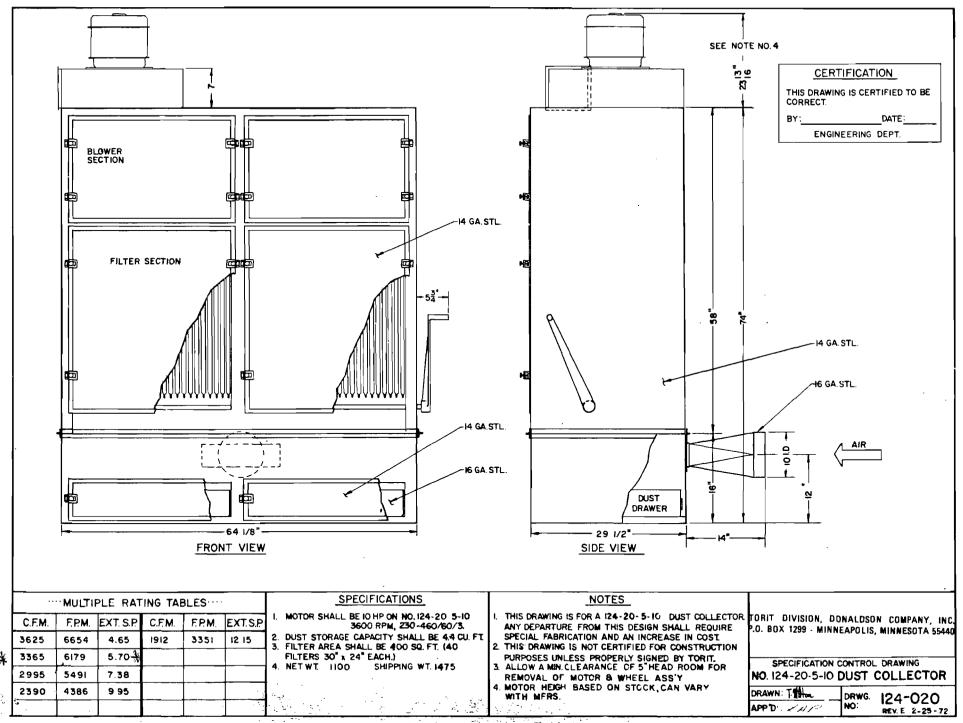
11. a) What will be the particulate and VOC emissions from emission point No. 2 mills?

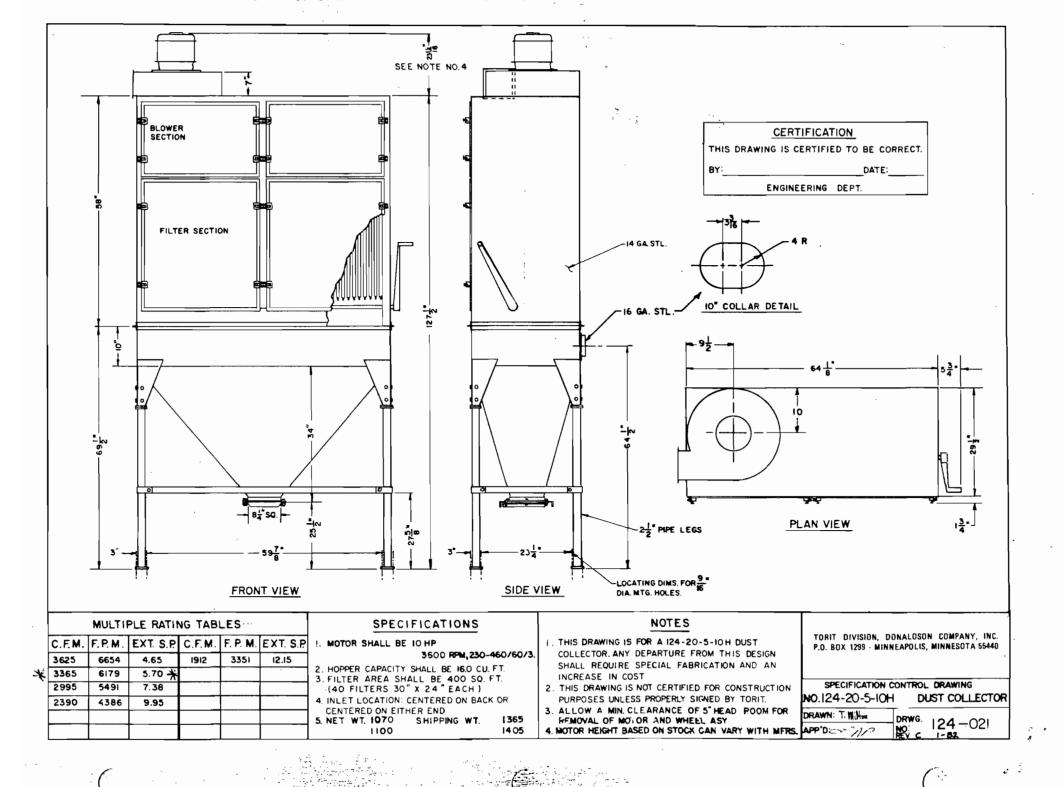
Particulate and VOC emissions combined from the milling operation would approximate 1.7 - 8.0 lbs/day. These numbers were generated by weighing all the individual components to be added to a batch before milling and then weighing the total batch post milling and noting the lost material. The plasticizers 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.

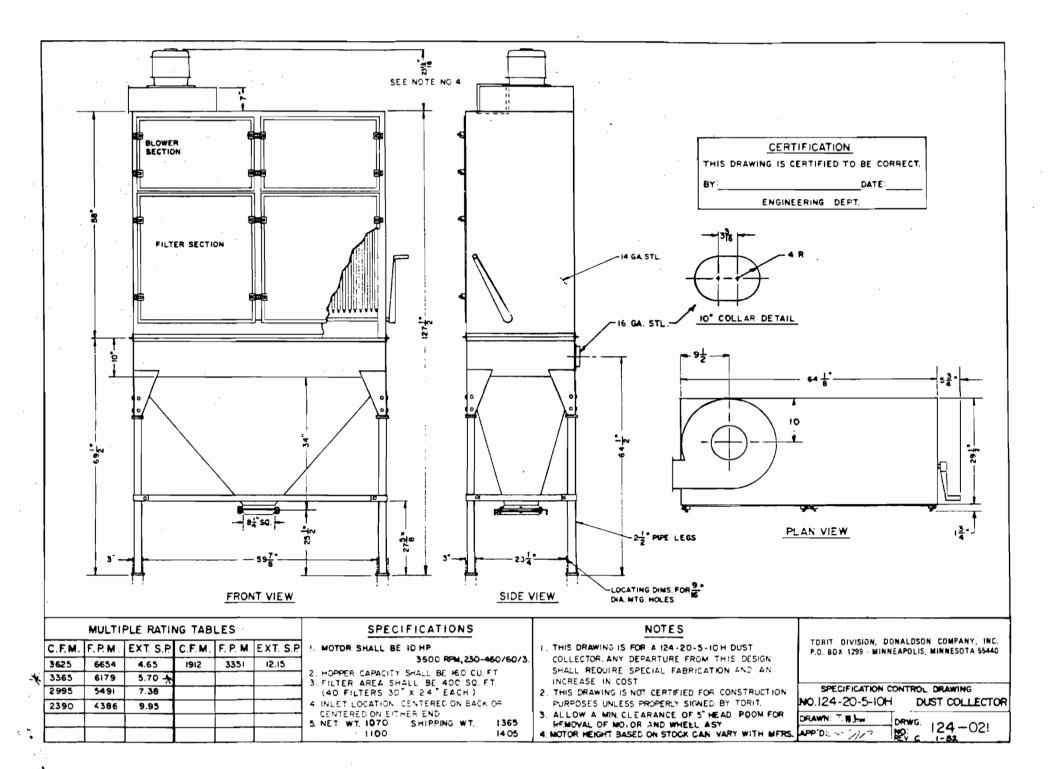
b) What control equipment will be used to limit particulate emissions from the mills?

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









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g bromobenzene and

lant disease control;

rystals: b.p. sublimes;

n hot water, alcohol,

powder; m.p. 73-

ery soluble in alcohol.

etic acid: synthesis of

. See diphenadione.

h crystals. Soluble in

hol, and ether; insolu-

s. 52.85°C; b.p. 302°C;

on temp. 1173°F; com-

mula weights of aniline

n autoclave. The prod-

chloric acid to remove

lined paper bags; fiber

kin. Tolerance, 10 mg

nd accelerators: solid

dyes; pharmaceuticals;

preservation of apples:

damsite: phenarsazine

stals. Sublimes readily.

410°C (dec): insoluble

xylene, carbon tetra-

enylamine with arsenic

ialation and ingestion:

Poison label. Not ac-

48-250°C; insoluble in-

ngth shifter in solution

soluble in water; slightly

atic hydrocarbons. Sen-

id fuming sulfuric acid.

Rail) Tear gas label.

sluble in toluene.

LC6H4NHC6H5.

See benzidine.

id nitrites.

alytical chemistry.

1)C6H2.

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residue is distilled.

e and fused.

niline) (C, H5)2NH.

ant for polyesters.

(C,H,):CHCN.

OH.

tion. m in air. nsfer agent: fungistat

ethyl acciaic Use: Polymerization inhibitor. diphenylbromoarsine (C₆H₅)₂AsBr.

Properties: White crystals. M.p. 54-56°C. Derivation: (a) Hydrobromic acid and diphenylarsenious 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 (C₆H₅NHNH)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 (C₆H₅O)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.

 $\label{eq:diphenylchloroarsine} \textbf{diphenylchloroarsine} \ (C_6H_5)_2AsCl.$ 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.

Hazard: Highly toxic by inhalation; strong irritant to Grade: Technical. tissue.

Use: Military poison gas. Shipping regulations: (Rail) Tear Gas label. (Au) Poison label. Not accepted on passenger planes,

diphenyldecyl phosphite $(C_6H_5O)_2POC_{10}H_{21}$. Properties: Nearly water-white liquid; sp. gr. 103 (25/15.5°C); m.p. 18°C; refractive index 1:5160/0 25/D). Combustible.

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

Properties: Colorless liquid; b.p. 305°C; f.p. 725 sp. gr. 1.19 (20°C): refractive index (n 25/D) 15703 flash point (COC) 288°F. Readily hydrolyzed by moisture, with liberation of hydrochloric Derivation: (a) Reaction of powdered silicon Combustible.

chlorobenzene in the presence of copper powde catalyst; (b) reaction of phenylmagnesium chloris with silicon tetrachloride.

Grade: Technical.

Hazard: Highly toxic: strong irritant to tissue. Use: Intermediate for silicone lubricants.

Properties: Colorless oil. BEST AVAILABLE COPYDerivation: Reaction of didodecyldichlorosilane with

phenyl lithium. Hazard: Probably toxic. Use: High-temperature lubricant.

diphenyldimide. See azobenzene.

diphenyldimethoxysilane (C6H3)/Si(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)

(C, H,), CHCH, 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; com-

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) C. H. CH. CH. C. H. 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) C.H.NHCH.CH.NHC.Hs.

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

Derivation: Treatment of aniline with cyanogen chlo-

C. H. HC: CHCH: CHCCH: CHC, H. Use: Wavelength shifter in solution scintillation

diphenyl isophthalate (DPIP) Ċ。H。ÖOCĊ。H₂COOC。H‹.

Properties: White solid; m.p. 138-139°C. Combus-

Use: Manufacture of polybenzimidazoles, high temperature-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; 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 isocvanate; 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(C, H₃)-

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 reguations: (Air) Corrosive label. (Rail) Corrosive material, n.o.s., White label.

diphenylmethylchlorosilane $(C_6H_5)_2(CH_3)SiCl$. 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 C10 H6 (NHC6 H5)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) OOC6H5:CC6H5. 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.

triethylborane (triethylborine; boron triethyl).

(C₂H₃)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: Igniterials.
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

Red label. (Air) Flammable Liquid label. (Air) 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 cg. solubility in oil 0.8

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 $N(CH_2CH_2)_3N$. 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 (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

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)₅OCC₇H₁₅.

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)

C₃H₁₁OCOCH₂(CH₂OCH₂)₂CH₂OCOC₅H₁₁.

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.

Use: Plasticizer.

triethylene glycol di(2-ethylhexoate)

C₇H₁₅OCOCH₂(CH₂OCH₂)₂CH₂OCOC₇H₁₅.
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 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 C₂H₃CO(OCH₂CH₂)₃OOCC₂H₃.

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 butoxytriglycol.

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

triethylenephosphoramide (tepa; tris-(1-aziridinyl)phosphine oxide; APO) (NCH2CH2), PO.

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[I-(2-methyl)aziridinyl]phosphine oxide.

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.

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triethylenetetramine NH₂(C₂H₄NH):C₂H₄NH₂.
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-ethylbexyl)phosphate [C₄H₉CH(C₂H₃)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 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/ triethy triethy etho

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جرة late! (Air) Legal

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solubility in oil 0.8

(COC): combustible.

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'20°C); b.p. 287.4°C;

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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

C, H, CO(OCH2CH2), OOCC, H, Properties: Crystals; b.p. 210-223°C; m.p. 46°C; flash point 457°F (TOC); sp. gr. 1.168. Combustible. Low

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

 $C_{\circ}\dot{H}_{19}COO(C_{2}H_{4}O)_{3}OCC_{9}H_{19}.$ 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)

CsH11OCOCH2(CH2OCH2)2CH2OCOC5H11 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, H₁₅OCOCH₂(CH₂OCH₂)₂CH₂OCOC, H₁₅. 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; re-fractive index (20°C) 1.5180; vapor pressure (225°C) 2.5; flash point 226°C; insoluble in water. Combustible. Low toxicity.

Use: Plasticizer.

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The Town for icity

Properties: Water-white liquid; mild ether odor; sp. 20/2000 0.0862; refractive index 1.4233 (n. 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.

Containers: Glass bottles; cans; 55-gal drums. Uses: Solvent for gases; coupling immiscible liquids.

triethylene glycol dioctoate. See triethylene glycol dicaprylate.

triethylene glycol dipelargonate $C_8 H_{17} COO(C_2 H_4 O)_3 OCC_8 H_{17}$. 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. Combustriethylene glycol dipropionate C2H3CO(OCH2CH3)3OOCC2H3

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-

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.

triethylenephosphoramide (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 tis-

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 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 eve damage.

Uses: Detergents and softening agents; synthesis of dyestuffs, pharmaceuticals and rubber accelerators. Shipping regulations: (Air) Corrosive label

tri(2-ethylhexyl)phosphate [C₄H₉CH(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 (C₆H₁₇O)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 C₆H₃(COOC₈H₁₇)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.

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

triethyl orthoformate (triethoxymethane) CH(OC₂H₅)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) (C2H5)3PO4

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). Com-

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 (C2H5)3PO3.

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) (C₂H₅O)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 (C2H5OCOCH2)2CHCOOC2H5. 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: solparts and precision

ethyl)trimethylammo-OHI C.H.O. ion): drums.

ichloroethylene (a.v.).

cobaltic oxide.

tricyclamol chloride

oluble in alcohol: in-9 (48°C); b.p. 234°C

ners.

COOH. A saturated in natural fats or oils crystalline solid: m.n. ed in medical research as chromatography.

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

phate; TCP)

of isomers. odorless liquid. Starefractive index 1.556); wt/gal 9.7 lb; crvs-Miscible with all the s. also with vegetable point 437°F; autoigni-

osphorus oxychloride. tank cars. ngestion and skin ab-

highly toxic; its tolerof air. chloride, polystyrene,

for plastics; air filter aterproofing: additive s; hydraulic fluid and

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

for plastics and resins.

H₈N · CH₃Cl. 1-Cyclo-

der: faint characteristic odor: soluble in alcohol and 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.

tricvclohexvl borate. See boric acid ester.

n-tridecane CH₃(CH₂)₁₁CH₃. Properties: Colorles 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 (tridecylic 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 C15 hydrocarbons. Grade: Technical

Containers: 55-gal drums; tank cars.

Uses: Esters for synthetic lubricants; detergents; antifoam 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.

tridecylic acid. See n-tridecanoic acid.

ployed.

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-

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 $C_6H_{11}C(C_6H_5)(OH)CH_2CH_2N(C_2H_5)_2 \cdot C_2H_5CI.$ (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.

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Properties: Liquid; refractive index 1.5181. Combustible

Hazard: May be toxic.

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

 $\begin{array}{ll} \textbf{tri(dimethylphenyl)phosphite} & (trixylenyl & phosphate) \\ & [(CH_3)_2C_6H_3O]_3PO. \end{array}$

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." Trademark for trimethadione, U.S.P.

tridodecyl amine. See trilauryl amine.

tridodecvl 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 CIC₃N₃[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

(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(C₂H₄OH)₃]₂. Hazard: Highly toxic by ingestion.

Use: Herbicide.

Shipping regulations: (Rail, Air) Arsenical compounds,

triethanolamine oleate. See trihydroxyethylamine ole-

triethanolamine stearate. See trihydroxyethylamine stearate

triethanolamine titanate. See titanium chelate.

1.1.3-triethoxybexane

CH(OC,H₁)-CH₂CH(OC₂H₃)C₃H₃.

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)(C₃H₃O)CHCH₃CH(OC₃H₃)₃

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

Grade: 99%.

Uses: Intermediate: crosslinking and insolubilizing

triethylaconitate

C,H,OOCCHC(COOC2H5)CH2COOC2H5.

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). $(C_2H_4)_3Al$.

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

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

triethylamine $(C_2H_5)_3N$.

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

Uses: Catalytic solvent in chemical synthesis; accelerator 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.

PLASTICIZERS AND PLASTICIZATION

Table 15.3

APPENDIX V

Properties of Commercially Available Plasticizers Arranged in Order of Ascending Number of Carbon Atoms

No.	. Name	Formula	Molecular Weight	۵° د.		20° C.	l so	dubility,	% at 20°	, c.	in	platility of Plasticiser in Air, eq. cm./hr.	
				Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)	
		Simple Pl	kısticize	4	_						·		
1 2 3 4 5 6 7 8 9	Acetin o- and p-Toluenesulfonamides (mixture) Diacetin N-Ethyl-p-toluenesulfonamide N-Ethyl-o- and p-toluenesulfonamides (mixture) Triacetin Dimethyl phthalate o-Nitrobiphenyl Diethyl phthalate Triethyl citrate Dimethyl sebscate	C ₅ H ₁₀ O ₄ C ₇ H ₉ O ₂ NS C ₇ H ₁₂ O ₅ C ₉ H ₁₃ O ₂ NS C ₉ H ₁₃ O ₂ NS C ₉ H ₁₄ O ₆ C ₁₀ H ₁₀ O ₄ C ₁₂ H ₉ O ₂ N C ₁₂ H ₁₄ O ₄ C ₁₂ H ₂₀ O ₇ C ₁₂ H ₂₂ O ₄	134 171 176 199 199 218 194 199 222 276	1. 190 at 25° C. 1. 353 solid 1. 178 1. 171 at 65° C. 1. 190 at 25° C, 1. 118 1. 136 at 25° C. 0. 986 at 30° C,	1. 445 at 25° C. 1. 438 at 25° C. 1. 822 at 65° C. 1. 540 at 25° C. 1. 613 at 25° C. 1. 613 at 25° C. 1. 613 at 25° C. 1. 441 at 25° C. 1. 441 at 25° C.	68.8 at 25° C. f.p. 105 38.7 at 25° C. 59.0 426 at 25° C. 15.1 at 25° C. 16.3 38 at 25° C. 12.6	0.15	0.7	0.5	0.34	0.55	101 >1000	

		Con	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticised to Equivalence												
No.	Name	Pasticiser p.b.r. at 23° C., 78% Elongation, 1000 p.s.	şth,	gation,	eter A	Stiffness at 23° C., p.s.i.		arature, °C.	% Loss, 4-mit Film		10 d 23° C.	% Extrac- tion, 10 days, 23° C., 4-mil Film		Supplier ^m	
		Plasticiser p.1. 75% Elongati	Tensile Strength, p.s.i. at 23° C.	Tensile Strength, p.s.i. at 23° C. Ultimate Elongation, % at 23° C.	% at 23° C. Shore Durometer at 23° C.		T4, °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	O;I	Spew, 2 days, 60° C., 100% Humidity b		
			Simple	Plastici	izera										
1	Acetin												_	Kee	
2	o- and p-Toluenesulfonamides (mixture)													Mon	
3	Diacetin													Kee	
4	N-Ethyl-p-toluenesulfonamide							1						Mon ·	
5	N-Ethyl-c- and p-toluenesulfonamides (mixture)													Mon	
6	Triscetin										,			Kes, TE	
7	Dimethyl phthalate													Mos, TE	
8	o-Nitrobiphenyl													Mon	
9	Diethyl phthalate											.		Mon, TE, USI	
10	Triethyl citrate													Pf	
11	Dimethyl sebacate	1												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,

PLASTICIZERS AND PLASTICIZATION

Table 15.3 (Continued)

Properties of Commercially Available Plasticizers Arranged in Order of Ascending Number of Carbon Atoms

No.	Name			20° C.		ن پ	So	lubility,	Volatility of Free Plusticises in Air, mg./eq. cm./hr			
	Namo	Formula	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscoeity, cp. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plasticiz	era (Cor	utinued)								
12	Di-n-butyl tartrate	C12H22O6	262	1.094	1.444 st 25° C.	67.7 at 25° C.	0.5	0.7	0.3	2.4	1.50	143
18	Tri-n-butyl phosphate	C12H27O4P	266	0.978	1.424 at 20° C.	3.4 at 25° C.	< 0.07	6.0	M	М	>4	502
14	Monomethyl phthulate ethyl glycolate ester	C13H14O6	266	1.227	1.504 st 25° C.	185	0.1	1.2	0.5	0.1	0.65	61
15	N-Cyclohexyl-p-toluenesulfonamide	C13H19O2NS	253	1.125 solid	Bt 25 C.	f.p. 87° C.					0.05	26
16	Monoethyl phthalate ethyl glycolate ester	C14H16O6	280	1.186	1.498 at 25° C.	68.0	<0.08	1.38	1.60	0.18	0.50	93
17	Di(methyl Cellosolve) phthalate	C14H18O6	282	1.169	1.501	at 25° C. 53	0.85	3.2	0.50	0.10	0.16	61
18	Triethyl acetylcitrate	C14H22O8	318	1.13	at 25° C.							> 525
19	Diisobutyl adipate	C14H26O4	258	0.957	at 23° C. 1.428	70						
20	Tributyria	C15H26O6	302	1.036	at 25° C. 1.433	at 25° C. 8. 9						
21	Methyl CELLOSOLVE laurate	C ₁₈ H ₃₀ O ₃	258	0.894 at 25° C.	at 25° C. 1.440 at 25° C.	at 25° C. 7.5 at 25° C.						,

		Con	parisor	of Pr	opertic	es of Vi	nylite F Equiv	Resin V' Mence a	ZNW-6	Specim	ens Pia	sticised	to,	
No.	Name	.r. at 23° C., on, 1000 p.s.i.	्य <u>्</u>	gation,	eter A	° C., p.s.i.		rature, °C.	% I 4-mil	оss, Film	% Extrao- tion, 10 days, 23° C., 4-mil Film		60° C.	Supplier "
		Plasticizer p.h.r. at 23° C., 75% Elongation, 1000 p.a.i.	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durome at 23° C.	Stiffness at 23°	T4, °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	වි	Spew, 2 days, 60° C., 100% Humidity	
		Simple	Plasticia	era (C	ontinu	ed)					_			
12	Di-л-butyl tartrate				•			·			<u>-</u>			Kes
13	Tri-n-butyl phosphate													C8
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				1						1			Mon
17	Di(methyl Chilosolve) phthalate	46 0							6.6		4.6	3.7	None	OALTE
18	Triethyl acetylcitrate													Pf ,
19	Dissobutyl adipate										ļ			Cab
20	Tributyrin												—	TE
21	Methyl Cellosolve 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.
 a 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute.
 Bee Reed. 25
 For key to supplier abbreviations see page 903.

PLASTICIZERS AND PLASTICIZATION

Table 15.3 (Continued)

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

V	Name			ر. د.		۴.	80	dubility,	°c.	in.	atility of Plasticiser a Air, q. cm./hr.	
No.		Formuls	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° P. (100° C.)	875° F. (191° C.)
		Simple Plasticis	era (Coi	itinued)				-				
22	Di-n-butyl phthalate	C16H22O4	278	1.048	1.493 at 20° C.	20.3	InsoL	0.46	34	30.0	0.98	148
23	Di(Cellosolye) phthalate	C16H22O6	310	1.120	1.491 at 25° C.	f.p. 31 °C.	0.2	2.0	2	0.64	0.15	30
24	n-Butyi laurate	C ₁₆ H ₃₂ O ₂	256	0.857 at 25° C.	Bt 23 C.	31 0.						
25	Methyl Czllosolve 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 ₁₆ 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	C18H22O	286	1.008	1.553 at 25° C.	29.0						'
29	n-Butyl cyclohexyl phthainte	C18H24O4	304	1.078	1.508 at 20° C.	117.0						75
30	Mouo-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	Triethylane glycol di(2-ethylbutyrate)	C18H34O6	346	0.995	1.440 at 20° C.	11.5	0.02	1.0	0.2	24.0	0.86	

		Con	Comparison of Properties of Vinylite Resin VYNW-5 Specimens Plasticised to Equivalence												
ło.	Name	on, 1000 p.s.i.	th.	Elongation, C.	eter A	23° C., p.a.i.		Temperature, °C.	% Loss 4-mil Fil		tic	strac- on, ays, , 4-mil lm	2 days, 60° C., Humidity	Supplier **	
		Plasticiter p.h.r. 75% Elongation,	Tensile Strength, p.s.i. at 23° C.	Ultimate Elor % at 23° C.	Shore Durometer at 23° C.	Stiffness at Z	T. °C.	Brittle Temp	10 days, 60° C.	24 hours, 70° C.	Water	750	Вреч. 2 days. 100% Humid		
		Simple	Plastici	eers (C	ontin	ed)									
22	Di-n-butyl phthalate	41.4 2				•			17.5		2.7	6.8	None	Bar, CS, Har, Mon, Nev, Pitt, TE, USI, DA	
23	Di(Crilosolve) phthalate							l						OA	
4	n-Butyl laurate													AH	
5	Methyl Czilosolve myristate													AH	
26	Di(butyl Carstrol) formal	55.3 °							10.1		15.6	14.5	None	Thi	
27	Triphenyl phosphate		· .										_	Dow, Mon, Pitt	
28	Bis(dimethylbensyl) ether			1									-	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°	3000	300					8.3		17	4.8	None	Mon	
31	Tri-n-butyl citrate	55.1	2800	300			-4	-82	8.1		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.²⁶
 For key to supplier abbreviations see page 903.

Table 15.3 (Continued)

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

No.	Name			0° C.		20° C.	So	lubility,	C.	Free Pt	lity of asticizer Air, cm./hr.	
110.		Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, ep. at 20	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plasticiz	ета (Соп	tinued)								
33	Butyl Cellosolve laurate	C18H36O3	300	0.882 at 25° C.	1.439 at 25° C.	6.5 at 25° C.						
34	Di-n-hexyl adipate	C18H34O4	314	(av.) 0.933 at 25° C.	1.439 at 25° C.	7.8 at 25° C.						
35	Di(1,3-dimethylbutyl) adipate	C18H84O4	314	0.926	1. 433 at 25° C.	5.6 cs. at 38° C.						
36	Di-n-butyl sebacate	C18H84O4	314	0.936	1.440 at 25° C.	9.8	Insol.	1.6	M	М		63
37	Di(butyl Cellosolve) adipate	C18H24O6	346	0.997	1.442 at 25° C.	12.5						58
38	n-Butyl myristate	C18H26G2	284	0.861 at 25° C.	at 20 C.							
39	Tri(butyl Cellosolve) phosphate	C ₁₈ H ₃₉ O ₇ P	398	1.022	1.434 at 25° C.	12.2	0.08	6.8	Almost	nuiscible	1	
40	Cresyl diphenyl phosphate	C10H17O4P	340	1.208 at 25° C.	1.563 at 20° C.	39.8						
41	n-Butyl benzyl phthalate	C19H20O4	312	1.116 at 25° C.	1.538 at 20° C.	39.8 at 25° C.					•	29
42	Bis(dimethylbensyl) carbonate	C19H22O8	298	1.083	at 20°C. 1.547 at 20°C.	155						32

M = Miscible.

		Con	npariso	of Pr	operti	es of Vi	nylite l Equiv	Resin V alence a	YNW-5	Specim	ens Plac	sticized	to	
No.	Name	i.r. at 23° C., on, 1000 p.s.i.	rth,	gstion,	ster A	° C., p.s.i.		rature, °C.	% I 4-mil	oss, Film	10 d	ttrac- on, laye, , 4-mil	2 days, 60° C., Humidity	Supplier **
		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 Durome at 23° C.	Stiffness at 23°	T4, °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	110	Spew, 2 days, 100% Humidi	
		Simple	Plastici	ета (С	ontinu	ed)								
33	Butyl Cellosolve laurate													Kes
34	Di-n-hexyl adipate													Kos
35	Di(1,3-dimethylbutyl) adipate)											Har
36	Di-n-butyl sebacate	37.8 °							7.2		0.4	10.5	None	Har, Pitt, RH
37	Di(butyl Cellosolve) adipate	41.4 °										18.0	None	Kes, OA
38	n-Butyl myristate	1												AH .
39	Tri(butyl Cmilosolve) phosphate	43.7°	ļ					1	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	Моц
	Bis(dimethylbenzyl) carbonate	71	3000	284	63		ı	ı	1	11.8	,	9.2	ı	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.²⁶
 For key to supplier abbreviations see page 903

Table 15.3 (Continued)

PROPERTIES OF COMMERCIALLY AVAILABLE PLASTICISERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

				J. C.		°.	So	lubility, '	% at 20°	C.	l in A	ility of asticiser Air, cm./hr.
No.	Name	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, ep. st 20° C.	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	875° F. (191° C.)
		Simple Plasticiza	ers (Con	tinued)								
43 44 45 46 47 48 49 50 51 52	Methyl pentachlorostearate Methyl ricinoleate Methyl Cellosolve palmitate Methyl hydroxystearate Diphenyl phthalate Dicyclohoxyl phthalate 2-Ethylhoxyl diphenyl phosphate Di-n-bexyl phthalate Di(2-ethylhutyl) phthalate Di(1,3-dimethylbutyl) phthalate	C19H23O2Cl6 C19H26O2 C19H28O2 C19H28O3 C20H14O4 C20H26O4 C20H27O4P C20H20O4 C20H20O4 C20H20O4	471 312 314 314 818 330 362 834 334	1. 204 0. 925 at 25° C. 0. 891 at 25° C. 1. 28 solid 1. 23 1. 092 1. 007 1. 016 0. 995	1.497 at 20° C. 1.402 at 25° C. 1.572 at 74° C. 1.511 at 20° C. 1.487 at 20° C. 1.488 at 25° C. 1.480	6,p. 48° C. 6,p. 69° C. 6,p. 63.5 21.0 29.5 56 28.8 cs. at 38° C.	Insol.	0.24	M	М	0.006	10 44 43 65 > 150

M = Miscible.

														· · ·
	•	Con	parison	of Pro	operti	es of Vi	nylite R Equiv	lesin V Mence a	/N W-5	Specim	ens Plac	sticized	to	
No.	. Name	Pasticiser p.h.r. at 23° C., 75% Elongstion, 1000 p.a.i.	gth,	ngation,	eter A	Z°C., pai		Temperature, °C.	% I 4-mil	oss, Film	tic	xtrac- on, lays, . 4-mil	2 days, 60° C., Humidity b	Supplier ^m
,		Pasticiser p.1 75% Elongst	Tensile Strength, p.s.i. at 23° C.	Oltimate Elongation, % at 23° C.	Shore Durom at 23° C.	P Stiffness at 2	T. °C.	Brittle Temp	10 days, 60° C.	24 hours, 70° C.	Water	ī o	Spew, 2 days, 100% Humid	
		Simple	Plasticis	ers (C	ontinu	ed)								
43	Methyl pentachlorostearate	69 d	2900	338	64	600	- 6	-19	0.2		0.2	17.1	None	Hook
44	Methyl ricinoleate					,			٠					Bak
45	Methyl Crilosolva palmitate								***				 —	AH
46	Methyl hydroxystearate													Bak
47	Diphenyl phthalate													Mon
48	Dicyclohexyl phthalate	64 *	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	45.30							4.4		0.2	8.4	None	TE .
52	Di(1,3-dimethylbutyl) phthalate													Har \widehat{v}_{i}

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. Dash (——) indicates no data available.

100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i, per minute. See Reed.**

Mixture of 1 part plasticiser to 2 parts Flaxol plasticiser DOP.

Mixture of 0.9 part plasticiser to 1 part Flaxol plasticiser DOP.

For key to supplier abbreviations see page 903.

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

				0° C.		ບູ	80	lubility, '	% at 20°	C.	Free Plain	lity of asticiser Air, em./hr.
No.	Name .	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, ep. at 20°	In Water	Wster in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	875° F. (191° C.)
		Simple Plasticiz	ers (Con	tinued)								-,-
53	Di(butyl Cellosolve) phthalate	C20H20O6	866	1.063	1.483 at 25° C.	42.0	Insol.	0.63	2.80	6.50	0.06	24
54	Di(Carbitol) phthalate	C20H30O8	398	1.150	1.492 at 25° C.	82.0	Insol.	2.80	0.09	0.20	0,08	18
55	Tri-n-butyl acetylcitrate	C20H34O8	402	1.046 at 25° C.	1.441 at 25° C.							61
5 6	n-Butyl palmitate	C20H40O2	313	0.865 at 25° C.	2020 0.							
57	Tricresyl phosphate	C21H21O4P	368	1.165	1.557 at 20° C.	120.0	.Insol.	0.87	5.5	2.0	0.01	7,
58	n-Butyl benzyl sebacate	C21H32O4	348	1.004	1.479 at 25° C.	8,6 cs.						
59	Methyl acetylricinoleate	C21H38O4	355	0.986	1.456 at 20° C.	at 38° C. 23.1						·
60	Methyl Cellosolve oleate	C21H40O3	341	0.902	1.453 at 25° C.	10.2					0.12	29
61	Di(2-ethylbutyl) aselate	C21H40O4	857	0.930	1.442 at 25° C.	11.4						43
62	Propylene glycol monoricinoleate	C21H40O4	357	0.950 at 25° C.								
63	Methyl Czllosolve stearste	C ₃₁ H ₄₃ O ₃	843	0.877 at 25° C.	1.443 at 25° C.	8.9 at 25° C.						30

		Соп	parison	of Pro	perti	se of Vi	nylite F Equiv	lesin V	YNW-6	Specim	ens Plas	ticised	to	• • • •
No.	Name	or, 1000 p.s.i.	tp.	gstion,	ster A	23°C., pai		Temperature, °C.	% I 4-mil	oss, Film	% E: tic 10 d 23° C. Fi	ttrac- on, ays, , 4-mil	60° C.,	Supplier **
		Pasticiser p.h.r. at 23° C., 75% Elongation, 1000 p.s.i.	Tensile Strength, pai at 23° C.	Oltimate Elongation, % at 23° C.	Shore Durometer A st 23° C.	Stiffness at 23	T. C.	Brittle Tempe	10 days. 60° C.	24 hours, 70° C.	Water	D.IO	Spew, 2 days, 60° C., 100% Humidity	
		Simple	Plastici	eere (C	ontinu	ed)								· · · · · · · · · · · · · · · · · · ·
53	Di(butyl CELLOSOLVE) phthalate	47.80							2,1		4.8	11.4	None	OA
54	Di(CARBITOL) phthalate	54.40							2.0		10.1	9.9	None	OA
55	Tri-n-butyl acetylcitrate				Ιì						·			Pf
56	n-Butyl palmitate										1	•	 —	AH
57	Tricresyl phosphate	75	2880	274	58	580	+ 5	- 8	0.1		0.7	6.1	None	Cel, Mon, OA, Pitt
58	n-Butyl benzyl sebacate			Ì '	1									Har ·
59	Methyl acetylricinoleate	47.80			H				8.8		9.6	11.0		Bak
60	Methyl Cellosolve oleate	50.5 a,f							8.6		8.0	10.8		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 Cullosolve stearate							'				1		AH, Kes
	•		1	I						l				

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
Mixture of 0.63 part plasticiser to 1 part tricresyl phosphate.
For key to supplier abbreviations see page 903.

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

				ပ န		ບູ	Soi	lubility, '	% at 20°	C.	Volati Free Plant in Mg./sq.	
No.	. Name	Formuls	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscosity, ep. at 20° C.	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	376° F. (191° C.)
· 		Simple Plastici	era (Con	tinued)								
64	n-Butyl ricinoleste	C22H42O3	355		ĺ						٠.	
65	Di(1,3-dimethylbutyl) sebacate	C22H42O4	871	0.911	1.439 at 25° C.	9.2 cs.	-					
66	Di(2-ethylhexyl) adipate	C22H42O4	871	0.927	1.447 at 20° C.	at 38° C. 18.7	Insol.	0.12			!	41
67	Diisoctyl adipate	C22H42O4	371	0.928	1.448 at 25° C.	17.7						88
68	Dicapryl adipate	C22H42O4	371	0.915	1.440 at 25° C.	8.3 cs. at 88° C.						
69	Diethylene glycol dipelargonate	C22H42O5	387	0.963	1.447 at 20° C.	19				. '		
70	Di(butyl Crilosolve) sebacate	C33H42O6	403	0.970	86 20 0.						0.04	
71	Triethylene glycol di(2-ethyl hexanoate)	C22H42O6	403	0.968	1.444 at 20° C.	16.1	Insol.	0.4	M	, M	0.18	50
72	Triethylene glycol dicaprylate	C22H42O6	406	0.973	a. 20 C.							
73	Di(butyl Carritol) adipate	C22H42O8	(av.) 435	1.02 at 16° C.		15 at 22° C.						,
74	n-Butyl stearate	C22H44O2	341	0.858	1.444 at 20° C.	7.9 at 25° C.						59

M = Miscible

			·.·						, .			<i>,</i>		
		Con	npariso	of Pr	operti	ies of V	inylite l Equiv	Resin V	YNW-5	Specim	ens Pla	sticized	to	
No.	Name	Pasticiser p.h.r. at 23° C., 75% Elongation, 1000 p.a.i.	th.	ngation,	eter A	23° C., p.e.i.		erature, °C.	% 1 4-mil	oss, Film	10 c	xtrac- on, lays, , 4-mil lm	2 days, 60° C., Humidity	Supplier ^m
		Plasticiaer p.1 75% Elongati	Tensile Strength p.s.i at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer at 23° C.	Stiffness at 23	T. °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	පි	Spew, 2 days, 100% Humid	
		Simple	Plastici	zers (C	ontini	ued)								
64	n-Butyl ricinoleate								ļ -					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,
67 68	Diisoõctyl adipate Dicapryl adipate													Kes, OA, Har Kes, MWi, OA, Pitt, RCA, Har, Cab 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	800	69	800	-17	-50	4.7	12.4	2.0	22.8	None	cccc
72	Triethylene glycol dicaprylate												-	RCA, Drew
78	Di(butyl Carrot) adipate		1											Thi
74	n-Butyl stearate												· ·	CS, AH, Har, Kes, OA

^{4 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.

^m For key to supplier abbreviations see page 903.

Table 15.3 (Continued) PROPERTIES OF COMMERCIALLY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

				ე გ		ູ່	So	lubility,	% at 20°	C.	Volati Free Planta in a mg./sq.	
No.	Name ,	Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, ep. st 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plasticiz	етв (Сот	utinued)								•
75	n-Butyl oleate	C22H42O2	339	0.864 at 25° C.	1.449 at 25° C.	7.7 at 25° C.						
76	Tetrabydrofurfuryl oleate	C23H42O3	366	0.925	1.466 at 20° C.	17.8						
77	Methyl Callosolve acetylricinoleate	C23H42O8	399	0.955	1.456 at 20° C.	42.1	Insol.	0.6	. M	М		12
78	o-Xenyl diphenyl phosphate	C24H19O4P	402	1.233	ca. 1.586 at 60° C.	30-55 at 60° C.	Insol.	0.35	2.22	0.8	0.005	20
79	Dibenayl sebacate	C24H30O4	382	1.055 at 30° C.	1.520 at 30° C.	13.0 cs. at 38° C.						
80	Di(2-ethylhexyl) phthalate	C24H38O4	390	0.986	1.486 at 20° C.	81.0	Insol.	0,20	М	М	0.02	24
81	Diisooctyl phthalate	C24H28O4	300	0.987	1.486	83.0			'			19
82	Dicapryl phthalate	C24H28O4	390	0.966	at 20° C. 1.479 at 20° C.	66.7						22
83	Di(2-ethylhexyl) tetrahydrophthalate	C24H42O4	395	0.969	1.466 at 20° C.	42.0	Insol.	0.26				80
84	Tetra-n-butyl thiodisuccinate	C24H42O8S	491	1.054	1.462 at 20° C.	63.6	Insol	0.22				6
Rō	Di(2-ethylhexyl) hexahydrophthalate	C24H44O4	897	0.956	1.461 at 20° C.	85.9			٠.			37
86	n-Butyl acetylricinoleate	C24H44O4	397	0.929	1.456 at 25° C.	35.3						30

		Con	nparisos	of Pr	oper ti	es of Vi	inylite I Equiv	Resin V	YNW-5	Specim	ens Plas	sticised	to	
lo.	Name	Pasticiser p.h.r. at 23° C., 75% Elongation, 1000 p.a.i.	gtb.	gration,	eter A	Z°C., pai		erature, °C.	% l 4-mi	Loss, Film	10 d 28° C.	itrac- on, lays, 4-mil	2 days, 60° C., Humidity b	Supplier ^{va}
		Plasticiser p.l 75% Elongsti	Tensile Strength, p.s.i. st 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer at 23° C.	Stiffness at Z	T. °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	8	Spew, 2 days, 100% Humid	
		Simple	Plastici	eers (C	ontini	ved)								,
5	n-Butyl oleate													AH, Kes
6	Tetrahydrofurfuryl oleate	59 d	2590	297	67	650	– 8	-34	0.8		0.3	17.2	None	AH, Em, Pitt
7	Methyl Cellosolve acetylricinoleate	50.8°							2.4		1.3	14.5	None	Bak, Dee
В	o-Xenyl diphenyl phosphate													Dow
9	Dibenzyl sebacate	45.3 °		•					0.7		0.8	11.8	None	Har
0	Di(2-ethylhexyl) phthalate	61	2360	253	67	700	- 3	~31	0.7	5.1	0	15.5	None	Bar, CCCC, Har, Mon, MWi, OA, Pitt, TE, DA, Gdr
1 2	Discord attacks	56 64	2700	300 270			- 1	-27 -35	0.6		0	33.8	None None	Bar, Cab, Har, MWI OA, Pitt, RCA RH, Har
3	Dicapryl phthalate Di(2-ethylhexyl) tetrahydrophthalate	66	2590	320	68	560	- 5 -13	-37	4.1	11.5	0.8	21.8	None	CCCC
,	Tetra-n-butyl thiodisuccinate	64	3200	318	62	760	- 13 2	-22	0	2.5	0.8	15.2	None	cccc
5	Di(2-ethylhexyl) hexahydrophthalate	64	2580	300	68	750	-10	-34	"	9.7	0.8	25.4	None	cccc
,	Ditation of the second	01	+000	500	, wo		1 40	-04	I	J	"		110116	1 0000

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 plasticiser to 2 parts Flaxot plasticizer DOP.
 m For key to supplier abbreviations see page 903.

Table 15.3 (Continued) Properties of Commercially Available Plasticizers Arranged in Order of Ascending Number of Carbon Atoms

No.	Name			20° C.		20° C.	So	lubility,	% at 20°	C.	in A	ility of asticiser Air, cm./hr.
No.	Name	Formula	Molecular Weight	Specific Gravity,	Refractive Index	Viscosity, sp. st 20	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plasticis	ora (Con	rtinued)								
88 89 90 91 92 93 94 95 96 97	Butyl Cellosolve oleate Dinonyl adipate n-Octyl n-decyl adipate (mixture) Polyethylene glycol di(2-ethylhexanoate) Butyl Cellosolve stearate Tri(2-ethylhexyl) phosphate Di(2-ethylhexyl) azelate n-Octyl n-decyl phthalate (mixture) Isoöctyl n-octyl n-decyl phthalate (mixture) Di(2-ethylhexyl) sebacate Disooctyl sebacate Dicapryl sebacate	C24H46O3 C24H46O4 C24H46O7 C24H48O3 C24H48O3 C24H48O4 C25H48O4 C26H42O4 C25H42O4 C25H42O4 C26H42O4 C26H42O4 C26H50O4 C26H50O4	383 398 398 (av.) 447 385 435 413 418 (av.) 427 427	0.885 at 25° C. 0.914 at 25° C. 0.920 0.989 0.882 0.926 0.918 0.975 0.978 0.912 0.917	1.454 at 25° C. 1.450 at 25° C. 1.450 at 26° C. 1.447 at 20° C. 1.446 at 25° C. 1.443 at 20° C. 1.485 at 20° C. 1.485 at 20° C. 1.485 at 20° C. 1.447 at 25° C. 1.444 at 25° C.	5.5 at 25° C. 18.5 C. 18.6 A. 25.1 13.3 14.1 20 72 42.9 19.9 20.4 at 25° C. 13.0 ca.	Insol.	1.4	М	Ж		44 26 39 20 8

M - Miscible.

		Co	mparisc	on of P	roper	ties of V	inylite Equi	Resin V valence	YNW-	Specin	nens Pl	neticized	to	
٧o.	Name	.hr. st 23° C., tion, 1000 p.a.i.	मुं	ngstion,	neter A	Z°C., pai		Temperature, °C.	% 4-mi	Loss, I Film	[ti	Extrac- ion, days, 4-mil ilm	2 days. 60° C., Humidity b	Supplier "*
		Plasticiser p.hr. 75% Elongation.	Tensile Strength, p.s.i. st 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 2	T. °C.	Brittle Temp	10 days, 60° C.	24 hours, 70° C.	Water	15	Spew, 2 days, 100% Humid	
		Simple	Plastic	zera (C	Contin	ued)	•	•				!		· · · · · · · · · · · · · · · · · · ·
7	Butyl Cellosolve oleste		Ι									<u> </u>	· .	
3	Dinonyl adipate													Kes
١	n-Octyl n-decyl adipate (mixture)	53	2550	295	78	1050	_ 7	-54		5.5	0.3	23.5	None	Pitt Herc, OA
	Polyethylene glycol di(2-ethylhexanoate)	62	2690	308	70	750	-12	-44	2.8	8.8	3.0	19.4	None	CCCC
l	Butyl Cellosolve stearate		.					**	•	0.0		18.4	14006	
۱ ا	Tri(2-ethylhexyl) phosphate	59	2640	300	66	980	16	-57	8.2	6.8	0.6	24.8	None	AH, Kes, OA
1	Di(2-ethylhexyl) aselate	57	2590	334	66	750	-17	-56	0.5	0.0	0.4		None	
	n-Octyl n-decyl phthalate (mixture)	62	2640	318	84	640	- 8	-88	0.2		0.3	21.8	None	Ein
	Isooctyl n-octyl n-decyl phthalate (mixture)	67	2720	334	66	470	-12	-39	J. 2	3.9	0.8		None	Hero Hero
	Di(2-ethythexyl) sebacate	60	2420	293	67	1700	-19	-56	0.08	4.8	0.3		None	
	Diisočetyl sebacate							"	0.00	1.0	0.3	30.2	14000	Dee, Har, MWi, RH
П	Dicapryl sebacate						' I						-	Har, Pitt, MWi

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

PLASTICIZERS	
AND	
PLASTICIZATION	
31	

No.	Name			Ü			So	olubility,	% at 20°	°C.	Volati Free Pla in A mg./sq.	Air, cua./hr.
No.	Name	Formula	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscosity, cp. at 20° C.	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plasticiz	етв (Сог	ntinued)								
99 100	2-Ethylhexanoic acid diester of N,N-bis(2-hydroxyethyl)-2-ethylhexan- amide Tri(p-tert-butylphenyl) phosphate	C ₂₈ H ₅₃ O ₅ N C ₃₀ H ₈₉ O ₄ P	484 495	0.956	1.458 at 20° C.	139. 2 f.p. 95–99. 5 °C.	Insol.	0.5				12
101 102	Glyceryl triricinoleate Glyceryl tri(acetylricinoleate)	C ₅₇ H ₁₀₄ O ₉ C ₆₃ H ₁₁₀ O ₁₂	933 1060	0.959 at 25° C. 0.964	1.477 at 25° C. 1.469 at 25° C.	°C.		,				0.9
_		Resinous F	lasticis	 T8								
103	Aroctor 1242, chlorinated biphenyl			1.378- 1.388	1.627- 1.629		Insol.	<0.1	М	М	2.50	146
104	FLEXOL plasticizer R-2H, polyester			at 25° C. 1.055	at 20° C. 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					, '	. 8
106	G.E. 2559, polyester			1.060	1.492 at 20° C.	4278						1
107 108	PARACRIL C, nitrile rubber PARAPLEX G-25, polyester		ca.	0.984 1.06	1.470	300,000						0.2

M = Miscible.

		Co	to											
No.	Name		gth.	ngation,	eter A	23° C., pai		rsture, °C.	% Loss, 4-mil Film		10	extrac- ion, days, 4-mil ilm	60° C.	Supplier ^m
		Plasticiter p.h.r. at 23° 75% Elongation, 1000 p	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer at 23° C.	Stiffness at 2	7. ℃.	Brittle Temperature,	10 daya, 60° C.	24 hours, 70° C.	Water		Brew, 2 days, 60° (100% Humidity b	
	·	Simple	Plastic	izera ((Contin	ued)			,					
99 00	2-Ethylhexanoic acid diester of N_rN -bis(2-hydroxyethyl)-2-ethylhexanomide Tri(p -tert-butylphenyl) phosphate	64	2730	827	63	650	- 8	-23	.0.4	1.7	1.1	11.0	None	CCCC Dow
01 02	Glyceryl triricinoleate Glyceryl tri(acetylricinoleate)	68. 6°, £		-					1.4		2.5	9.5	None	Bak Bak
		Rest	nous Pi	lasticiz	sra									
03	ABOCLOB 1242, chlorinated biphenyl	56.2 A							4.5		0.6	0.6	None	Mon
)4	FLEXOL plasticizer R-2H, polyester	90	3080	360	57	440	+2	-10	0	0	0	2.4	None	,
5	O.E. 2557, polyester	72	2730	310	63	580	– 2	-18	0.9	8.5	8.1	17.0	Very	CCCC
16	G.E. 2559, polyester	90 °	2110	215	72	740	6	- 6	0.6	2.3	2.7	7.1	slight Very	GE GE
)7)8	PARACRIL C. nitrile rubber PARAPLEX (I-25, polyester	106 82	2500 2600	410 310	60	480	+ 7 - 1	-54 -25	0.5 0.2		0.2	0.6 0.6	slight None None	Nau RH

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.s.i. per minute.

b Dash (——) indicates no data available.

100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed. 26

Mixture of 0.9 part plasticizer to 1 part tricresyl phosphate.

h Only composition tested, does not yield standard elongation.

h Highest plasticizer content tested. Elongation at 1000 p.s.i. = 04%.

For key to supplier abbreviations see page 903.

DESCRIPTION OF TABLES AND FIGURES

PROPERTY OF COMMERCIALLY					_			37		Cinnon	Among	١.
			A	TAT	ODIVED	OF	ASCENDING	NUMBER	OF	CARBON	VIOWO	1
DOODWING OF COMMERCIALLY	AVAILABLE	PLASTICIZERS	ARRANGED	IN	OUDER	O.D	TEDCMIDING	2100000		-		

				ú		20° C.	Bol	ubili ty, 9	C	Volatili Free Pla in A mg./sq.	sticiser ir.	
No.	Name	Formula	Molecular Weight	Bpecific Gravity, 20°	Befractive Index	Viscosity, ep. st 20'	In Water	Wster in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
!	h	esinous Plastici	sers (Co	ntinued)								
09 10	Paraplex G-40, polyester Paraplex G-50, polyester Plastolrin 9720, polyester		ca. 6000 ca. 2200 ca. 850	1.15 1.084 1.043	1.472 at 20° C. 1.470 at 20° C. 1.462 at 20° C.	290,000 8000 431						0. 1 2
		Hydrocarbon-T	ype Plas	ticisers						<u>,</u>	· -	
112	Dow 276-V2, polymolecular product derived from α-methylstyrene HB 40, partially hydrogenated mixture of isomeric terphenyls Kenplex L, hydrocarbon resin made by reaction of formaldehyde + dimethylnaphthalenes		240	1.021 1.007 1.01 at 15.6° C. 1.075	1.582 at 20° C. 1.570 at 25° C.	112.8 f.p. -1° C. 23,500						115
15 16	NEVILLAC 10° (PHO), a hydrindyl phenol derivative NEVINOL		(av.)	to 1. 10 at 15.6° C. 1.03 to 1.08 at 15.6° C. 0.955	at 25° C. 1.596 (av.) at 25° C. 1.561	at 25° C. 65 to 110 at 25° C.						
17 18	Panaplex BN-1, alkylated aromatic hydrocarbon mixture Sovaloid C, alkylated aromatic hydrocarbon mixture			1.038	at 20° C.	105						,

	The state of the s	Соп	to	1											
o.	Name	on, 1000 p.s.i	tth,	gation,	eter A	Z°C., pri		rature, °C.	% I 4-mil	Oss, Film	10 d 23° C.	xtrac- on, lays, , 4-mil	60° C.	- 1	Supplier ^m
		Plasticiser p.b.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 . at 23° C.	Stiffness at 23	T. °C.	Brittle Temperature,	10 days, 60° C.	24 hours, 70° C.	Water	8	Spew, 2 days, 60° C 100% Humidity b		
Ċ		Resinous	Plastic	izers (C	Contin	ued)									
۱ و	PARAPLEX G-40, polyester	67 j	2780	310	69	760	1	-15	0.3		1.0	5.2	Slight	RH	٠,
Ò	PARAPLEX G-50, polyester	75	2630	320	62	680	0	16	0.2		1.3	6.5	None	RH	
ı	Plastolein 9720, polyoster	69	2850	312	66	900	- 7	-23	0.3	1.8	1.9	14	None	Em	
!		Hydroc	arbon-T	yps Pl	asticis	et 8						!			
,	Dow 276-V2, polymolecular product derived from α-methylstyrene	74 E	2420	315	62	650	5	-24	1.4		0.6	22.6	None	Dow	,
	HB 40, partially hydrogenated mixture of isomeric terphenyls	62 d.	2890	298	66	940	~ 1	23	8.4		0.5	15.5	None	Mon	
١	$\mathbf{Kenplex}\ \mathbf{I}_{*}$ hydrocarbon resin made by reaction of formulduhyde + dimethylnaphthalenes												-	Ken	
	Nuvillac 10° (PHO), a hydrindyl phenol derivative	56.3°, h							13.3		0.1	0.1	None	Nev	
	Nevinol												None	Nev	
	Panaplex BN-1, alkylated aromatic hydrocarbon mixture	63 d	2800	310	66	920	2	-21	6.1		1.8	16.8	None	Pan	
8	Sovalom C, alkylated aromatic hydrocarbon mixture	61.4 4	3100	280			+ 4	-16	11.5		1.7	9.0	None	87	

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 Dush (----) 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 Flexon plasticizer DOP.
d Mixture of 1 part plasticizer to 2 parts Flexon plasticizer DOP.
d Mixture of 1 part plasticizer to 1 part plasticizer DOP.

^k Mixture of 1 part plasticiser to 3 parts Flexol plasticiser DOP.

Mixture of 0.85 part plasticiser to 1 part Flexol plasticiser DOP.

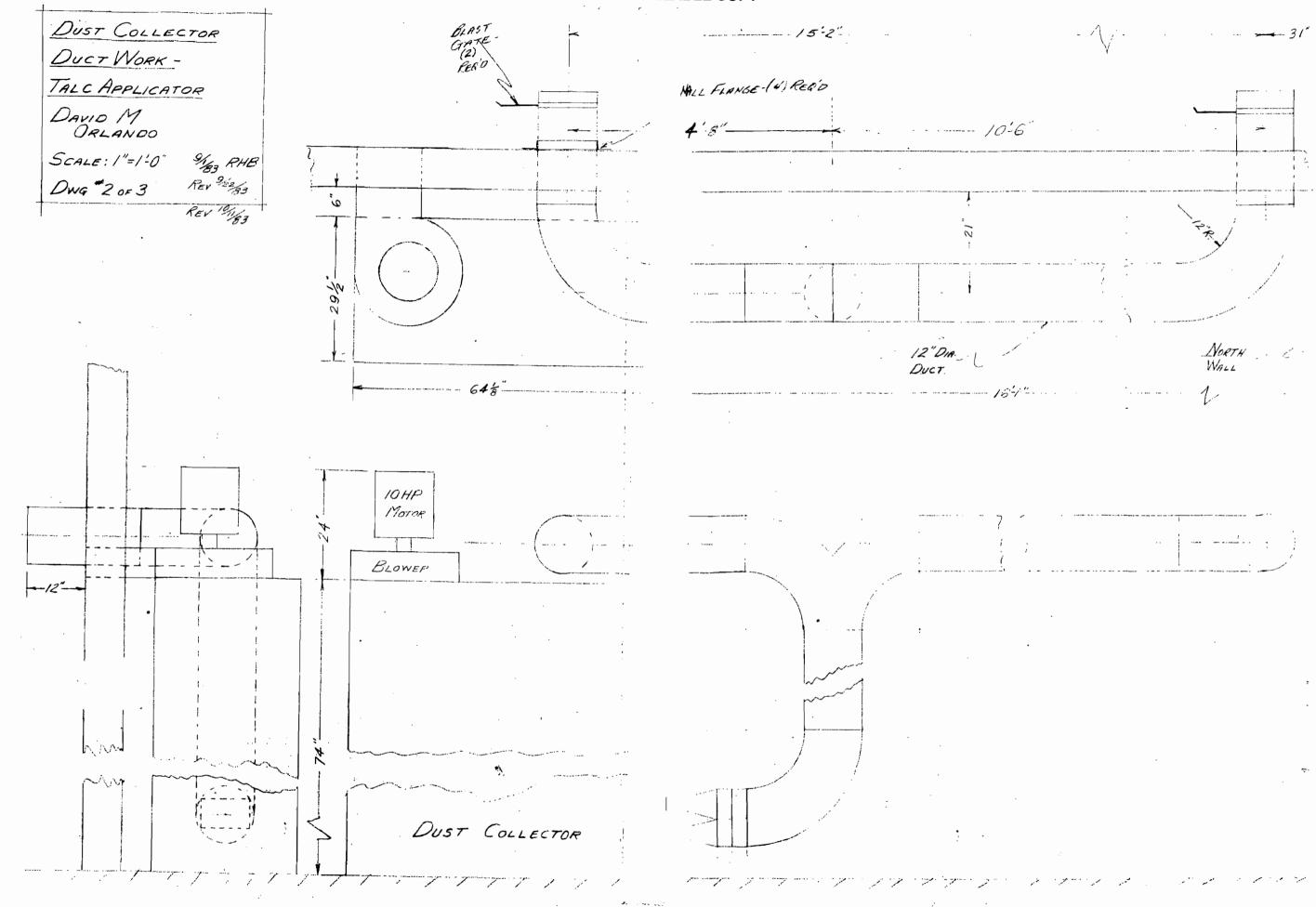
For key to supplier abbreviations see page 903.

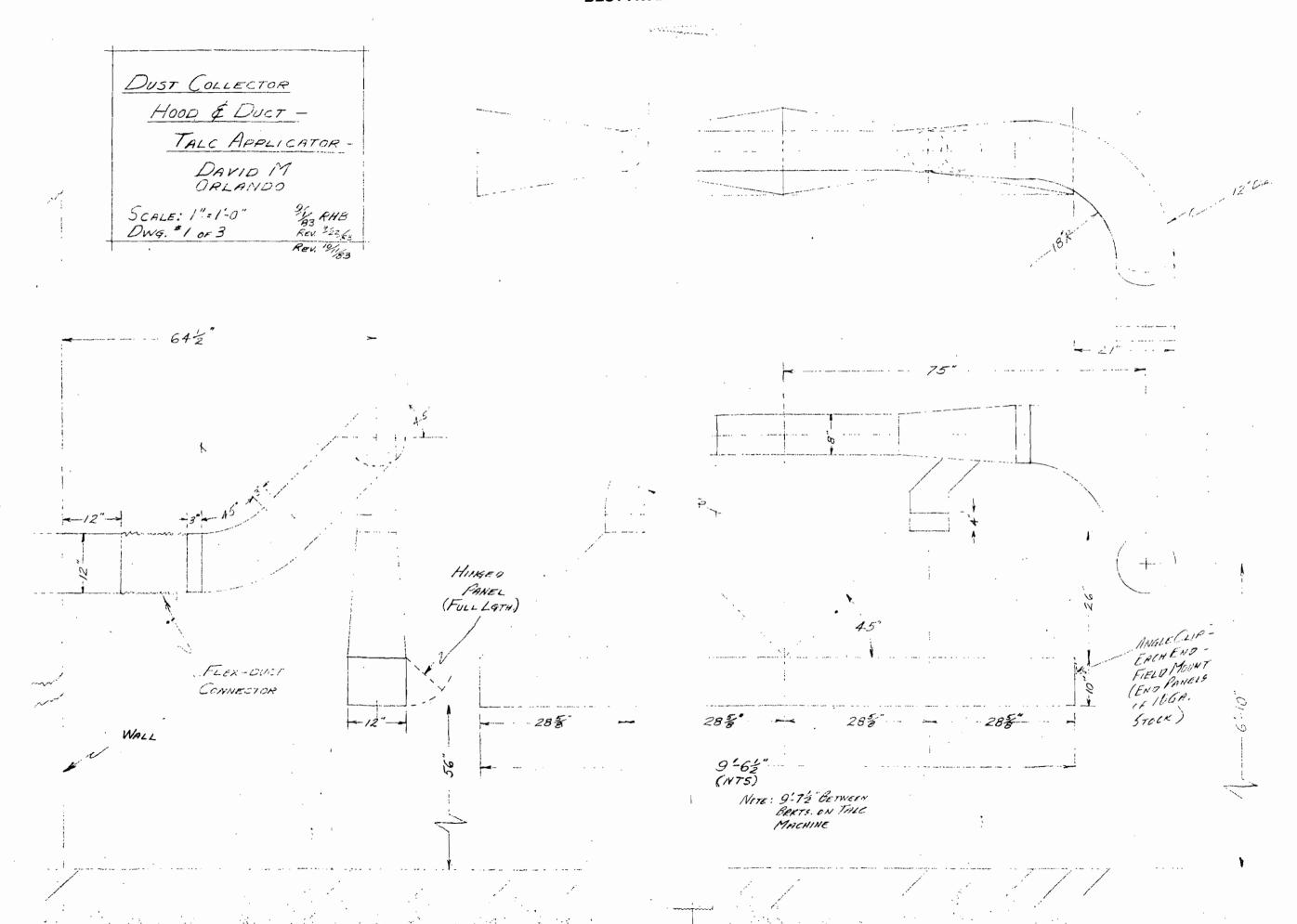
Table 15.3 (Continued) PROPERTIES OF COMMERCIALLY AVAILABLE PLASTICIZERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

				ာ ၁		J.	So	lubility,	C .	Volatility of Free Plasticis in Air, mg./sq. am./l		
No.	Name	Formula	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscosity, cp. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	875° F. (191° C.)
		M iscellaneous	 Plastic	izer e							·	
119 120 121 122 123 124 125 126 127 128 129 130 131	ARNEEL TOD, mixture of olcio, linolcio, and cyclic nitriles n-Butyl acetyl polyricinoleate, PG16 Chloriunted paraffin 40 Flexol plasticizer B-400, polypropylene glycol mono-n-butyl ether Glyceryl polyricinoleate, No. 15 oil Harflex 500, monomeric ester type Hercoflex 600, a pentacrythritol ester Ohopex Q-10, octyl fatty-phthalic acid esters l'Araflex G-00, high-molecular-weight cotor l'lasticizer DP-520, chlorinated aliphatic product Polyethylene glycol dilaurate Santicizer 180 Staflex KA, monomeric ester type		Ca. 278 362 (av.) 538 (av.)	0.910 0.913 at 25° C. 1.164 0.995 1.031 at 25° C. 0.932 1.006 0.952 0.994 1.126 at 15.6° C. 0.97 at 25° C.	1.460 st 25° C. 1.505 st 23° C. 1.449 st 20° C. 1.483 st 25° C. 1.483 st 20° C. 1.475 st 20° C. 1.474 st 20° C. 1.474 st 25° C.	30.2 3100 at 25° C. 152 17.0 48.3 41 240 to 300 66.4 cs. at 54° C.	0.1	3.3				oa. 60 0.9

		Con	nparison	of Pr	operti	es of V	inylite I Equiv	Resin V alence ^a	YN W-5	Specim	ena Plas	sticised t	to	
No.	Name	h.r. at 23° C., ion, 1000 p.s.i.	gth,	Elongation, C.	eter A	23° C., p.s.i.		Temperature, °C.	% I 4-mil	ASS, Film	10 d 23° C.	xtrao- on, lays, , 4-mil lm	60°C.	Supplier ^m .
		Plasticizer p.h.r. a 75% Elongation, 1	Tensile Strength, p.s.i. at 23° C.	Ultimate Elo % at 23° C.	Shore Durometer A at 23° C.	Stiffness at 2	Tt, °C.	Brittle Temp	10 days, 60° C.	24 hours, 70° C.	Water	150	Spew, 2 days, 60° C 100% Humidity b	٠,
		Misc	ellaneou	e Plas	icizer									
119	ARNEEL TOD, mixture of oldic, limber, and cyclic nitriles	52.8	2000	290				-44	11.7		0.7	14.8	None	Arm
120	n-Butyl acetyl polyricinoleate, PG16	1	րշուրո 	l tible a	t 55.3	p.h.r.								Bak
121	Chlorinated paraffin 40	68 .	2820	335	67	900	- 5	-28	1.0		.0	20	None	Halo, Dia, Herc, Hook
122	FLEXOL plasticizer B-400, polypropylene glycol mono-n-butyl ether							·				'	-	cccc
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	HERCOPLEX 600, a pentacrythritol ester	58	2910	327	66	800	- 2	-28	0	1.1		12.8	None	Here
126	Ohopex Q-10, octyl fatty-phthalic acid esters												-	OA
127	PARAPLEX 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	STAPLEX KA, monomeric ester type												—	Des

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 Dush (——) indicates no data available.
 A Mixture of 0.9 part plasticizer to 1 part Piexon plasticizer DOP.
 For key to supplier abbreviations see page 903.





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 % 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.

michael U- Ware

MAW:bjw Attach.

muke 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 May 15, 1984 Vs. Recovery July 1, 1984 Final Specification Development September 1, 1984 Final Specification/Quotations and Analysis December 1, 1984 Appropriations Request Development and Approval March 1, 1985 Oven Enclosure and Exhaust System Installation August 1, 1985 Solvent Recovery Delivery and Installation Operational Shakedown, Debugging and Compliance December 1, 1985 Testing



SUTCLIFFE SPEAKMAN INC

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

TELEPHONE: 301-337-2800 TELEX: 908020 240189



YOUR REF:

OUR REF: E

E 0379 S 7212 DATE 2 August 1983

PROPOSAL

F 0 R

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

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL

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.

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 injet for regulation of the airflow. 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 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

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, gaivanized 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

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-hazard-ous 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. 5 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.

Proposal No. 5 7212

David M. Company

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.

COMPRESSED AIR................20 scfh.

SPACE

The equipment, we estimate, would require a floor area of approximately 45' \times 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.

Proposal No. S 7212

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 Unasion Ichibilas S.?. Chronoly

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

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 perlod 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.



Proposal No. S 7212

BUDGETARY PRICE

(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 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.

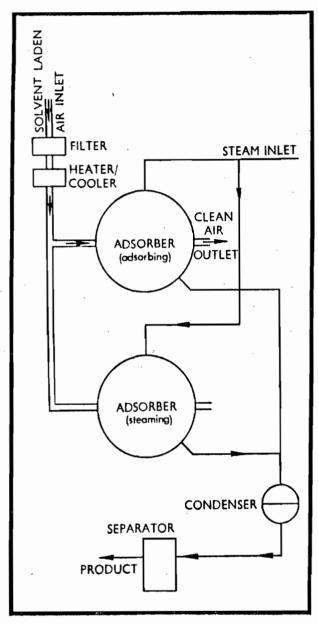
WHEN IS IT WORTHWHILE?

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 LOSS VALUE LBS/HOUR/1000 SCFM PER ANNUM X1000 CONC. % V/V 0.0 40. 50 60 0.8 0.7 0.5 0.4 175 200 0.3 225 0.2 EXAMPLE: 0-5% CON. METHYL ETHYL KETONE -M.W. 72-1 LOSS 56-5 LBS/HOUR/1000 SCFM OF AIR VALUE PER ANNUM (OF 8000 HOURS) AT 10 PENCE/LB IS £18,800 VALUE PER ANNUM (OF 8000 HOURS) AT 11-5 CENTS/LB IS \$52,000

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	В		A	Ð
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.	Perchlorethylene	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 Acetata	88.10	2.25	Tetrachlorethane	167.86	Non-flam.
Ethyl Alcohol	48.07	3.3	Tetrahydrofuran	72.10	1.84
Ethylene Dichloride	98.97	6.2	Toluena	92.13	1.3
Furfural	98.08	2.1	Trichlorethylena	131.4	Non-flam.
Heptane (Normal)	100.2	1.0	Xylene	106.16	1.0
Hexane (Normal)	86.17	1.25			

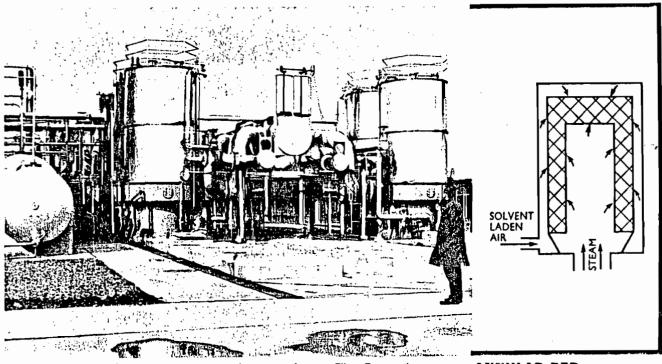
A Mol. Wt.

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.

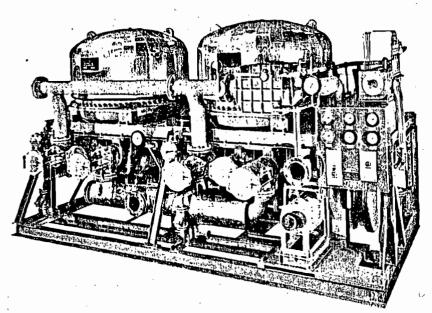
STATIC BED PLANT

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.

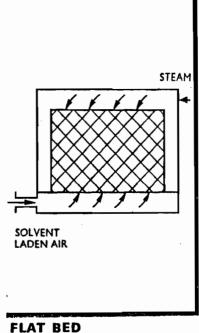


Adsorption section of a large recovery complex at an Acetate Fibre Factory in Belgium. The plant is arranged for future extension.

ANNULAR BED



Packaged type fully automatic plant of cylindrical flat bed design.

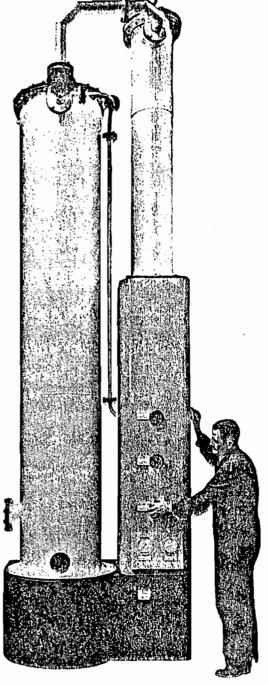


LIQUID PHASE ADSORPTION

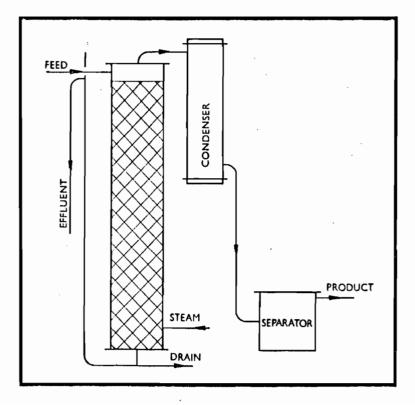
This system uses active carbon for the adsorption of small amounts of certain water-miscible solvents from solutions with water. The weak solvent/water liquor is passed through an active carbon bed and, when the carbon has adsorbed the designed charge of solvent, the solvent is distilled off by our patented 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.

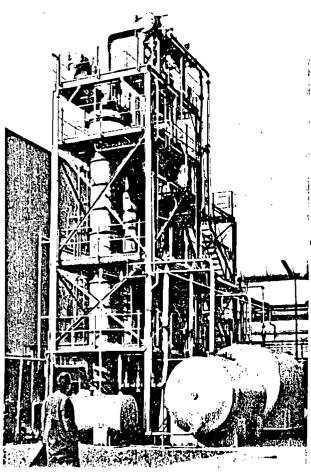
A particular feature of this system is the extreme simplicity of operation and control which can be manual or automatic.



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

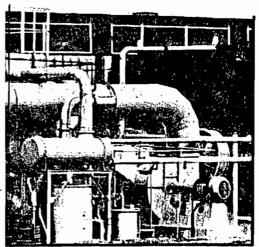
SUITE 200, HEAVER PLAZA, 1301 YORK RD. LUTHERVILLE, MARYLAND 21093 TELEPHONE: 301-337-2800 TELEX: 908020

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 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	Concen- tration (%V/V)	*Initial adsorp- tion (%W/W)	§Cyclic adsorp- tion (%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	6 6
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 3, 3, 5-trimethyl cyclohexyl	0.3	35.6	11.9	3.7	153-193
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.

SENSOR CONTROLS

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:

(a) Inlet concentration 1000 ppm Outlet concentration 35 ppm

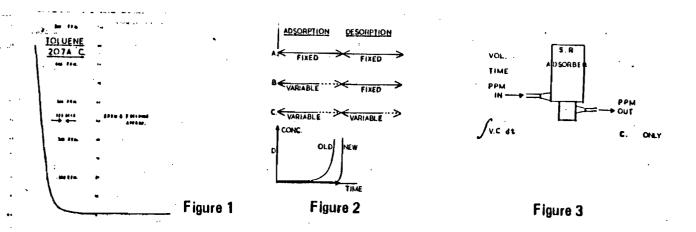
$$\Upsilon = \frac{1000 - 35}{1000} \times 100\% = 96.5\%$$

(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.



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

. . .

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

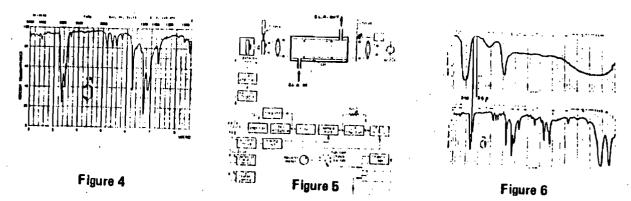
as solvent laden air enters the adsorber we can measure solvent concentration and flow rate, and compute, by integration with respect to time, the total quantity of solvent passed to the adsorber. When a predetermined amount of solvent has been passed a signal is generated which is used to change over 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 1-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



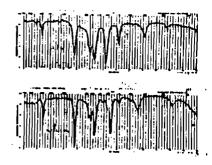
being sensitive to background absorptions caused by fouling of the sample cell windows, broad band absorptions in the sample gas (such as that caused by water vapor), and temperature differences between sample and reference beams. However, most of the disadvantages of the two beam system can be eliminated 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 - 1.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.





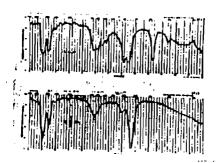


Figure 7

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 stronly, 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 absorbtion 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 absorbtion bands.

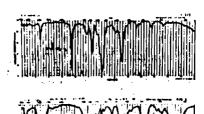
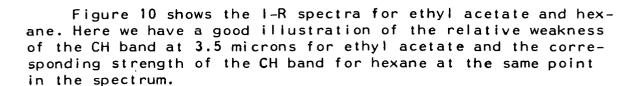


Figure 10



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.

- (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.



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

TELEX: 9

988020 240189

YOUR REF:

OUR REF:

DATE: 4 August 1983

DESIGN FOR EXPANDINATION !

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 galions 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



Wheelabrator-Frye Inc. **Chemicals & Coatings Group**

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

October 10, 1983

H. Fancy, P.E. Deputy Chief Bureau of Air Quality Management Twin Towers Office Building

DER

DCT 1 3 1983

9000 Blair Stone Road Tallahassee, FL 32301-3241

BAOM

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.

Michael A. Ware

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

nechael a. Ware

MAW: bj w

cc: Wayne Brady

R. H. Baddeley

J. Spangler

H. Mycroft

T. Lucas

J. Seabury

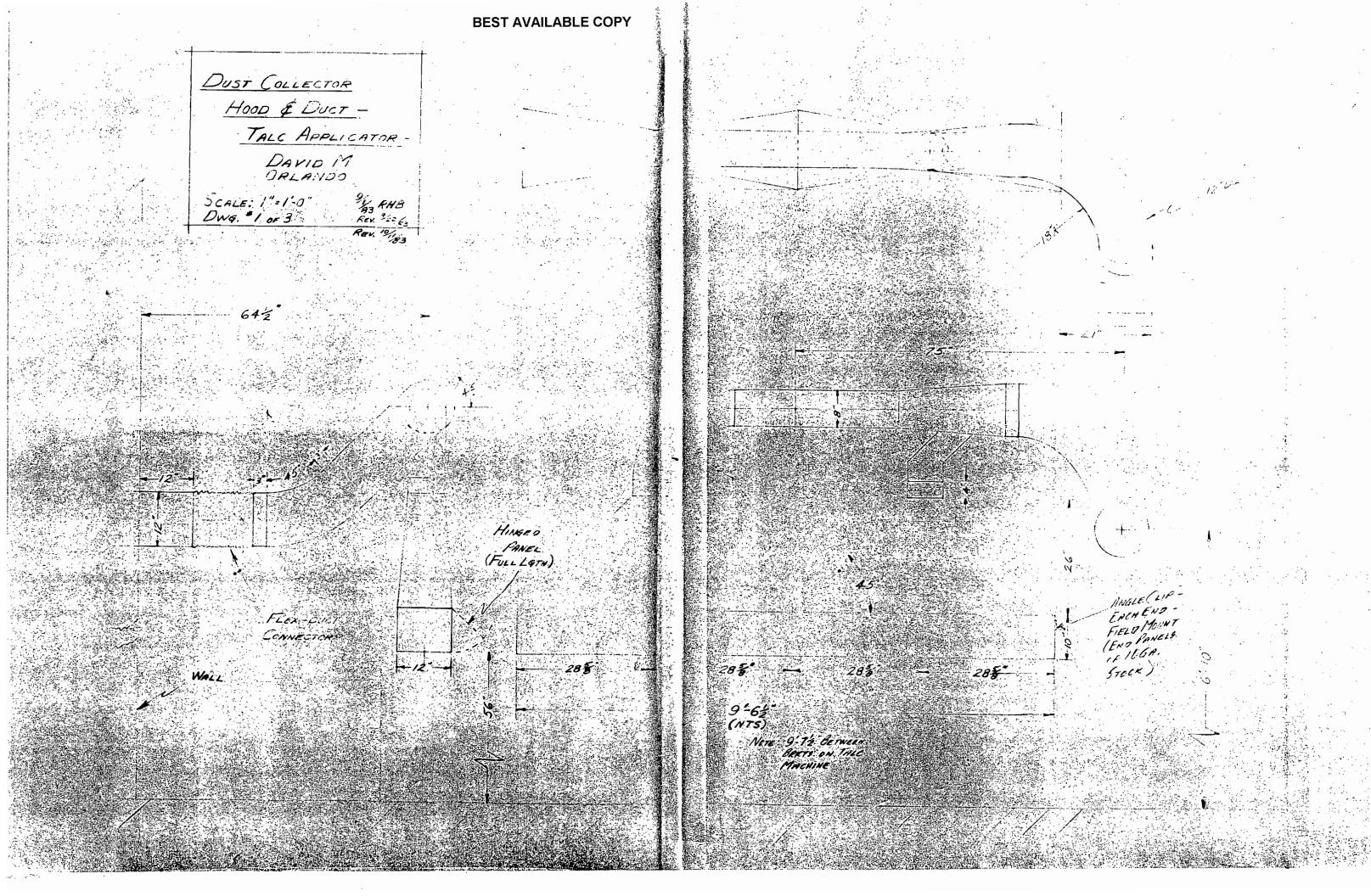
C. Collins





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

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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 1 2 1983 BAOM

SUBJECT: Your Memorandum of August 12, 1982 Concerning VOC Permits, Boiler Permitsand 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. 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.

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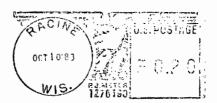
MAW: bjw

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)

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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.
- 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 (1b/1b)?
- 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

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

008020 240189 TELEX:



YOUR REF:

OUR REF:

E 0379 S 7212

DATE: 2 August 1983

PROPOSAL

F 0 R

SOLVENT RECOVERY SYSTEM

TO BE LOCATED AT:

DAVID M. COMPANY

201 VALENTINE WAY

LONGWOOD, FLORIDA



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

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL

598 the 2000# 16x250h 299 MA

(304/hu)



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.

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. (' ' ' '

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.

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 \times 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-hazard-ous area, assumed within 50' of the adsorbers.

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

David M. Company

Proposal No. 5 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.



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. 5 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.

Proposal No. S 7212

David M. Company

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.

POWER.....100 KwH.

COMPRESSED AIR..........20 scfh.

SPACE

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

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.

Proposal No. S 7212

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 yoursen Inhibitor is? Chumolis

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.

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.

Proposal No. S 7212

BUDGETARY PRICE

(ONE HUNDRED SIXTY THOUSAND DOLLARS)

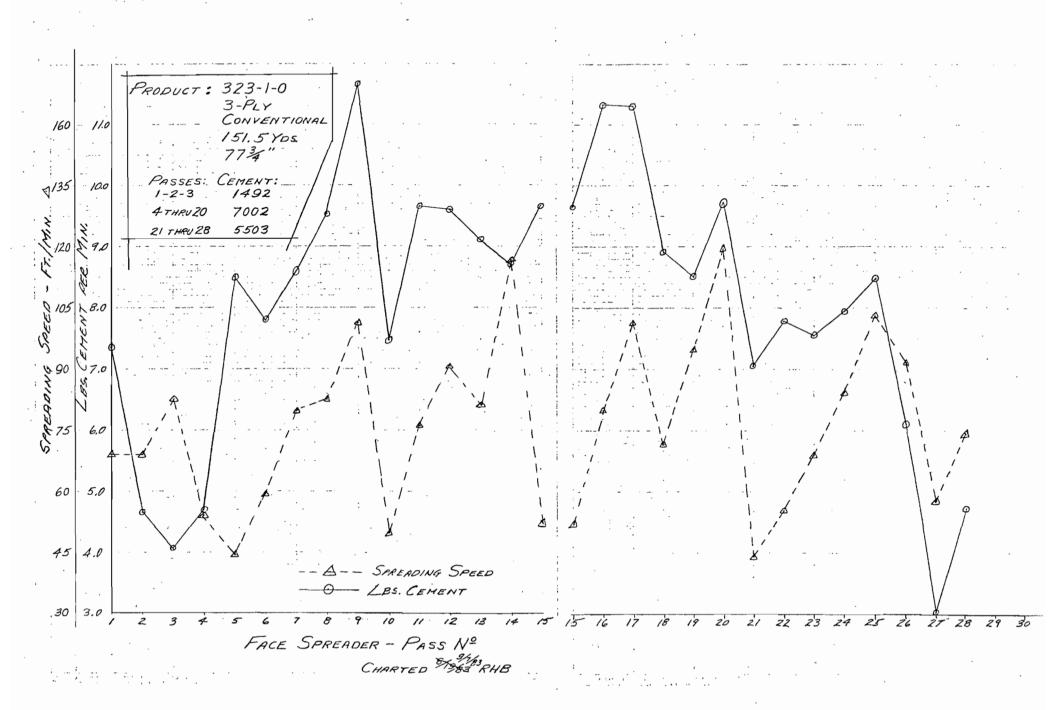
COMPLETION

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.

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41,000 fer paid STATE OF FLORIDA ENVIRONMENTA ENVIRONMEN ST. JOHNS RIVERY BOB GRAHAM AUG 1 2 1983 GOVERNOR VICTORIA J. TSCHINKEL SECRETARY 3319 MAGUIRE BOULEVARD SAINT JOHNS SUITE 232 ORLANDO, FLORIDA 32803 ALEX SENKEVICH DISTRICT MANAGER RIVER DISTRICT APPLICATION TO OPERATE/CONSTRUCT AIR POLLUTION SOURCE SOURCE TYPE: Blanket Spreading Operation [x] New [] Existing AUG 24 1983 APPLICATION TYPE: [x] Construction [] Operation [] Modification COMPANY NAME: DAVID "M" COMPANY Identify the specific emission point source(s) addressed in this application (i.e. Lime Spreaders #1,#2,#5 Kiln No. 4 with Venturi Scrubber; Peaking Unit No. 2, Gas Fired) Festooning and Mixing are SOURCE LOCATION: Street 201 Valentine Way City Longwood North 3177160 UTM: East 17-470106 Longitude 81° 18' Latitude 28 ° 43 ' 26 "N 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 to the city or a sali APPLICANT ut a unification o 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 contro 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 of legal transfer of the permitter establishment.

*Attach letter of authorization

Signed: Président & 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 hav: 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)

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	,	John W. Seabury
		None (Please Type) (2) (2) (2) (3) (3) (4) (4) (4) (4) (4) (4) (4) (4) (4) (4
		Seabury-Bottorf Associates, Inc. 2000 Company Name (Please 1999)
		3702 Silver Star Rd., Orlando, Florida 32808
•		Mailing Address (Pless Type)
lda Registration No	8719	Date: August 11,1983 Telephone No. 305/298-0846
		GENERAL PROJECT INFORMATION
•		of the project. Refer to pollution control equipment,
necessary.		in full compliance. Attach additional sheet if
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	this is a new source or major modification, answer the following quest	ions.
ı.	Is this source in a non-atteinment area for a particular pollutant?	No
	e. If yee, hes "offset" heen applied?	No
	b. If yes, has "Lowest Achievable Emission Rate" been applied?	No
	c. If yee, list non-attainment pollutents.	No
2.	Does best svailable control technology (BACT) apply to this source? If yes, see Section VI.	Yes
3.	Dase the State "Prevention of Significant Deterioriation" (PSD) requirement apply to this source? If yes, see Sections VI and VII.	No
4.	Do "Stenderds of Performance for New Stationary Sources" (NSPS) apply to this source?	No
5.	Do "National Emission Standards for Hezardous Air Pollutants" (NESHAP) apply to this source?	No
	"Reasonably Aveilable Control Technology" (RACT) requirements apply this source?	No
	a. If yes, for what pollutents? N.A.	

Attach all supportive information related to any answer of "Yes". Attach any justification 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).

any information requested in Rule 17-2.650 must be submitted.

SECTION III: AIR POLLUTION SOURCES & CONTROL DEVICES (Other them Ingineratore)

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

Description	Contami	lnent =	Utilization			
	Type	% Wt	Rete - 1be/hr	Relate to Flow Diagram		
Rubber	None Known		Variable	Used to formulate cements		
Rubber Cement	V.O.C.	30-70%	Variable	Mixing (2), Spreading (3,4,5), Festooning (6		
	MEK Toluene			(At step #6 99%+ of V.O.C. has been flash		
			-	off)		
	Territoria de la composición della composición d		*			

*The process utilizes rubber cements which are mixtures of various rubber compounds i.e.(nitrile, e 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.

- 1. Total Process Input Rate (lbe/hr): Variable
- 2. Product Weight (lbe/hr): Variable
- C. Airborne Conteminante Emitted: (Information in this table must be submitted for each emission point, use additional sheets as necessary)

Name of	Emios	full fuer	Allowed ² Emission Rate per	Alloweble ³ Emission	Potentia Emissionia	14 16 Jan Jan	Relate to Flow
Conteminent	Meximum lba/hr	Actuel T/yr	Rule 17-2	lbs/hr	lbs/yr	T/yr	Diagram
VOC - MEK/	103	84	BACT	To be determined	420,000	210	1
" Toluene	5	4	BACT	To be determined	20,000	10	3
11	94.5	75.6	BACT	To be determined	378,000	189	4
11	94.5	75.6	BACT	To be determined	378,000	189	5
	297.0				morar some	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.

²Reference applicable emission standards and unita (e.g. Rule 17-2.600(5)(b)²2. Table II, E. (1) - 0.1 pounds per million BIU heat input)

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Calculated from operating rate and applicable standard.

AEmission, if source operated without control (See Section V, Item 3)

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O. Contrai	Devices:	(See Section	٧,	Item	4)
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Neme and Type (Model & Seriel No.)	Conteminent	Efficiency	Range of Particles Size Collected (in microns) (If applicable)	Basis for Efficiency (Section V Item 3)
'1				
<u></u>			,	:

F. Funla

	Consum	ption*	
Type (Be Specific)	avg/hr	mex./hr	Meximum Hest Input (MMSTV/hr)
			-
		The second se	

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

Fuel Analysis:

Porcont Sulfur:_	N.A.		Percent	Aeh:	N.A.	·	
Deneity:	N.A.	lbe/gel	Typicel	Percent	Nitrogen:	N.A.	·
Heat Capacity:	N.A.	8TU/1b		N.A.	·	·	BTU/ge1
Other Fuel Conte	minente (which way	cause sir p	ollution;):	N.A.		
	,				· · · · · · · · · · · · · · · · · · ·		

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

Annual Average	N.A.	Maximum	N.A.	

8. Indicate liquid or eolid 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.

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tack Height:					1			
es flow Rate:	ACFH	DS(CFH Ga	Gae Exit Temperatures				
ater Vapor Cente	ntı <u> </u>		* Y.	locity:		<u></u>		
						4 9		
	SECT	ION IA! INC		•	OM	· · · · · · · · · · · · · · · · · · ·		
		Not Ap	opl ica b	Le	<u> </u>	<u> </u>		
Type of Type (Plast	O Type I (Rubbieh)	Type 11 Ty (Refuee) (Ga	ype IIE erb ege)	Type IV (Patholog- ical)	Type V (Liq.& Gea By-prod.)	(Solid By-prod.)		
Actual lb/hr nciner- ated								
Uncon- trolled lbe/hr)			-					
otal Waight Inci	nersted (1be/h	-				/hr)		
otal Waight Inci pproximate Numbe	nersted (lbs/h	Operation per	r dey _	day/	wk	wke/yr.		
•	nersted (1be/h	Operation per	r day	day/	wk	wke/yr.		
otal Waight Inci pproximate Numbe anufacturer	nersted (lbs/h	Operation per	r day	day/	wk	wke/yr.		
otal Waight Inci pproximate Numbe anufacturer	r of Hours of Volume (ft)	Operation per	r day	dsy/	wk	wke/yr		
otal Waight Inci pproximate Numbe enufacturer ate Constructed	r of Hours of Yolume (ft)	Operation per	r day	dsy/	wk	wke/yr		
otal Waight Inci pproximate Numbe anufacturer ata Constructed Primary Chamber Secondary Chambe	ref Houre of Volume (ft)	Operation per Heat Relac (BTU/hr)	Model	No. Fuel	BTU/hr	Temperature (°f)		
proximate Number and Foregree Chamber Secondary Chamber Lack Height:	volume (ft)	Heat Relace (BTU/hr)	Model	No	BTU/hr Stack T	Temperature (°F)		
pproximate Number and Flow Reter	volume (ft)	Heat Relea (BTU/hr) Stack Diamter ACFM	Model	JSCFM*	BTU/hr Stack T	Temperature (°F)		
pproximate Number and Footurer Ate Constructed Primary Chamber Secondary Chamber tack Height: Ate Flow Rate:	volume (ft) ft.	Heat Relea (BTU/hr) Stack Diamter ACFM ign capacity, ed to 50% exc	Model	DSCFM*	Stack T Velocity:	Temperature (*F) empFF		

20.25 PT H. Emission stack geometry and flow characteristics: From estimates Emission Point <1> (SEE process floor diagram): Stack Dimensions: 4.5 x 4.5 square (ft.) Stack Height: 18 ft. Gas Flow Rate: 12,600 ACFM 9960 DSCFM Cas Exit Temp.: 150 (F) Velocity: 699 FPS Water Vapor Content Ambient (variable)% 622.2 FPM Emission Point <3> (See process flow diagram): Stack Dimensions: 1.5 x 1.5 square (ft.) Stack Height: 12 ft. Gas Flow Rate: 1575 ACFM 1410 DSCFM Gas Exit Temp.: 95 (°F) 700 FPM Water Vapor Content Ambient (variable)% Velocity: 700 FAS Emission Point 4 (See process flow diagram): Stack Dimensions: $(2(1/33 \times 1.33 \text{ outlets}))$ (ft.) Stack Height: 12.5 ft. Gas Flow Rate: 1415 ACFM 1120 DSCFM Gas Exit Temp.: 150 (°F) 400 1685 Water Vapor Content: Ambient (variable)% Velocity: 800 FRS Emission Point (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 EPS = 622,2 FEM Natraily = 12,600 FT3

Discharge points (2) (Mills) missing

Brief deacription of operating character	ri stics of contro	ol devices:	Not A	ppliable	
		1			
			,		
				,	
·	A Thair and Title against a the Control of the Cont	***************************************	· i · ·		
Ultimate disposal of any effluent other eah, etc.):	then that emitte	d from the	eteck	(ecrubber wet	.et,
Not 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)]

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

- NOT APPLICABLE

 To a construction application, attach basis of amission astimate (e.g., design calculations, design drawings, pertinent manufacturer's test data, atc.) and attach proposed mathods (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 control co
- NOT APPLICABLE

 5. With construction permit application, attach derivation of control device(a) afficiency. Include test or design data. Items 2, 3 and 5 should be consistent: actual emissions a 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 rew meterials enter, where solid and liquid waste exit, where gaseous emissions and/or sirborns particles are evalved and where finished products are obtained.
- SEE APPENDIX 2

 7. An 8 1/2" x 11" plot pien showing the location of the establishment, and points of sirborns 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 abowing the location of manufacturing processes and outlate for airborna amissions. Relate all flows to the flow diagram.

SEE APPENDIX 3

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1

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9.	The appropriate	application fee	in accord	ance with	Rule 17~	4.05. The	check	should	be
	made payable to	the Department	of Environ	mental Reg	ulation.	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

4

SECTION VI: BEST AVAILABLE CONTROL TECHNOLOGY

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	* F 19)					
N.A.	N.A						
		711					
B. Has EPA declared the best available cont yes, attach copy)	rol technology for this class of sources (If) - 4 k - 25 N					
[] Yes [X] No		*					
Contaminant	Rate or Concentration						
BACT has not been declared for this	class of sources.						
		. •					
C. What smission levels do you propose as be	st available control technology?	. 1					
Contaminant	Rate or Concentration						
VOLATILE ORGANIC COMPOUNDS	AVERAGE COATING VOC CONTENT OF \$5.411	s./ga					
	VOC/gal. OF COATING DELIVERED TO COAT	TING					
	APPLICATOR.						
		ξ.,					

- 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

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Useful Life: Operating Costes Energy Maintenance Cost: Emissions: Conteminent Rate or Concentration 10. Stock Peremeters Height: ft. Diameter: ft. Flow Rete: . ACFH d. Temperature: Velocity: FPS Describe the control and treatment technology available (As many types as applicable, use additional pages if necessary). LOW SOLVENT TECHNOLOGY CONTROL VOC EMISSION Operating Principles: BY FORMULATION Control Device: FORMULATE COATING TO b. GIVEN SPECIFICATION Efficiency: NOT APPLICABLE Cepitel Coet: ESTIMATE \$10,000.00 Useful Life: CONTINUING Operating Cost: \$20,000.00 , f. g. Energy: 2 NOT APPLICABLE Maintenance Cost: NOT AVAILABLE Availability of construction materials and process chemicals: AVAILABLE Applicability to manufacturing processes: APPLICABLE Ability to construct with control device, install in available epace, and operate within proposed levels: ABLE TO INITIATE 2. CATALYTIC INCINERATION Control Device: CATALYTIC INCINERATOR b. Operating Principles: CATALYZED INCINCERATION Efficiency: 1 90% + Capital Cost: \$500,000.00 Operating Cost: NOT AVAILABLE AT THIS WRITIN Useful Life: 10-15 YEARS ſ. Energy: 2 NOT AVAILABLE FROM BUDGET h. Meintenance Coet: NOT AVAILABLE QUOTATION Availability of construction materials and process chamicals: AVAILABLE Explain method of determining efficiency.

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²Energy 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
- e. Control Device: CARBON ADSORBER
- b. Operating Principles: PHYSICAL ADSORPTION

c. Efficiency: 1 85-90%

- d. Cepitel Cost:\$600,000.00-\$700,000.00
- e. Ueeful 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 chamicals: 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:

. Usoful Life:

f. Operating Cost:

q. 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 epace, 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. Cepitel Coet: \$10,000.00
- 4. Ueeful Life: CONTINUING
- 5. Operating Coat: \$20,000.00
- 6. Energy: NOT APPLICABLE
- 7. Maintenance Coet: NOT APPLICABLE
- 8. Manufacturer: NOT APPLICABLE
- 9. Other locations where employed on similar processes: NONE KNOWN
- e. (1) Company: NOT APPLICABLE
- (2) Mailing Address:
- (3) City:

(4) State:

1Explain method of determining efficiency.
2Energy to be reported in units of electrical power - KWH design rate.

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(5) Environmental Manager:	
(6) Telephone No.:	
(7) Emissions:1	
Conteminent	Rate or Concentration
(8) Process Reter ¹	
b. (1) Company:	
(2) Heiling Address:	en e
(3) City:	(4) State:
(5) Environmental Manager:	
(6) Telephone No.:	
(7) Emissions:1	
Conteminent	Rate or Concentration
(8) Process Rate: 1	
10. Remon for melection and descripti	on of systems:
Applicant must provide this information we available, applicant must state the reason SECTION VII - PREVENTION NOT APPLICA	O(a) why. OF SIGNIFICANT DETERIORATION
A. Company Monitored Data	
1no. eiteeTSP	() SO ² Wind spd/dir
Period of Monitoring month	day year month day year
Other date recorded	
Attach all data or statistical summarie	a to this application.
*Specify bubbler (B) or continuous (C).	
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:

	2.	Inetrumer	ntetion, Field a	nd Laboratory				
•	٠.	Was insti	rumentation EPA	referenced or its	equivalent?	[] Yes	[] No	
	b.	Was Insti	rumentation cali	brated in accorda	nce with Dep	artment p	raceduree?	
		[] Yes	[] No [] Unk	กงพก				
8.	Het	eorologica	al Data Used for	Air Quality Mode	ling		·	•
	1.	Yee	er(e) of data fro	om // month day ye	to manth	/ / day yaa	,	
	2,	Surface d	lata obtained fro	om (location)				
	3.	Upper qir	(mixing height)) data obtained f	rom (locatio	n)	·	
	4.	Stability	wind rose (STA	R) data obtained	from (locati	on)		
c.		puter Mode					•	
	1.		<u> </u>		Modified?	lf yes,	attach des	cription.
	Att		of all final me	odel rune showing				
D.	App.	licante He	ximum Allowabla	Emission Date				
	Pol	lutent		Emission Rate			·	
		TSP			gr	988/880		
		50 ²	1		gr	ams/sec		•
Ε.	Emi	ssion Data	Ueed in Modelin	ng				1
•	pole	nt source		ces. Emission de number), UTM cooi				
F,	Att	ach all ot	her information	supportive to the	e PSD review	•		
G.				mic impact of the oa, payroli, pro				

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H. Attach acientific, engineering, and technical material, reports, publications, jour-nals, end other competent relevant information describing the theory and application of

assessment of the environmental impact of the sources.

the requested best svailable 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. = $\frac{77.4 \text{ tons/yr.}}{}$

Spreaders #2 and #3

Maximum VOC loading in 1bs./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

24 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 \(\frac{1}{2} :

Emission point \bigcirc 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.

Emission Point 1 Total = (spreader (# 1: 77.4 T/yr.) + (spreaders (2 and 3) 504 T/yr. x .25) + (festooning 5 tons/yr.) + (mixing general 2.0 T/yr.) = 210 T/yr.

Emission Point 🔇 :

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 T/yr. x .75 =
$$189 \text{ tons/yr}$$
.

Emission Point <5 :

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

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

Maximum Lbs./Hr. Calculation

Emission Point 🗘 :

- 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 min./hr. x 4.3 lbs./min x 1.0 = 38.7 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 min./hr. x 14.0 x .25 = 31.5 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 min./hr. x 14.0 x .25 = 31.5 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.

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%

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%

Actual Tons/Yr.

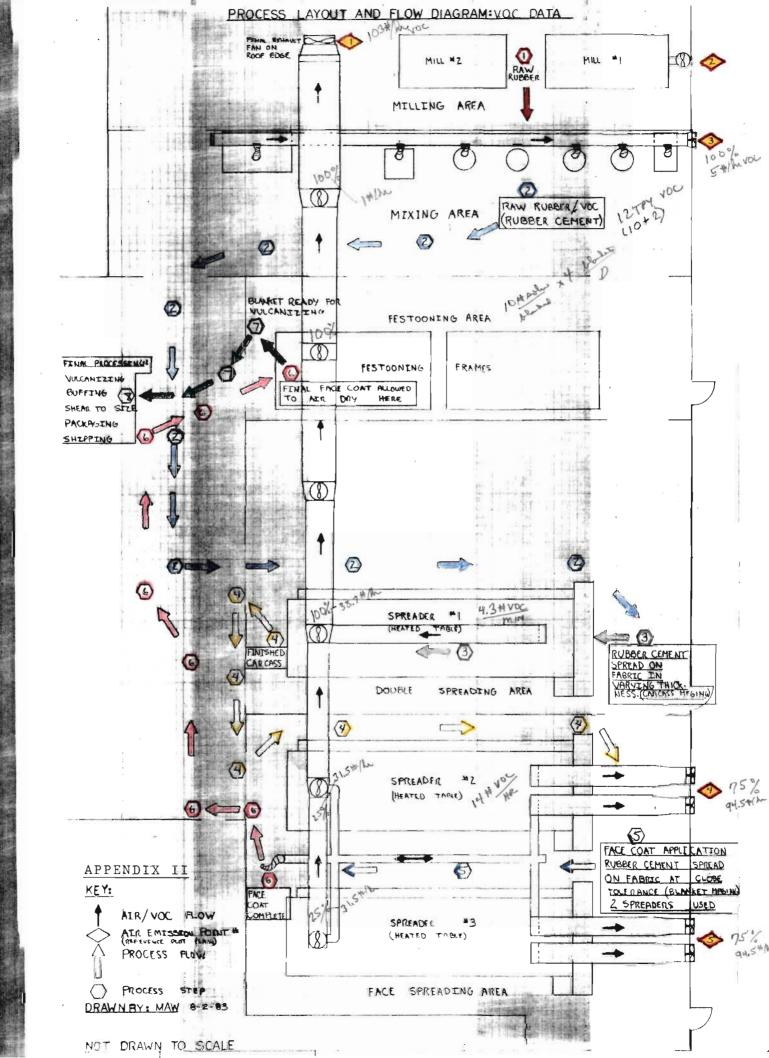
The facility currently operates at approximately 40% of its emitting potential. Therefore, potential T/yr. are multiplied by .40 to obtain actual emissions.

Emission Point	Potential Emissions	Actual Emissions	
	210 T/yr.	84 T/yr.	
③	10 T/yr.	4 T/yr.	
4>	189 T/yr.	75.6 T/yr.	
\$	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.

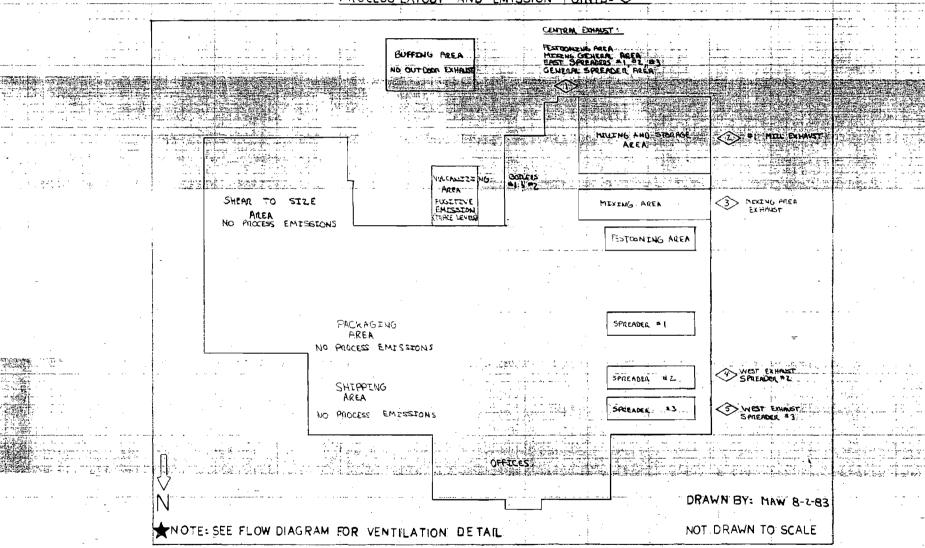
APPENDIX II CONT. PROCESS FLOW EXPLANATION

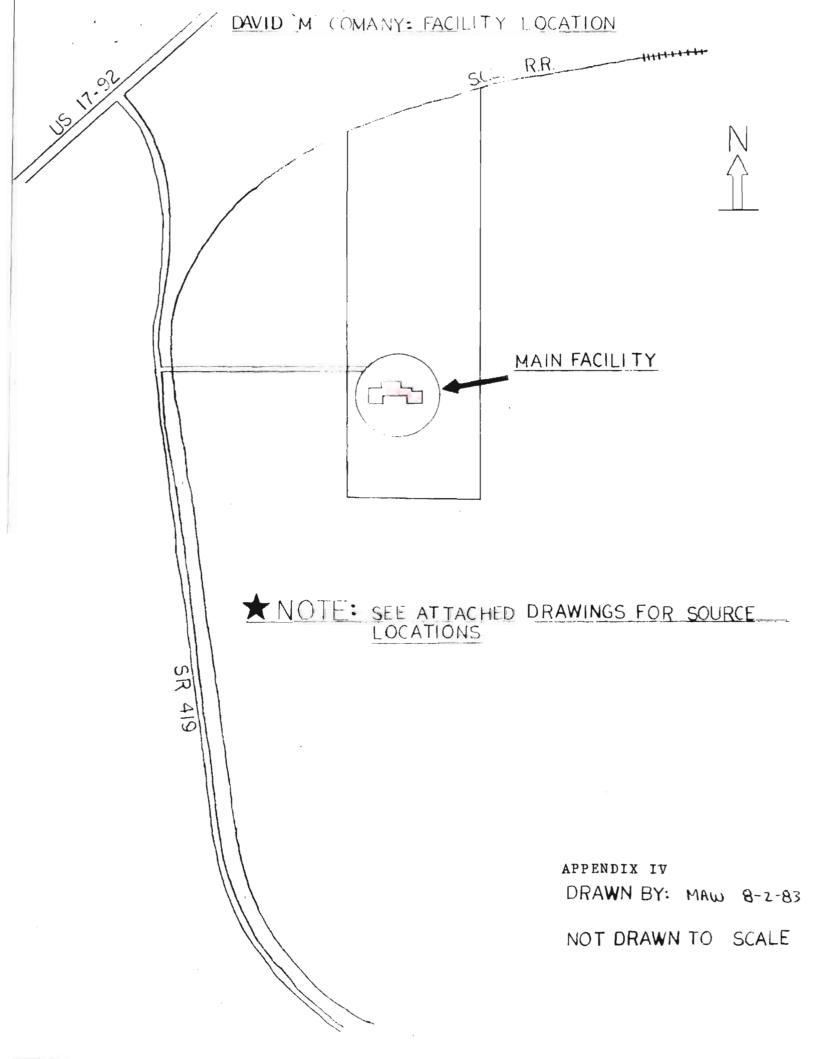
Step ①	Rubber is milled into sheets for use in cement making. No calculation of emissions is included in this application since only particulate emissions would be regulated and this source is a minor one which will be dealt with separately from VOC emissions.
Step Q	Rubber is combined with VOC (MEK/Toluene) and mixed to the de-
Step 🐧	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 🕜	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. 50°
Step (8)	Following final cure the blanket may be buffed or sheared and then packaged for shipping.



Best Available Copy

APPENDIX III PROCESS LAYOUT AND EMISSION POINTS:







COMPOUND	# SAMPLES ANALYZED	AVG LBS. TOLUENE/100 LBS.	AVG LBS RUBBER/
7 0 02	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.

COMPOUND	<u>ĤI</u>	LO	AVERAGE	NUMBER TESTED
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.

ofor row

4100g

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 > .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.

Falc

c)

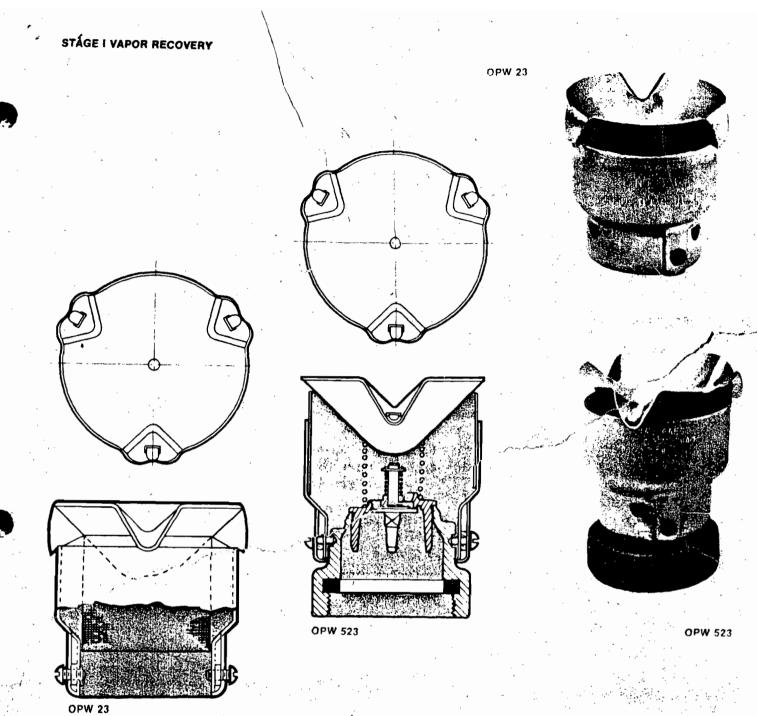
2 mill

How mu

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

Ho LWILLS NE

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



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 $1\frac{1}{2}$, 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 $\frac{1}{2}$ 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.



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 0:3 1983 CAQM

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

Graphics Div. Env. Coord. Chemicals & Coatings Group

nichael a. Wave

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:

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

no process in

30 Loursym

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

Proceed Tolvene x 7.24/gd = 298.8 Tims

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 1bs/1b?

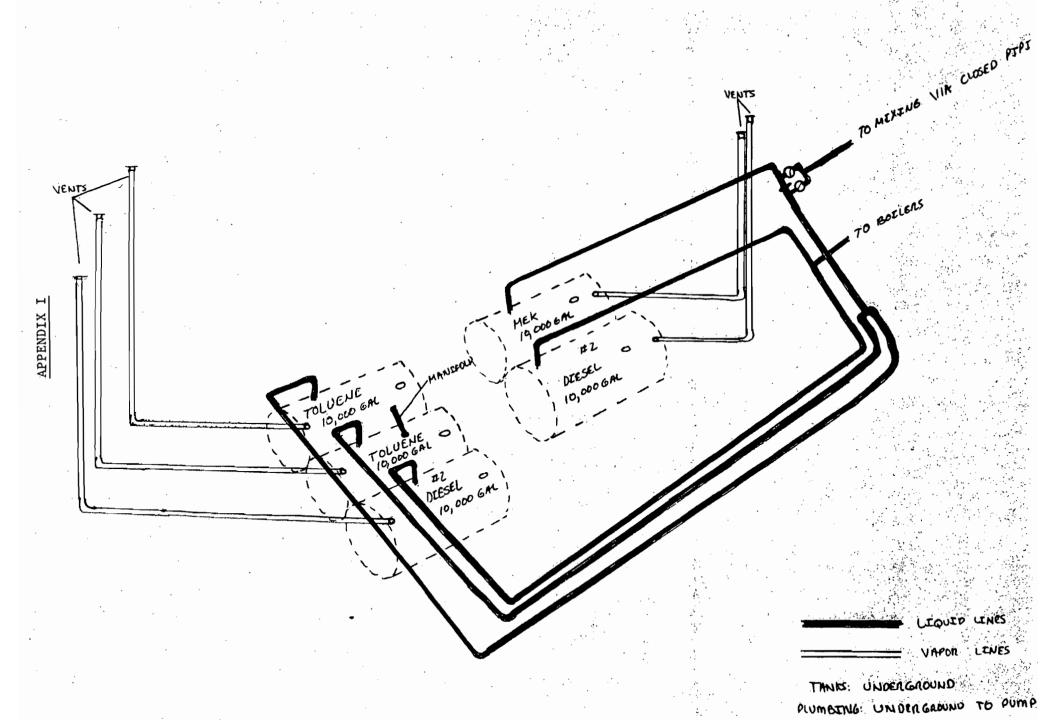
The chart below outlines the mixing ratio of solvent to rubber on a 100 lb. batch basis.

Sterney founds

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COAXIAL

SECTION B

Coaxial Vapor Recovery Equipment For use in stage I and stage II systems

System C

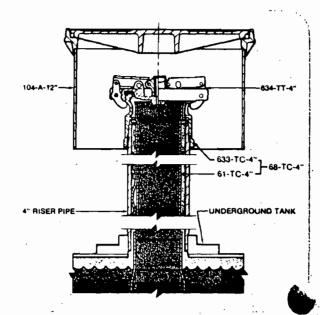
TYPE 1 - 68-TC - 3" & 4" Series Fittings
(Non Poppeted)

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

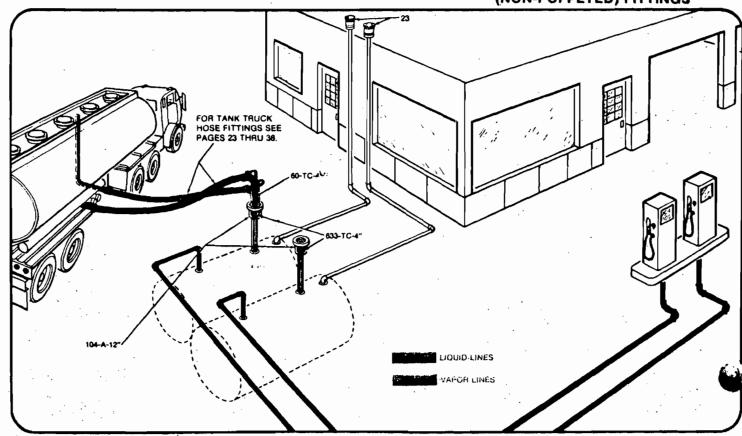
An easy, inexpensive means to convert existing conventional 3" & 4" size underground storage tank fill pipes to coaxial installations. Normally little to no digging or tearing up of concrete is necessary. Simply remove existing tight fill adaptor and fill pipe (if installation is so equipped), inserting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

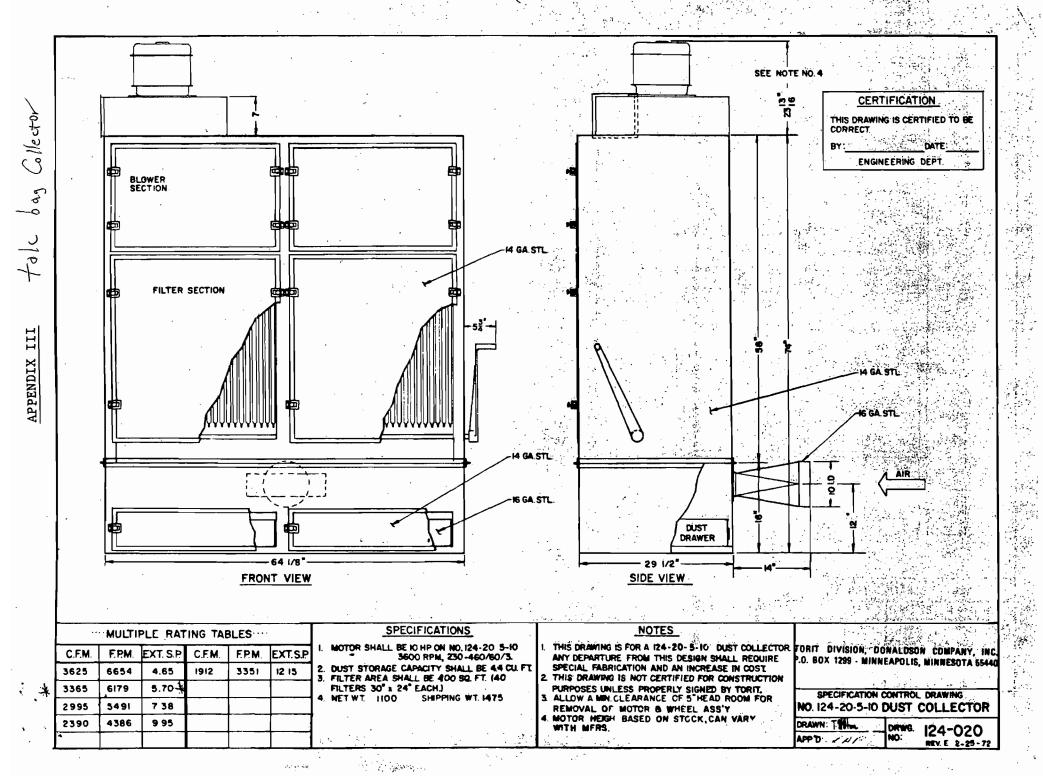
NOTE: Type 2 poppeted fittings are certified by California Air Resources Board (CARB)

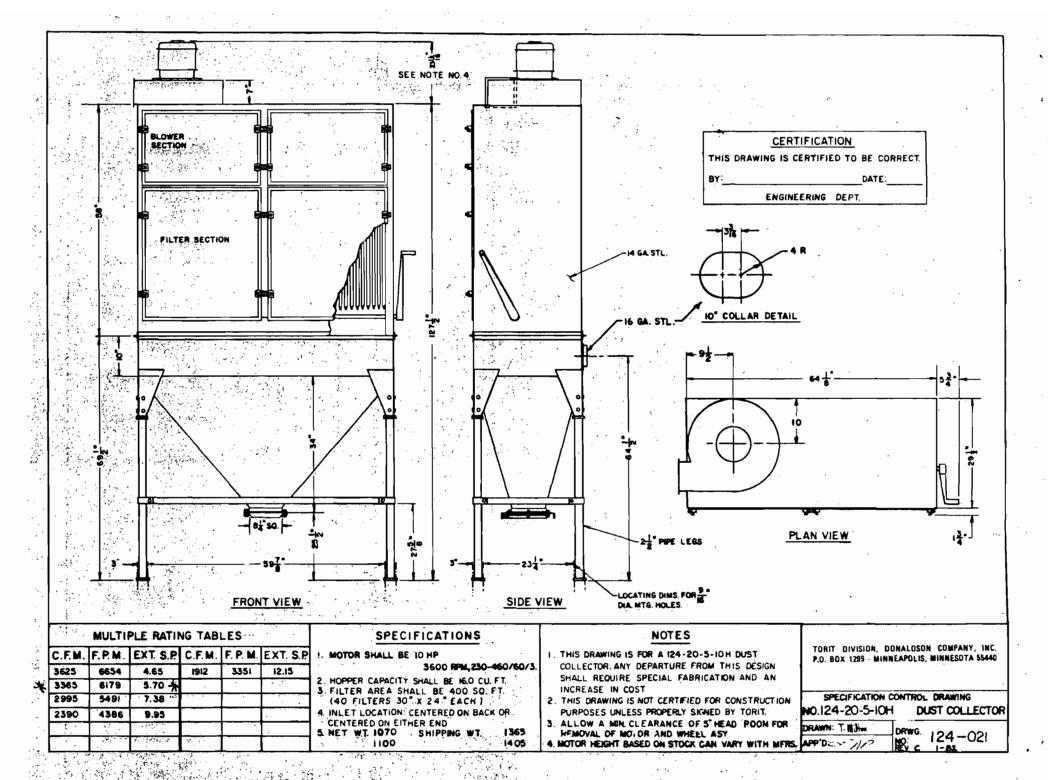
SHOWN BELOW IS A TYPE 1 - 68TC-4" TYPICAL INSTALLATION



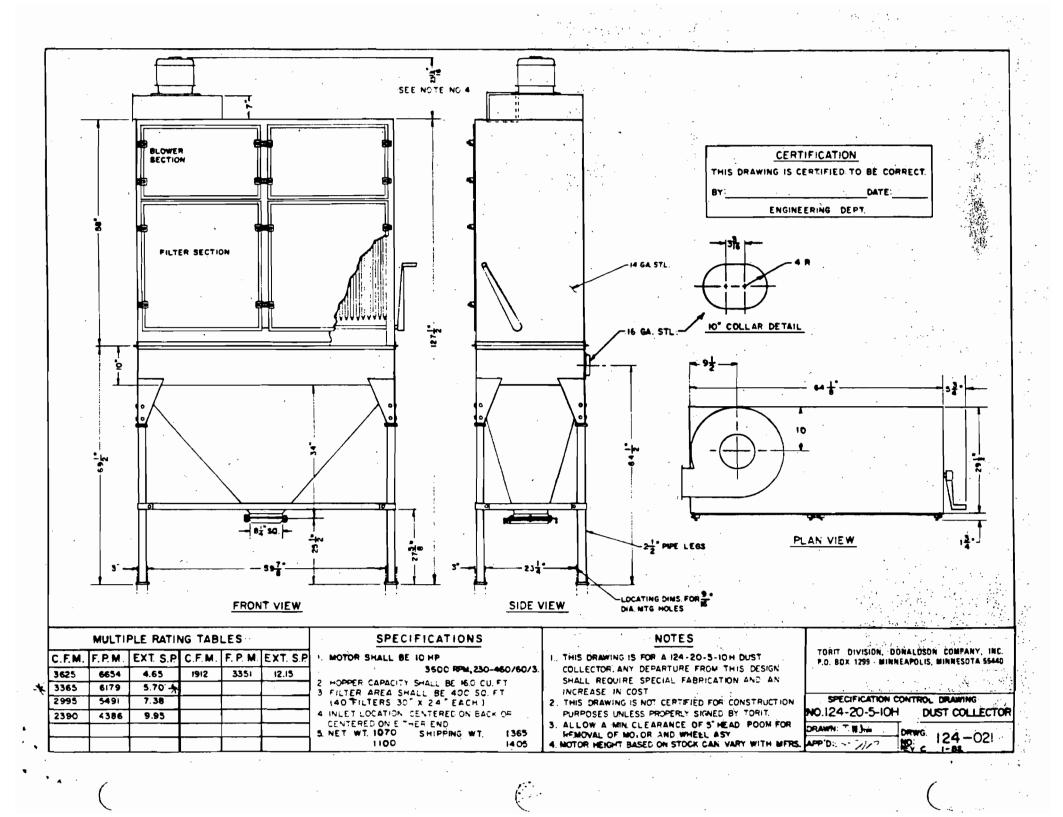
TYPE 1 68-TC - 4" SERIES (NON-POPPETED) FITTINGS







en E



ansfer agent: fungistat plant disease control: tant for polyesters.

OOH. crystals; b.p. sublimes; in hot water alcohol.

O (C. H.)-CHCN. e powder, m.p. 73very soluble in alcohol. cetic acid; synthesis of

e. See diphenadione.

aniline) (C. H.)-NH. sh crystals. Soluble in ahol, and ether; insolup. 52.85°C: b.p. 302°C: on temp. 1173° F: com-

rmula weights of aniline in autoclave. The prodochloric acid to remove residue is distilled. ce and fused. -lined paper bags; fiber

skin. Tolerance, 10 mg

and accelerators; solid dves: pharmaceuticals: preservation of apples: nalytical chemistry.

adamsite: phenarsazine H)C.H.

stals. Sublimes readily. 410°C (dec); insoluble xylene carbon tetra-

envlamine with arsenic

halation and ingestion:

od treating. Poison label. Not ac-Rail) Tear gas label.

48-250°C: insoluble inpluble in toluene.

ength shifter in solution

LC, HANHCAH,

soluble in water; slightly iatic hydrocarbons. Sen-

d fuming sulfuric acid.

See benzidine. d nitrites.

diphenylbromoarsine (C, H,), AsBr. Properties: White crystals, M.p. 54-56°C. Derivation: (a) Hydrobromic acid and diphenylarsenious 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.

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

1.3-diphenyl-2-buten-1-one. See dypnone.

diphenylcarbazide (C.H.NHNH)-CO.

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.

dinhenyl carbonate (C, H,O),CO

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.

diphenylchlorograine (C, H,). AsCl.

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 CO: 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. (And Poison label. Not accepted on passenger planes.

diphenyldecyl phosphite (C6H5O); POC10H21. Properties: Nearly water-white liquid; sp. gr. P. .. (25/15.5°C); m.p. 18°C; refractive index 1.3160 25/D). Combustible.

Uses: Chemical intermediate: stabilizer for polyect and polyolefin resins.

diphenyldichlorosilane (C, H,)2SiCl2.

Properties: Colorless liquid; b.p. 305°C; f.t sp. gr. 1.19 (20°C); refractive index (n 25/D) flash point (COC) 288°F. Readily hydroly moisture, with liberation of hydrochloric Combustible.

Derivation: (a) Reaction of powdered silled chlorobenzene in the presence of copper por catalyst; (b) reaction of phenylmagnesium with silicon tetrachloride.

Grade: Technical.

Hazard: Highly toxic; strong irritant to tissue Use: Intermediate for silicone lubricants.

with wallon, Reaction of didodecyldichlorosilane with nhenyl lithium Hazard: Probably toxic

Use: High-temperature lubricant.

dinhenyldimide. See azobenzene

dinhenyldimethoxysilane (C.H.)-Si(OCH.); Properties: Liquid; sp. gr. 1.080 (25°C); b.p. 191°C (53 mm); refractive index 1.54(14 (25°C). Soluble in acetone, benzene, methyl alcohol; combustible, Hazard: Probably toxic

Uses: Treatment of powders, glass, paper, and fabrics,

dinhenvleneimine. See carbazole

alpha-diphenylenemethane. See fluorene

diphenylene oxide (dibenzofuran) CoH,O (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.H.) CHCH

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 acetaldchyde on benzene in presence of concentrated sulfuric acid. Uses: Solvent for nitrocellulose; organic synthesis,

sym-diphenylethane (bibenzyl; dibenzyl; 1.2-diphenylethane) C. H. CH. CH. C. H.

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 henzyl chloride on ben-Almagnesium chloride. Use Organic synthesis

diphenyl ether. See diphenyl oxide.

diphenylethylene. See stilbene.

N.N-diphenylethylenediamine (ethyl diphenyldiamine) C.H.NHCH.CH.NHC.H.

Properties: Cream-colored solid; sp. gr. 1.14; softening point 54°C; insoluble in water; soluble in acctone, ethylene dichloride, benzene, and gasoline. Use: Antioxidant in rubber compounding.

diphenylglycolic acid. See benzilic acid.

diphenylguanidine (DPG: melaniline) HN:C(NHC, Hs)2. Properties: White powder: bitter taste; slight odor: sp. gt. 1.13; m.p. 147°C; decomposes above 170°C; soluble in ethyl alcohol, carbon tetrachloride, chlotoform, hot benzene and toluene; slightly soluble in

Derivation: Treatment of aniline with cyanogen chlo-

·™C.C#CH:CHCCH:CHC.Hc Use: Wavelength shifter in solution scintillation

diphenyl isophthalate (DPIP) C'H'OOCC'H'COOC'H'

Properties: White solid: m.p. 138-139°C. Combus-Use: Manufacture of polybenzimidazoles, high tem-

perature-resistant polymers.

diphenylketone. See benzophenone.

diphenylmethane (benzylbenzene) (C, H,)2CH, 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-

Derivation: Condensation of benzyl chloride and benzene in presence of aluminum chloride. Hazard: Probably irritant and narcotic in high con-

Uses: Organic synthesis; dyes; perfumery.

diphenylmethane-4,4'-dlisocyanate (MDI; methylene-diparaphenylene isocyanate; methylenebis(phenyl iso-, cyanate) CH (C. H. NCO).

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 phospene

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(C.H.)

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 reguations: (Air) Corrosive label. (Rail) Corrosive material, n.o.s., White label.

diphenylmethylchlorosilane (C,H,)2(CH,)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 C10 H6(NHC6H5); 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) OOC, Hs: CC, Hs 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., corebinding resins); corrosion inhibitor; propellant. Shipping regulations: (Rail) Flammable liquids, n.o.s., Red label. (Air) Flammable Liquid label.

triethylborane (triethylborine; boron triethyl). (C₂H₃), B.

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) C1H3(COOC2H3)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)1H.

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 ib (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
CH₃COOCH₂CH₂OCH₂CH₂OCH₃CH₂OOCCH₃.
Properties: Colorless liquid. Sp. gr. (25°C) 1.112; re-

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) C₁H₁₃COO(CH₂CH₂O)₃OCC₁H₁₅.

Properties: Clear liquid; sp. gr. 0.973 (20°C); acidity 0.3% max. (caprylic); moisture 0.5% 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 CoH10COO(C2H4O)3OCC0H10.

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)
C₃H₁₁OCOCH₂(CH₂OCH₂)₂CH₂OCOC₃H₁₁.

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)
C₂H₁₃OCOCH₂(CH₂OCH₂)₂CH₂OCOC₂H₁₃.
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. Com-

bustible. Low toxicity. Use: Plasticizer.

triethylene glycol dihydroabletaté C₁₀H₃₁COO(C₂H₄O)₃OCC₁₀H₃₁.

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 dlmethyl ether CH₃(OCH₂CH₂)₃OCH₃.

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 C₈H₁,COO(C₂H₄O)₃OCC₈H₁,.

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 C₂H₃CO(OCH₂CH₃),OOCC₂H₃. 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 butoxytriglycol.

triethylenemelamine (tretamine; TEM; 2,4,6-tris(1-azi-ridinyl)-s-triazine) 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.

triethylenephosphoramide (tepa; tris-(1-aziridinyl)phosphine oxide; APO) (NCH₂CH₂), PO.

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-

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 trist 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 NH₂(C,H₂NH)₂C₂H₂NH₂.

Properties: Moderately viscous vellowish 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-ethylbexyl)phosphate [C₄H₀CH(C₂H₃)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 (C₃H₁,O₃), P. 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 C₆H₃(COOC₆H₁₁)₃.

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

Superior numbers refer to Manufacturers of Trade Mark

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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.

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Uses: Plasticizer for vinyl resins; adhesives.

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triethylene glycol di(2-ethylhexoate) C-H₁₅OCOCH₂(CH₂OCH₂),CH₂OCOC₇H₁₅. 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 C10H11COO(C2H4O)3OCC10H31 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 CH₁(OCH₂CH₂)₃OCH₃. 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.

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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.

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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-

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(l-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 eve damage.

Uses: Detergents and softening agents; synthesis of dyestuffs, pharmaceuticals and rubber accelerators. Shipping regulations: (Air) Corrosive label.

tri(2-ethylhexyl)phosphate [C₄H₉CH(C₂H₅)CH₂],PO₄ 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 (C₄H₁₇O)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 C₄H₁(COOC₄H₁₁)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.

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

triethyl orthoformate (triethoxymethane) CH(OC, H₃)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) (C₂H₅)₃PO₄. 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); w1/gal 8.90 lb (68°F). Com-

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 (C2H5)3PO3. 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) (C2H4O),PS.

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 (C2H4OCOCH2)2CHCOOC2H5. 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.

A Ch. flash N AV Biu lb. _ macible with and boron

Core-

Services, no se

Sec. 3.

1-1 msk Ignites malation: and oxi-

Action organest fuel and whitermedi-

work house,

Soo Hydrolyzes in ve oded crys-رة 10.864 (20 مارية عام 10.86 1. wi sat 7,20 lb.

knock agent. Lisk Moderately

mable Inquid, n.o.s., clane (Air) Legal

 $_{\alpha}(H_{1})(X)(Y)$ J. Bitter taste; b.p. Sp. gt. 1 136 (25° C); pour point -50°F; solubility in oil 0.8 (CCC), combustible.

acid w tank cars.

nitrocellulose and emovers; agglutinant; 1 over 0.25%).

Catalyst used in Combustible.

,(r),H. pic, practically odor-່ງທີ່(), b.p. 287.4°C; un (20°C); flash point 20°C), treezing point (20°C). Autoignition immiscible with benabustable, low toxicity. Avgen, as a by-product

I dřums; tank cars. i vinyl, polyester and tion of natural gas; action solvent ("Udex"

ier intermediate; solparts and precision

thyl)trimethylammo-ϽͰϳϧ·ϹͼͰͱϬ·ͺ on); drums.

hloroethylene (q.v.).

obaltic oxide. tricyclamol chloride

oluble in alcohol; in-) (48°C); b.p. 234°C

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

C.H.),BO, sp. gr. 1.065 (25°C); x 1.5480 (24°C); flash in all proportions in

hydrolyzes on contact

hate; TCP) f isomers. odorless liquid. Starefractive index 1.556); wt/gal 9.7 lb; crvs-Miscible with all the s, also with vegetable point 437°F; autoigni-

iosphorus oxychloride. : tank cars.

ingestion and skin abhighly toxic; its toler-

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

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

for plastics and resins.

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der; faint characteristic odor; soluble in alcohol and

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)(1)CH3. Properties: Colorles 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 (tridecylic acid; tridecoic acid) CH₁(CH₂)₁₁COOH. A saturated fatty acid usually prepared synthetically.

Properties: Coloriess 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 C15 hydrocarbons. Grade: Technical Containers: 55-gal drums; tank cars.

Uses: Esters for synthetic lubricants; detergents; antifoam 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.

tridecylic acid. See n-tridecanoic acid.

tri(decyl) orthoformate CH(OC10 H21)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 (C10 H21O)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 $C_0H_{11}C(C_0H_2)(OH)CH_2CH_2N(C_2H_3)_2 \cdot C_2H_3CI$. (3-

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

[(CH₁)₂NCH₂]₃C₄H₂OH. Properties: Liquid; refractive index 1.5181. Combus-

Hazard: May be toxic.

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

tri(dimethylphenyl)phosphite (trixylenyl phosphate) [(CH₁); C, H₂O]; PO.

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." 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 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 CIC₃N₃[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)

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 (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

triethanolamine methanearsonate CH₃As(O)[ONH(C₂H₄OH)₃]_{2.7}

applications, especially shampoos.

Hazard: Highly toxic by ingestion.

Use: Herbicide. Shinning regulations: (Rail Air) Arsenical compounds. triethanolamine oleate. See trihydroxyethylamine ole-

triethanolamine stearate. See trihydroxyethylamine

triethanolamine titanate. See titanium chelate,

1.1.3-triethoxyhexane

CH(OC:Ha):CH:CH(OC:Ha)CaHa. 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) (CH₂O)(C₂H₂O)CHCH₂CH(OC₂H₃)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

triethylaconitate C2H4OOCCHC(COOC2H4)CH2COOC2H4.

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). $(C_2H_1)_1AI$.

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 (C2H3)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

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.

APPENDIX V

Volatulity

TABLE 15.3

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

No.	Name			ν° C.		20° C.	Sc	olubitity,	% at 26	*C.	in	tility of Pasticisor Air, cm./hr.
		Formula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	Viscosity, cp. at 26	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	875° F. (191° C.)
		Simple P	kuticise	тө	·							•
1 2 3 4 5 6 7 8 9 10 II	Acetin o- and p-Toluenesulfonamides (mixture) Discetin N-Ethyl-p-toluenesulfonamide N-Ethyl-o- and p-toluenesulfonamides (mixture) Triscetin Dimethyl phthalate o-Nitrobiphenyl Diethyl phthalate Triethyl citrate	C _b H ₁₀ O ₄ C ₇ H ₉ O ₂ NS C ₇ H ₁₇ O ₆ C ₉ H ₁₃ O ₂ NS C ₉ H ₁₃ O ₂ NS C ₉ H ₁₄ O ₆ C ₁₀ H ₁₀ O ₉ C ₁₂ H ₉ O ₂ N C ₁₂ H ₂₀ O ₇ C ₁₂ H ₂₂ O ₇ C ₁₂ H ₂₇ O ₄	134 171 176 199 199 218 194 199 222 276	1.190 at 25° C. 1.353 solid 1.178 1.171 at 65° C. 1.190 at 25° C. 1.191 1.203 at 25° C. 1.118 1.136 at 25° C. 0.988 at 26° C. 0.988 at 30° C.	1. 445 at 25° C. 1. 448 at 25° C. 1. 522 at 65° C. 1. 540 at 25° C. 1. 613 at 25° C. 1. 613 at 25° C. 1. 442 at 26° C. 1. 442 at 26° C. 1. 442 at 26° C. 1. 442 at 26° C. 1. 443 at 26° C.	66.8 at 25° C f.p. 106 35.7 at 22° C, f.p. 59.0 426 at 23° C, 15.1 at 23° C, 16.3 at 25° C, 12.6	0.15	0.7	0.5	0.34	0.55 >4.0 3.40	101 >1000 > 500

		C	mperieo	a of Pr	roperti	es of Vi	inylite I Equiv	Resin V' alesco a	YN₩-6	Specim	ens Pla	sticised	10	. ,
No.	Name	Pasticier p.h.r. at 23° C., 75% Elongation, 1000 p.a.i.	4	ngation.	eter A	23°C., p.k.i.		erature, °C.	% l	Com.	10 d	istrac- on, laya, d-mil	2 days. 60° C. Humidity	Supplier **
		Placticiner p. 75% Elongat	Tennile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer at 23° C.	Stiffness at 2	1. °C	Brittle Temperature,	20 da ya 20°C.	24 bours, 70° C.	Water	් ස	Spew, 2 days 100% Humid	
	,		Simple	Plastic	iters,									
1	Aostin		T	Ī·	1							.	_	Kee
2	o- and p-Toluenesulfonamides (mixture)													Mon
8	Discetia		ŀ											Kee
4,	N-Ethyl-p-toluenesulfonamide	-:.	. `	ļ.,	li		İ							Mon
5	N-Ethyl-s and p-toluenesulfonamides (mixture)			-	-		Ì						—	Mon
6	Trinoctin	.]	1											Kes, TE
7	Dimethyl phthalate								٠					Mon, TE
8	o-Nitrobiphenyl												-	Mon
9	Diethyl phthalate		:	:			l	٠.						Mon, TR, USI
10	Triethyl citrate	3, 1											<u> </u>	Pf .
11	Dimethyl sebacate													Her
		1 "		1	1		1							

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

" ()ash (----) indicates no data availa

PLASTICIZERS AND PLASTICIZATION

TABLE 15.3 (Continued)

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

No.	Name			چ د.		20° C.	So	lubility,	% at 20°	c.	Free Pi	itity of lasticisor Air, om./br.
	Nemb	Formula	Molecular Weight	Specific Gravity, 20°	Befractive Index	Visconity, cp. at 20	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	876 F. (191 C.)
		Simple Plasticis	ers (Cos	utinued)					•			-
12	Di-n-butyl tartrate	C12H22O4	262	1.094	1.444	67.7	0.5	0.7	0.3	2.4	1.50	143
18	Tri-n-butyl phosphate		266	0.978	at 25° C.	at 25° C.	<0.07	6.0	U.S	# #		501
	• • •	C12H27O4P		}	1.424 nt 20° C.	8.4 at 25° C.			' '		>4	
16	Monomethyl phthalate ethyl glycolate ester	C14H14O0	266	1.227	1.504 at 25° C.		0.1	1.3	0.5	0.1	0.65	e u
15	N-Cyclohexyl-p-toluenosulfonamide	C14H19O2NS	253	1.125 solid		f.p. 87 C.					0.05	26
. 16	Monoethyi phthaiate ethyi giyeolate ester	C14H16O6	280	1.186	1.498 at 25° C.	68.0 at 25° C.	<0.08	1.38	1.60	0. 18	0.50	93
17	Di(methyi Cmilosouva) phthalate	C14H18O6	282	1.169	1.501	53	0.85	3.2	0.50	0. 10	0.16	62
18	Triethyl acetylcitrate	C14H22O8	318	1.13	at 25° C. 1.439							> 525
19	Disobutyl adipate	C14H26O4	258	0.957	at 23° C. 1.428	70						ĺ
20	Tributyrin	C16H26O8	302	1.035	at 25° C.	at 25° C. 8.9						
21	Methyl Callocolva laurate	Слензоов	258	0.894 at 25° C.	at 25° C. 1.440 at 25° C.	at 25° C. 7.8 at 25° C.						

M = Miscible.

		Con	nperiece	of Pr	perti	ne of Vi	equive	Resia V'	4-WAY	Specim	ens Plas	rticised	اد 	
ía.	Name Name	og, 1000 p.e.i.	4 .	Elongstion, C.	eter A	23° C. pai.		brakure, °C.	% I 4-mil	Loss, Film	% E. 10 d 23° C. Fri	ĸa.	2 days, 60° C., Humidity	Supplier **
		Plasticiser p.hr. 76% Elengation	Tensile Strength pai at 23° C.	Ultimate Elor	Shore Durometer A at 23° C.	Buffrom at 22	T. C.	Brittle Temperature,	10 days, 60° C.	24 bours, 70° C.	Water	3	Bpew, 2 days, 100% Humid	
		Simple	Plastici	tere (C	ontin	ed)		-			٠.			
2	Di-n-butyi tartrate	1.		,	•	,								Kee
3	Tri-n-butyl phosphate		İ			· .						^		C8
	Monomethyl phthalate ethyl glycolate enter	55.4¢			`				17.5		6.1	5.9		Moo
4	· · · · · · · · · · · · · · · · · · ·													Mon
	N-Cyclobexyl-p-toluenesulfonamide				. .	۰		l	l .			1	L	
5						***	:	! 			-		_	Moo
16	N-Cyclobexyl-p-toluenesulfonsmide	46 *							6.6	٠.	4.6	8.7		OVALE Woo
5	N-Cyclohexyl-p-toluenesulfonamide Monosthyl phthelate sthyl giyoolate ester	46 *			2 W 25				6.6		4.6	8.7	None	
15	N-Cyclohexyl-p-toluenesulfonamide Monosthyl phthalate sthyl glycolate ester Di(methyl Czigosogwa) phthalate	46 *			100000000000000000000000000000000000000	·表情等等于 · · ·					4.6	8.7	None	OALTE Pf Cub
14 15 16 17 18 19 20	N-Cyclobexyl-p-tolucnesulfonsmide Monosthyl phthelate ethyl glycolate ester Di(methyl Chilosoliva) phthelate Triethyl acstylcitrate	46 *			1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Salar Barrell					4.6	8.7	None	OANTE

^{9 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.

Por key to conciler abbreviations are page 902.

Dash (---) indicates no data available.

PLASTICIZERS AND PLASTICIZATION

TABLE 15.3 (Continued)

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

No.	N				30° C.		j.	Su Su	olubility,	% at 20	· C.	Free Pi	ility of lasticises Air, . con./hr.
No.	Name		Formula	Molecular Weight	Specific Gravity, 28	Befractive ladex	Viecosity, cp. at 20°	lo Water	Water in	In Mineral Oil	Mineral Oil in	217° P. (100° C.)	876" P. (191" C.)
			Simple Plasticis	ero (Cos	utinued)								
22	Di-n-butyl phthalate		C16H22O4	278	1.048	1.493 at 20° C.	20.3	Insol	0.46	34	30.0	0.98	148
23	Di(CELLOSOLYE) phthalate		C14H22O6	310	1.120	1.491	1.p. 31 °C.	0.2	2.0	2	0.64	0.15	30
24	n-Butyl laurate		C16H32O3	256	0.857	at 25° C.	31° C.						
25	Methyl Crilosolve myristate		C17H34O3	236	at 25° C. 0.895 at 25° C.								
26	Di(butyl Carror) formal		C17H36O6	336	0.97								
27	Triphenyl phosphate	•	CisHisO4P	326	1.268 at 60° C.	1.552 at 60° C.	f.p. 49.9° C.					0.02	22
28	Bis(dimethylbensyl) ether		C13H23O	286	1.008	1.553 at 25° C.	29.0						
29	n-Butyl cyclohexyl phthalate		C ₁₅ H ₂₄ O ₄	304	1.078	1.508 at 20° C.	117.0				.		75
30	Mono-n-butyl phthalate n-butyl glycolate ester		C16H24O6	336	1.103	1. 490 at 25° C.	64.9	Insol.	0.37	4.65	2.70	0.08	81
31	Tri-n-butyl citrate		C15H32O7	360	1.045	1. 443 at 25° C.		Insol	<0.2	10.0	14.5	0.16	4
32	Tristhylane glycol di(2-ethylbutyrate)		C16H16O6	348	0.995	1. 440 at 20° C.	11.5	0.02	1.0	0.2	24.0	0.88	

		Con	mperiso	of Pr	operti	ea of V	inylite I Equiv	Resin V alence	YN W- 5	Specim	eas Plac	sticised	to, ,	÷.
No.	Name	r. at 23° C., 00, 1000 p.a.i.	4	estion,	ther A	23° C., p.a.i.	··	Temperature, °C.	% I 4-mil	oss, Film	10 d 23° C.	xurac- on, lays, . 4-mil	66 °C.	Bupplier =
		Placticiser p.hr. at 23° C., 75% Blongation, 1000 p.a.i.	Tennile Strength p.s.i. at 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer /	Beid new at 23		Brittle Tompe	10 days. 80° C.	M hours. 70°C.	Water	8	Brew, 2 days, 60° C., 100% Hamidity	*
	<u> </u>	Simple	Plastici	г ат е (С	estin	ed)								
22	Di-a-butyl phthalate	41.4 *				. •			17.5		2.7	6.8	None	Bar, CB, Har, Mon Nev. Pitt, TR, USI, DA
23	Di(CELLOSOLVE) phthalate							- 1					_	OA AH
24	n-Butyl laurate													AR
25 26	Methyl Callosolva myristate Di(butyl Calastrol) formal	55.30							10.1		15.6	14.5	None	Thei
27	Triphenyl phosphate								••••					Dow, Moo, Pitt
28	Bis(dimethylbensyl) ether											, .	. _	OA
29	n-Butyl cyclohexyl phthalate	58.8					+8	~20	0,5		0.6	8.8	None	Ber
30	Mono-n-butyl phthalate n-butyl glycolate ester	58.2°	3000	309					8.8		17	4.8	None	Mon
31	Tri-a-butyl citrate	55.1	2900	300			4.	-23	8.1		8.1	14.6	None	Pf
82	Triethylene glycol di(2-ethylbutyrate)		,				. . ·				,		·	ccco

¹⁰⁰⁰ p.a.i. teusile modulus at 75% elongation Dash (——) indicates no data available. 2 100% elongation, 1000 p.a.i., constant rate of For key to supplier abbreviations see page 90

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TABLE 15.3 (Continued)

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

No.				20° C.		, c	So	lubility,	% at 20°	° C.	Free Pt	ility of lesticise Air, . cm./hr
No.	Name	Pormula	Molecular Weight	Specific Gravity, 20	Refractive Index	Viecosity, cp. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	375° F. (191° C.)
		Simple Plastici	sera (Co	ntinued)								
33	Butyl CRILOSOLVE laurate	C18H26O2	300	0.882 at 25° C.	1. 439 at 25° C.	6.5 at 25° C.						
34	Di-n-hexyl adipate	C14H34O4	314	(av.) 0.933	1.439	7.8						
35	Di(1,3-dimethylbutyl) adipate	C18H34O4	314	at 25° C. 0.926	at 25° C. 1.433	5.6 ca.						
38	Di-n-butyl sebacate	C18H34O4	314	0.936	at 25° C. 1.440	at 38° C. 9.8	lasol.	1.6	H	м		6
37	Di(butyl Callosolva) adipate	C18H34O4	346	0.997	at 25° C. 1.442	12.5						5
38	s-Butyl myristate	C18H26O2	284	0.861	# 32. C							
39	Tri(butyl CELLOSOLVE) phosphate	C15H29O7P	398	at 25° C. 1.023	1.434	12.2	0.08	6.8	Almost	nuiscible		
юί	Creayl diphenyl phosphate	C19H17O4P	340	1.208	at 25° C. 1.563	39.8						
,	n-Butyl bensyl phthalate	C ₁₉ H ₂₀ O ₄	312	at 25° C. 1.116	at 20° C. 1.538 at 20° C.	39.8						2
12	Bis(dimethylbensyl) carbonate	C19H22O2	298	at 25° C. 1.063	at 20° C. 1.547 at 20° C.	at 25° C. 156						3

M = Miscible

	Coa	o per isot	of Pr	op or ti	ee of Vi	nylite I Equiv	Rasia V Menos *	YNW-5	Specius	ns Plan	nicised	lo	
. Name	Plarticiser p.b.r. at 23° C., 76% Elongation, 1000 p.a.i.	q	agation,	eter A	23° C., p.s.i.	·.	crature, °C.	% I 4-mil	oes, Film	10 d 23° C.	ktrac- on, lays, . 4-mil	2 days, 60° C., Humidity	Supplier **
	Plasticiser p. 75% Elongat	Tensile Strength, p.s.i. at 23° C.	Ultimate Elongation, % at 23 ° C.	Shore Durometer at 23° C.	Suffness at 2	T. °C.	Brittle Temperature,	10 daya, 60° C.	24 hours, 70° C.	Water	Ŋ.	Srew, 2 days 100% Humid	
	Simple	Plastici	era (C	ontinu	ed)								
Butyl Cellosolve laurate	Ī			,									Kos
Di-n-hexyl adipate						,							Kes
Di(1,3-dimethylbutyl) adipate							1	:					Har .
Di-n-butyl sebacate	37.8°			.		ľ	ł	7.2		0.4	16.5	None	Har, Pitt, RH
1•		1									18.0	None	Kos, OA
Di(butyl Chilosolva) adipate	41.4	l .											
	41.46				∵,							_	AH
Di(butyl Chilosolva) adipate s-Butyl myristate	43.7 4				· .			1.1	_	14.2	13.5	None	OA
Di(butyl Cullocolve) adipate n-Butyl myristate		3000	292	64	475	•	-15	1.1	6.1	14.2	13.5 14.8	None None None	

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

6 100% elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed. 25.

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TABLE 15.3 (Continued)

PROPERTIES OF COMMERCIALLY AVAILABLE PLASTICISERS ARRANGED IN ORDER OF ASCENDING NUMBER OF CARBON ATOMS

Name	Pormula	Molecular Weight	Specific Gravity, 20° C.	Refractive Index	ity, cp. st 20° C.	b		35	.д		
		·		2	Viscosity,	In Water	Weter in	In Mineral Oil	Mineral Oil	212° F. (100° C.)	876° F. (101° C.)
	Simple Plasticizer	re (Com	tinued)								
sthyl pentachlorostearste sthyl ricinoleste sthyl Callosouva palmitate	С ₁₉ Н ₂₂ О ₂ Сl ₈ С ₁₉ Н ₂₆ О ₂ С ₁₂ Н ₂₆ О ₂	471 312 314	1.204 0.925 at 25° C. 0.891	1.497 at 20° C. 1.462 at 25° C.	975.0						7
sthyl hydroxystearate phenyl phthalate syclohexyl phthalate	C ₁₉ H ₁₈ O ₂ C ₂₉ H ₁₄ O ₄ C ₂₀ H ₂₈ O ₄	814 818 330	1. 29 solid 1. 23	1, 579 s4 74° C.	1.p. 48° C. 1.p. 69° C.					0.006	10
Sthylhexyl diphenyl phosphate	C ₂₀ H ₂₀ O ₄ C ₂₀ H ₂₀ O ₄	363 834 334	1.002 1.007 1.016	2.512 st 20° C. 1.487 st 20° C. 1.488 st 25° C.	29.5 56	Insol.	0. 94 <0.05	M	M	6.20	44 43 44 1
pi oj	henyi phthalate vdohezyi phthalate shyihezyi diphenyi phosphate s-hezyi phthalate	henyi phthalate CzoH14O4 CzoH2zO4 chinkayi phthalate CzoH2zO4 chinkayi phthalate CzoH2zO4 CzoH2zO4 CzoH2zO4 CzoH2zO4 CzoH2zO4 CzoH2zO4	henyl phthalate CzoH14O4 318 CzoH24O4 330 CzoH24O4 330 CzoH24O4 330 CzoH24O4 363 CzoH24O4 363 CzoH24O4 364 CzoH24O4 334 CzoH24O4 234 CzoH24O4 234 CzoH24O4 234 Czo	hyl hydroxystearate benyl phthalate czoHi4O4 318 1.28 solid rciohexyl phthalate czoHi2O4 330 1.33 hylhexyl diphenyl phosphate chexyl phthalate czoHi2O4 331 1.003 1.007 ctoHylbutyl) phthalate czoHi2O4 334 1.007	hyl hydroxystearate condition c	hyl hydroxystearate C19H1#O2 314	hyl hydroxystearate benyl phthalate	hyl hydroxystearate C1pHagO2	hyl hydroxystearate C_{10}H_{16}O_{0} 814	hyl hydroxystearate C19H1002	hyl hydroxystearate C_{1pHagO_2} 314

M - Miscible

	A STATE OF THE STA	·						·					4.	P.
•		Coa	mperison	a of Pr	operti	ana of ∀i	nylite E Equiv	Resia V	YN W -4	Specim	sca Pla	sticised	to.	
No.	Name	h.r. st 25° C. ion, 1000 p.s.i.	ą.	aration,	estér A	25°C, pai.		ersture, C.	% l 4-mil	oes, Pilm	10 d	strae- on, laya, 4-mil		Bupplier =
		Pasticies phr. s 75% Ecognica, 1	Tensile Strength, pai at 23° C.	Oltimate Bong	Shore Durom at 23° C.	P Buiffness at 3	F. C.	Brittle Temp	10 days.	N bours	Water	8	Spee, 2 days, 100% Humidis	
		Simple	Plastici	sera (C	ontina	ed)					. ;	•.		
43	Methyl pentachlorostesrate	69 4	2900	388	54	600	- 6	-19	0.2		0.2	. 17. 1	None	Hack
4	Methyl ricinoleste Methyl Carrocorve palmitate			,					YS _C .					Bak AH
44	Methyl hydroxystearate		<u> </u>					'					_	Bak
47	Diphenyl phthalate												_	Moa
4	Disyslohexyl phthalate	64.5	3006	290		t gr	+ 5	.~ S	. 120		0.	4.5	None	Bar, OA
19	2-Ethylhexyl diplomyl phosphate:	60	2580	260	64	540	-, 3	-28	1.6		0.2	14.8	None	Moa
50	Di-a-hexyl phthalate	57	2720	293	63	570	- 1	-81	. 3.8	11.6	0	18.5	None	0000, TE
51	Di(3-othylbutyl) phihahtis	45.3.*	:- <u>/:</u> :		٠,			18	4.4		0.3	3.4	Nome	TR. v
. 53	Di(1,8-dimethylbutyl) phthainte		:		,		A Second						_	Her

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

Mixture of 1 part plasticises to 2 parts Flaxos, plasticises DOP.
 Mixture of 0.9 part plasticises to 1 part Flaxos, plasticises DOP.

^{100%} elongation, 1000 p.a.i., constant rate of loading of 810 p.a.i. per minute. See Reed.

Table 15.3 (Continued)

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

				, C.		ü	80	lubility,	% at 20°	· C.		lity of nationer Air, em./hr,
No.	Name	Formula	Molecular Weight	Bpecific Gravity, 20°	Befractive Index	Viscouity, ep. at 20°	In Weter	Weter in	In Mineral Oil	Mineral Off in	812° P. (100° C.)	875 P. (191° C.)
		Simple Plasticis	ers (Cos	tin ud)								
-53 54	Di(butyl Callosolve) phthalate Di(Carretol) phthalate	C ₂₀ H ₂₀ O ₆ C ₂₀ H ₂₀ O ₈	366 398	1.063 1.150	1.483 at 25° C. 1.492	42.0 82.0	Insol.	0.63 2.80	2.80 0.09	6.50 0.20	0.08	24 18
55 56	Tri-n-butyl acetyloitrate n-Butyl palmitate	C ₂₀ H ₂₄ O ₈ C ₂₀ H ₄₉ O ₂	402 313	1.046 at 25° C. 0.865	at 25° C. 1.441 at 25° C.							. 41
57	Tricrenyl phosphate	C21H21O4P	368	at 25° C. 1.165	1.557 at 20° C.	120.0	Insol.	0.87	8.8	3 .0	0.01	7.
58 59	n-Butyl benzyl sebacate Methyl scetylricinoleste	C21H22O4 C21H22O4	348 855	1.004	1.479 at 25° C. 1.456	8.6 ca. a4.88° C. 33.1						
60	Methyl Callosolus oleste	C21H40O2	. 341	0.903	at 20° C. 1.453	10.8					0,13	29
61	Di(2-ethylbutyi) aselate	C21H40O4	857	0.930	at 25° C. 1.442 at 25° C.	11.4						44
82	Propylene glycol monoricinoleste	C11H40O4	857	0.950 at 25° C.	~							
63	Methyl Callosolve stearate	C ₈₁ H ₄₃ O ₃	848	0.877 at 25° C.	1.443 at 25° C.	8.9 a4 25° C.						30

	•	Con	operiso	a of Pr	opert	ies of Vi	inytite I Equiv	Resin V alence	YNW-6	Specim	ens Pho	sticized	to .	
No.	Name.	Lr. at 23° C.,	4	Stion,	ster A	°C, pai		rature, °C.	% 1 4-mi	oes, Film	l '- tir	strac- on, lays, . 4-mil	60°C.	Supplier.
		Particies phr. 178% Bongston,	Tennie Strength pal na 23° C.	Ohmste Flongstion, % at 13° C.	Shore Durometer A at 23° C.	Buffness at 23°	7° 7°	Brittle Temper	30 th 30 th	24 km	Water	8	Sper. 2 days, 60° C., 160% Hamidity	
_		Simple	Plantici	eero (C	ontin	ud)				·			: .	
53	Di(butyl CELLOSOLVE) phthalate	47.8*			[·				2,1		4.8	11.4	None	OA.
54	Di(Campron) phthalate	54.4*							2.0		10.1	9.8	None	OA.
55	Tri-a-butyl acetyleitrate					-			٠. :			. <i>'</i>	—	Pf
56	n-Butyl palmitate		1							.· .	::		<u> </u>	AH.
57	Tricresyl phosphate	75.	2880	274	58	880	+ 5	- 3	0.1	:	0.7	6.1	None	Col, Mon, OA, Pitt
58	n-Butyi bensyi sebacate								1					Har
59	Methyl scetylricinoleste	47.8%				r.			8.8		9.5	11.0	: 	Bak
80	Methyl Czzzosorva oleste	80.5 W			· .				8.6	1.	8.0	10.3	_	AH, Kes, CA
61	Di(3-ethyibutyi) ascinte	44.5	2900	260	ľ	:		-40	8.3		1.5	14.0	Nome	B
63	Propylens giyent monoricinolests										٠		<u> </u>	Belt
62	Mothyl CRILOSOLVE steerate		٠.						. ,		V ** *		 _	AH, Kes

TABLE 15.3 (Continued)

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

				ပ န		ರ	So	lubility,	% at 20°	C.	Volati Free Pl in mg_/sq.	Air.
No.	Name	Formula	Molecular Weight	Specific Gravity, 20°	Refractive Lodes	Viscosity, op. st 20*	In Water	Water in	In Mineral Oil	Mineral Oil in	212° P. (100° C.)	876° F (191° C.)
		Simple Plasticis	era (Con	tinued)							ı .	
64	n-Butyl ricinolests	C12H42O8	855	· .								
65	Di(1,3-dimethylbutyi) sebacate	C12H43O4	· 871	0.911	1.439 at 25° C.	9.3 ca.						
66	Di(2-ethylhexyl) adipate	C22H42O4	871	0.937	1.447 at 20° C.	at 88° C. 18.7	Insol.	0.12				41
67	Discoetyl adipate	C22H42O4	871	0.928	1.448 at 25° C.	17.7					_	88
68	Dicapryl adipate	C22H42O4	371	0.915	1. 440 at 25° C.	8.3 cs. a4.38° C.						
69	Diethylene glycol dipelargonate	C22H42O4	387	0.963	1.447 at 20° C.	19 ~						
70	Di(butyl Casaconouva) sebscate	C22H42O4	408	0.970	2.30 C.					١.	0.04	
71	Triethylene giyeol di(2-ethyl hexanoste)	C22H42O4	408	0.968	1.444 at 20° C.	16.1	Insol	0.4	M	M	0.18	50 ·
72	Triethylene glycol dicaprylata	C22H42O0	406 (av.)	0.978	- W W						.,	
78	Di(butyl Carreron) adipate	Ç23H43O3	486	1.09 at 16° C.		15 at 22° C.		,				
74	s-Butyl stearste	C22H44O2	341	0.858	1.444 at 20° C.	7.9 at 25° C.				·		59

M - Missible

		Con	nparison	of Pr	operti	es of Vi	oylite F Equiv	Resin V alence 4	YNW-5	Specia	ens Pia	Micised	LO .	
No.	Namo state de la companya della companya della companya de la companya della comp	on, 1000 p.s.i.	ą	e tide.	otor A	'C, pai		rature, °C.	% I 4-mil	oss, Film	10 d 22° C.	zirso- on, lays, 4-mil	60° C.	Supplier =
		Placticiser p.h.r. at 28° C., 75% Elongation, 1000 p.e.i.	Tennile Strength, pai at 23° C.	Ultimate Bongation % at 23° C.	Shore Durometer . at 23° C.	Buffness at 28°	1. °C	Brittle Temps	10 days.	24 hours, 70° C.	Water	8	Spew, 2 days, 60° C., 100% Humidity b	
		Simple	Plastici	sers (C	ontin	ed)								
64	a-Butyl ricinoleate									•			_	Doe
65	Di(1,3-dimethylbutyi) sebacate								•					Har
66	Di(3-ethylhexyl) adipate	51	2730	318	67	600	~15	- 53	3.5	12.2	2.2	21.0	None	CCCC, Gdr, MWi,
68	Dissoctyl adipate Disspryl adipate												_	Ken, OA, Har Ken, MWi, OA, Pitt, RCA, Har, Cub Har
. 69	Diethylene glycol dipelargonate	51.3	2800	330			-15	48	6.9		6.0	22.0	Yes	16m
70	Di(butyl Calacactes) sebacate	٠.								7.5				Dee
71	Triethylene glycol di(2-sthylhexanoste)	69 ·	2689	800	56	800	-17	-50	4.7	12.4	2.0	22.5	None	occc
73	Triothylene glycol dicapsylate		Ì				,			· .			-	BCA, Drew
78	Di(butyi Casarrot) adipate	·	:							in.				Thi
74	n-Butyl steerate	1	Ì						. 2	V	1,			CB, AH, Har, Kea, OA

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 misute

Destr. (---) indicates no data available.

DESCRIPTION OF TABLES AND FIGURES

PLASTICIZERS AND PLASTICIZATION

TABLE 15.3 (Continued)

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

				ۍ د.		ڻ ن	Bo	lubility,	% at 20°	c .	ia.	ility of asticiaer Air, em./hr.
No.	Name	Formula	Mobeular Weight	, Specific Gravity, 20°	Refractive lades	Viscouity, op. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	312° F. (100° C.)	876° P. (191° C.)
		Simple Plasticis	ers (Cos	tinued)								
75	a-Butyl cleate	C22H47O2	339	0.864 at 25° C.	1.449 at 25° C.	7.7				,		
76	Tetrahydrofurfuryl oleate	C22H42O3	366	0.925	1.466 at 20° C.	at 25° C. 17.8						
π	Methyl CELLOSOLYB acetylricipoleste	C22H42O5	399	0.955	1.456 at 20° C.	42.1	Insol.	0.6	M	M		12
78	e-Xenyl diphenyl phosphate	C34H19O4B	402	I. 288	ca. 1.586	80-85 a4 60° C.	InsoL	0.35	2.23	0.8	0.005	20
79	Dibensyl sebacate	C24H30O4	382	1.055 at 30° C.	1.520 at 30° C.	13.0 ca. at 38° C.						
80	Di(2-ethylhexyl) phthalate	C24H28O4	390	0.980	1.486 at 20° C.	81.0	Insol.	0.20	M	M	0.02	24
81	Diisootiyl phthalate	C24H28O4	390	0.987	1.486 at 20° C.	83.0						19
82	Dicapryl phthalate	C24H38O4	390	0.966	1.479 at 20° C.	66.7						23.
88	Di(2-ethylhexyl) tetrahydrophthalete	C24H42O4	395	0.969	1.466 at 20° C.	42.0	Insol.	0,26				30 ·
84	Tetra-n-butyi thiodisuocinate	C24H42O3B	491	1.054	1. 463 at 20° C.	53.6	Insol	0.22				6
85	Di(2-ethylhexyl) hexabydrophthalate	C24H44O4	897	0.958	1.461 at 20° C.	85.9						87
86	a-Butyl acetylricinoleste	C24H44O4	397	0.929	1.456 at 25° C.	35.2						80

		Ca	mperiso	of Pr	operti	es of Vi	nylite I Equiv	Resin V sistace *	YNW-4	Вресіл	os Pla	ticised	to	
No.	Name	or, at 23° C.	4.	gation,	oter A	28°C, pat		watere, *C.	% i	Loss, I Film	10 c	strac- on, lays, 4-mil	,	Supplier **
		Plasticiser p.h.r. 75% Elongation,	Tensile Strength pai at 23°C.	Ultimate Elongation, % at 23° C.	Shore Durameter A	Bifform at 20	2.14	Brittle Temperature,	80 00 100 100 100 100 100 100 100 100 100	22 bount	Weter	පි	Spew, 2 days, 60° (
		Simple	Plastici	ters (C	ontina	ud)								
75	s-Butyl oleste													AH, Kes
76	Tetrahydrofurfuryl oleate	59 4	2590	297	67	650	- 8	-34	0.8		0.8	17.2	None	AH, Em, Pitt
77	Methyl Childeolym acetylricinoleste	50.8¢					,		2.4	ĺ	1.3	14.5	None	Bak, Dee
.78	o-Xenyl diphenyl phosphate													Dow
79	Dibenayi sebacate	45.84		•					0.7		0.8	11.8	None	Har
80	Di(2-ethylhexyl) phthalate	61	2360	253	67	706	- 1	-81	0.7	5.1	0	15.5	None	Bar, CCCC, Har, Mon, MWi, OA,
81	Discoutyl phthalate	56	2700	300		٠,	- 1	-27					None	Pitt, TE, DA, Odr Bar, Cab, Har, MWi,
82	Dicapryi phthalate	64·	2500	270		÷. 、	~ 5	-35	0.6		0.	23.8	None	Bar, Cub, Har, MWi, OA, Pitt, RCA RH, Har
83	Di(2-othylhexyl) tetrahydrophthalata	66	2590	320	58	560	~13	-37	4.1	11.5	6.8	21.8	None	cocc
84	Tetra-a-butyl thiodisuccinate	64 .	3200	318	62	760	~ 2	~22	o.	2.5	0.8	15.2	None	ococ
85	Di(2-ethylhexyl) hazahydrophthalata	64	2580	300	68	750.	-10	-34		9.7	0	25.4	None	0000
		55.4*												

DESCRIPTION OF TABLES AND FIGURES

Properties of Commercially Available Plasticizers Arranged in Ohder of Ascending Number of Carbon Atoms 8

				U S		ü	Boh	ıbili ty , 9	% at 20°	C.	Volatil Free Pla in / mg./sq.	Lir.
o.	Name	41	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscouity, op. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	876° F. (191° C.)
_		Simple I	Plasticisers (C	ontinued)								
,	Butyl Callosolve cleate	CsaH		at 25° C.	1.454 at 25° C.	5.5 at 25° C.						 -
,	Dinonyl adipate	C ₂₄ H ₄	6 O₄ 898	at 25° C.	1.445 at 25° C.	18.5 at 25° C.			ļ	1		'
1	n-Octyl n-decyl adipate (mixture)	C14H	4O ₄ 398	1	1.450 at 20° C.	15.4		1.4		,		44
ŀ	Polyethylene glycol di(2-ethylheranoate)	C24H.	407 447	0.989	1.447 at 20° C.	25.1	Insol.	1.4			1	26
١	Butyl Carrosorva stearate	C ₂₄ H.	30 ₃ 38	1	1.446 at 25° C.	13.3	١	١.,	ı,	<u> </u>] .	36
١	Tri(2-ethylhexyl) phosphate	C24H	10,P 43	0.926	1.448 at 20° C.	14.1	Insol.	1.4	, A	_		20
	Di(2-ethylhexyl) azelnie	C ₁₆ H.	18O4 413	0.918	1.450 at 20° C.	20						~
	n-Octyl n-decyl phthalate (mixture)	CasH.	12O4 41		1.484 at 20° C.	72]					Ι '
	Isooctyl a-octyl a-decyl phthalate (mixture)	C ₂₆ H		8 0.978	1.485 at 20° C.	42.9		l				۱ ا
	Di(2-ethylhexyl) schacate	CaeH		0.912	1.451 at 20° C	19.9					'	١ .
ı	Dimocctyl sebacate	C ₂₄ H	10O4 42	7 0.917	1.447 at 25° C	20.4 25° C.	1					
6 7						13.0 cs.						

1 1 1 1 1 1 1 1 1 1	•	Cor	nparisos	of Pro	operti	ee of Vi	nylite F Equiv	Resin VI alence ⁴	YNW-6	Specim	na Plas	ticised	to	
Simple Plasticiaers (Continued) Simp		or, 1000 pai	£ .	gation,		ぴ			% [. 4-mil	oss, Film	tic	ND:	eo C. ity	Supplier ^m
Butyl Cultocolve cleate Dinonyl adipate Di		Planticiser p.h. 75% Elongati	Tensile Streng p.a.i. at 23° C	Ultimate Eloc	Shore Durons	4	f. "C.	Brittle Tamp	10 days. 60° C.	24 hours, 70* C.	Weter	78	Spew. 2 days. 100% Humid	
Dinonyl adipate n-Outyl n-decyl adipate (mixture) 53 2550 295 78 1050 - 7 -54 5.5 0.3 23.5 None Here, OA Polyethylens glycol di(2-ethylhexanoate) Butyl Cellosolve stearste Tri(2-ethylhexyl) phosphate 59 2540 300 66 980 -16 -67 8.2 6.8 0.6 24.8 None CCCC Di(2-ethylhexyl) seelate 50 2540 318 64 640 - 8 -38 0.2 0.3 21.8 None Here Fitt	Simple	Plastic	sero (C	ontina	red)									
n-Octyl n-decyl adipate (mixture) 53 2550 295 78 1050 -7 -54 5.5 0.3 23.5 None Here, OA Polyethylens glycol di(2-ethylhexanoste) 62 2690 308 70 750 -12 -44 2.8 8.8 3.0 19.4 None CCCC Butyl Crizzosolve stearste 59 2640 300 65 990 -16 -67 8.2 6.8 0.6 24.8 None CCCC Di(2-ethylhexyl) seelste 57 2590 334 66 750 -17 -56 0.5 0.4 22.5 None Em. A -Octyl n-decyl phthalate (mixture) 63 2400 318 64 640 -8 -88 0.2 0.3 21.8 None Here B Isodetyl n-decyl phthalate (mixture) 67 2720 334 66 470 -12 -39 3.9 0 22.4 None Here B Di(2-ethylhexyl) selects 60 2420 298 67 1700 -19 -56 0.08 4.6 0.8 30.2 None Dec, Har, MW	Butyl Chilosolve oleste													Kee :
Polyethylens glycol di(2-ethylhexanoate) 62 2690 308 70 750 -12 -44 2.8 8.8 8.0 19.4 None CCCC	Dinonyl adipate				•									Pitt
Butyl Cellosolve stearste 59 2640 300 66 980 -16 -67 8.2 6.8 0.6 24.8 None CCCC	n-Octyl n-decyl adipate (mixture)	63 .	2550	295	78	1050	- 7	54		5.5	0.8	23.5	None	Here, OA
Tri(3-ethylhexyl) phosphate: 59	Polyethylene glycol di(2-ethylhexanoste)	62	2690	308	70	750	-12	-44	2.8	8.8	8.0	19.4	None	OCCC
Di(2-ethylhexyl) secisie 57 2590 334 56 750 -17 -56 0.5 0.4 22.5 None Em.	Butyl CELLOSOLVE stearate]											 —	AH, Kee, OA
s-Octyl s-decyl phthalate (mixture) 62 2540 318 54 540 - 8 -38 0.2 0.8 21.8 None Here Isooctyl s-octyl s-decyl phthalate (mixture) 57 2720 334 56 470 -12 -39 8.9 0 23.4 None Here Di(2-cthythexyl) sehecate 60 2420 298 57 1700 -19 -55 0.08 4.6 0.8 30.3 None Dea, Har, MW	Tri(2-ethylhexyl) phosphate	549	2640	300	66	980	-16	-57	8.2	6.8	0.6	24.8	None	0000
IsoSctyl s-octyl s-decyl phthelate (mixture) 87 2720 834 66 470 -12 -89 8.9 6 23.4 None Here	Di(2-ethylhexyl) assists	57	2590	334	66	750	-17	-56	0.5		0.4	22.5	None	B _m
Di(2-ethythexyt) sehecate 00 2420 293 67 1700 -19 -56 0.08 4.6 0.8 30.2 Nome Des, Har, MW	s-Octyl n-decyl phthalate (mixture)	62	2540	318	64	640	- 8	-88	0.2		0.8	21.8	None	Here
	Isooctyl a-octyl a-decyl phthelate (mixture)	67	2720	334	66	470	-13	-89		8.9	0	23.4	None	Here
Diisotetyt sebacata	Di(2-ethythexyl) sebecate	90	2420	298 .	67	1700	-19	-56	0.08	4.6	0.8	30.2	Nome	Dee, Har, MWi, RI
	Diisooctyl sebacate												_	Har, Pitt, MWi

	,	· ·			
PROPERTIES OF COMMERCIALLY AVAI			A	Maragana on C	ADDION ATOMS
C A7747	TABLE DIAGRICIVEDS	ARRANGED IN URDER OF	ASCENDING	NUMBER OF C.	ARBON ATOMS
Properties OF COMMERCIALLY AVAI	INDER I PUSITOIEN	TEMPORAL TO COMPANY			

				່ວ		ບູ	Sol	ubility, 9	% at 20°	C.	Volatil Free Pla in A mg./sq.	eticiser ir.
ía.	Name	Formula	Molecular Weight	Specific Gravity, 20*	Refractive Index	Viscosity, cp. at 20° C.	In Water	Water in	In Mineral Oil	Mineral Oil in	212° P. (100° C.)	875°F. (191°C.)
_		Simple Plasticis	ers (Con	tinued)								
11.	2-Ethylhexanoic acid diester of N.N-bis(2-hydroxyethyl)-2-ethylhexan- amide Tri(p-tert-butylphenyl) phosphate Glyceryl triricinoleate Glyceryl tri(acetylricinoleate)	C ₂₆ H ₈₂ O ₆ N C ₂₀ H ₈₉ O ₄ P C ₆₇ H ₁₀₄ O ₉ C ₆₂ H ₁₁₀ O ₁₃	484 495 933 1060	0.956 0.959 at 25° C. 0.964	1.458 at 20° C. 1.477 at 25° C. 1.469 at 25° C.	139. 2 f.p. 95-99. 5 °C.	Insol.	0.5		,		12 0. 0.
	-	Resinous F	laticise	78								
03 04 05 06 07	AROCLOR 1243, chlorinated biphenyl FLEXOL plasticizer B-2H, polyester G.E. 2557, polyester G.E. 2559, polyester PARACELL C, nitrile rubber PARAPLEX G-25, polyester		ca.	1.378- 1.388 at 25° C. 1.055 1.028 1.060 0.984 1.06	1.627- 1.629 st 20° C. 1.469 st 20° C. 1.459 st 20° C. 1.492 st 20° C.	ca. 20,000 306 4278	Insol.	1.0	м -	, M	2.50	146 0. 3 1

ľ		Co	mperiso	a of Pr	operti	ee of Vi	aylite I Equiv	Rosia V	ANM-1	Specim	ons Plac	ticised :	lo.	
No.	Nams	phr. st 23° C., pation, 1000 pai	म्	Elongston, C.	eter A	23°C, pai		erstern, "C.	% 1 4-mi	oss, Film	10 d	trac- io, iaya, i-mil	eo c.	Supplier **
		Plasticiser p. 75% Elongst	Tensile Strength	in a sa	Shore Durometer et 23° C.	Stiffners at 2	7. ℃	Brittle Temperature,	10 days C. C.	24 bours,	Weter	. i3	Spew, 2 days, 60° (100% Humidity b	•
``		Simpl	Plaetic	isera (Contin	wd) ·					¢			
» »	2-Ethylhexanoic acid diester of N.N-bis(2-hydroxyethyl)-2-ethylhe anamide Tri(p-tert-butylphenyl) phosphate	5- 64	2730	827	63	650	- 8	-28	0.4	. 1.7	1.1	11.0	None	CCCC Dow
122	Glyceryl triricinoleste Glyceryl tri(acetylricinoleste)	68.64.4							-1,4		2.5	9.5	None	Bak Bak
		Re	rinous F	laticis	ere			•						
œ	AROCLOB 1242, chlorinated biphenyl	. 56.2 h		1		,			4.5		0.6	0.6	None	Mon
94	PLEXOL plasticiner B-2H, polyester	90	3080	360	57	440	+3	-10	0	0	0	2.4	None	cccc
06 06	O.E. 2557, polyester O.E. 2559, polyester	72 .	2730 2110	810 215	63 73	580 740	- 3	-18 - 6	0.9	3.5 2.3	3.1 2.7	17.0 7.1	Very slight Very	GE GE
07	PARACREL C, nitrile rubber PARAPLEK G-25, polyester	106	2500 2500	410 310	50	480	± ;	-54 -25	0.5	•	0.2	0.6	alight None None	Nau RH

ant rate of loading of 15,000 p.s.i. per minute.

Name	Resinous Plastici	Molecular Weight	Specific Gravity, 20°	Befractive Indax	Vaccerty, ep. at 20°	In Water	ä	ra Oil	di ii		
	Resinous Plastici	sera (Co			.>	4	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	875° F. (191° C.)
APLEX (1-40 tuplyonter			ntinued)								
APLEX G-50, polyester	•	6000 68. 2200 ca. 850	1.15 1.084 1.043	1.472 at 20° C. 1.470 at 20° C. 1.462 at 20° C.	290,000 8000 481						0. (1 2
	Hydrocarbon-T	ype Plas	ticitere		•						
w 276-V2, polymolocular product derived from a-methylstyrene 40, partially hydrogenated mixture of isomeric terphenyls FLAX L, hydrocarbon resin made by reaction of formaldehyde + di- bethylnaphthalenes FLAX C 10° (PHO), a hydrindyl phenol derivative		240	1.021 1.007 1.01 at 15.6° C. 1.075 to 1.10	1.582 at 20° C. 1.570 at 25° C. 1.597 at 25° C.	112.8 f.p. -1° C. 23,500 at 25° C.						118
riziol.			1.03 to 1.08 at 15.6° C. 0.955		65 to 110 at 25° C. 151.6						
,	ntac 10° (PHO), a hydrindyl phenol derivative	nlac 10° (PHO), a hydrindyl phenol derivative mol	DILLC 10° (PHO), a hydrindyl phenol derivative DIOL APLEX BN-1, alkylated aromatic hydrocarbon mixture	15.6° C. 240 (av.) MLAC 10° (PHO), a hydrindyl phenol derivative 240 (av.) 240 (av.) 240 1.075 1.03 10.1.08 241 1.03 241 1.03 242 1.03 243 245 245 245 245 246 247 247 247 247 247 247 247 247 247 247	15.6° C 1.597 1.597 1.598 1.56° C 1.075 1.597 1.598 1.56° C 1.075 1.598 1.56° C 1.081 1.08 1.08 1.56° C 1.081 1.08 1.08 1.56° C 1.58°	15.6° C 1.075 1.597 23,500 at 25° C 1.075 at 25° C 15.6° C. 1.59° C. 1.59° C. 1.59° C. 1.59° C. 1.00° C.	15.6° C. 1.597 1.597 1.597 1.596 1.596 1.596 1.08 1.596 1.597 1.596 1.	15.6° C 1.675 1.597 23,500 24,500 25.6° C 1.075 25.6° C 1.075 25.6° C 25.6	18.6° C. 1.05 1.597 23.500 24.00 24.00 24.00 25° C. 24.00	15.8° C. 1.597 23,500 23,500 24,000	

	and the service of th	Con	n paris o	a of Pr	operti	ies of V	inylite l Equiv	Resin V	YNW-	Specim	ees Ph	ticised	to.		
ła.	Name	hr. at 23° C.,	4	Elongation, C.	eter A	28° C., pai.		ersture, *C.	% 1 4-mi	Lons, I Pilm	10 c 23° C	xtrac- on, lays, , 4-mil	dayr. 60° C., lumidity b	Supplier **	
		Plantiniser p.h.r. 76% Elongstion,	Tensile Strength, p.s.l. at 23° C.	Ultimate Elo	Shore Durometer	Briffness at 22	2	Brittle Tempe	30 days.	24 bount	Water	8	Spew, 2 days, 100% Humid		
		Rosinous	Plastic	isere ((Contin	med) ·	· .					•			
Q	PARAPLEX G-40, polyester	67 ^j	2780	310	69	760	1	-15	0.8	-	1.0	8.2	Slight	RH .	
0	PARAPLEX G-50, polyester	75:	2630	320	62	680	0	-16	0.8		1.8	6.5	None	RH	
1	Plasfolein 9720, polyester	69	2850	312	66	900	- 7	-23	0,8	1.8	1.9	14	None	Bm.	
		H ydroc	arbon-T	ype Pl	anticis	at	J			<u> </u>			<u> </u>	<u> </u>	-
2	Dow 276-V2, polymolecular product derived from a-methylatyrene	74 5	2420	815	62	650	- 5	-24	1.4	1.	0.6	22.6	None	Dow	•
8	His 40, partially hydrogenated mixture of isomeric terphenyls.	63.4	2890	298	66	940	- 1	-23	8.4		0.5	15.5	None	Mon	
4	Kamplex L. hydrocarbon resin made by reaction of formaldshyde + dimethylmaphthalence				.,								<u> </u>	Ken	٠,
5	Navellac 10° (PHO), a hydrindyl phenol derivative	56.8 ª A	ļ.					† . 	13.3	. :- :	0.1	0.1	None	Nev	
8	Navinol				ψ.								None	Nev	
7	Panapan BN-1, alkylated aromatic hydrocarbon mixture	53 4	2800	310	66	920	3	-21	6.1	'	1.8	16.8	None	Pan	
8	Sovalom C, alkylated aromatic hydrocarbon mixture	61.4	8100	280	.		+4	-16	11.5	.	1.7	9.0	None	8V	

^{* 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 minuta.

^{6 100%} elongation, 1000 p.s.i., constant rate of loading of 810 p.s.i. per minute. See Reed.26

Table 15.3 (Continued)

Properties of Commercially Available Plasticizers Arranged in Order of Ascending Number of Carbon Atoms

		ń,		° C.		° C.	80	lu bility,	% at 20	° C.	l in	ility of assicisor Air, om./ar.	
No.	Name	Formula	Molecular Weight	Specific Gravity, 20°	Refractive Index	Viscosity, cp. at 20°	In Water	Water in	In Mineral Oil	Mineral Oil in	212° F. (100° C.)	876° F. (161° C.)	PLASTICIZERS
		M iscellaneous	Plastic	isere			•						AND
119 120 121 122 123 124 125 126 127 128	ARNESI. TOD, mixture of olsic, linolaic, and cyclic nitriles n-Butyl acetyl polyricinoleate, PG18 Chlorinated paraffin 40 Flexol plasticiser B-400, polypropylene glycol mono-n-butyl ether Glyceryl polyricinoleate, No. 15 oil Harseax 500, monomeric ester type Hercoriax 600, a pentaerythritol ester Ohorax Q-10, octyl fatty-phthalic soid esters Parafrax G-60, high-molecular-weight ester Plasticiser DP-520, chlorinated aliphatic product Polyethylene glycol dilaurate		538 (av.)	0.910 0.913 at 25° C. 1.164 0.995 1.031 at 25° C. 0.932 1.005 0.952 0.994 1.125 at 2.0.97 at 25° C.	1.460 at 25° C, 1.505 at 23° C. 1.449 at 20° C. 1.483 at 20° C. 1.478 at 20° C. 1.478 at 20° C. 1.478 at 20° C.	30.2 3100 at 25° C. 153 17.0 48.3 41 340 to 260 60.4 os. at 54° C.	0.1	5. 5				0.9	PLASTICIZATION
130 131	SANTICIER 180 OTAFIEE KA, monomeric ester type			1.071	1.506 at 20° C.	234	,						:

·		Con	n perison	of Pro	operti	es of Vi	aylite B	Regio V	NW-5	Specime	ens Plas	ticised (ا ما	
No.	Name	on, 1000 pai	ų,	ogstion,	eter A	23° C., pai.	edma	arture, C.	% I	one, Film	10 0	tirso- on, lays, , 4-mil	2 days, 60° C Humidity	Supplier **
		Particine: p.h.r. at 23° 75% Elongstion, 1000	Tensile Burength, pai. st 23° C.	Ultimate Elongation, % at 23° C.	Shore Durometer at 23° C.	Stiffness st 22	£.°C.	Brittle Temperature,	10 days.	24 bours, 70° C.	Weter	70	Spew, 2 days, 100% Humid	
		Miss	ellansou	e Plas	ticien									
119	Annent TOD, mixture of oleis, limbers, and syclic nitriles	52.8	2900	290				-44	11.7		0.7	14.8	None	Arm
120	n-Butyl acetyl polyricinoleate, PG16	1	Tuccustre 	tible s	55.3	p.b.r.		•	1					8ak
121	Chlorinated paraffin 40	68 * .	2820	335	57	900	- 5	28	1.0		0	20	None	Halo, Dia, Here, Hook
122	FLEXOL plasticizer B-400, polypropylene glycol mono-s-butyl ether													cccc
123	Chyceryl polyricinoleste, No. 15 oil												_	Bak
124	HARFLEE 500, monomeric ester type	- 68	2400	319	67	980	-10	-81	1,1		0.4	23.0	None	Har
125	Hamcortax 600, a pentacrythritol ester	58	2916	327	66	900	- 3	-28	٥.	1.1		12.8	None	Hero
. 126	Onorex Q-10, octyl fatty-phthalic acid enters	· · ·					İ			. •				OA.
127	PARAPLEX G-60, high-molecular-weight ester	69	2800	317	67	880	- 2	-21	0.2	ļ	0.6	11.1	None	RH
128	Plasticiser DP-520, chlorinated aliphatic product:													Drow
129	Polyethylene glycol dilaurate	-			1							,		Cly
130	SARYERER 180	78	2670	308	67	480	- 3	-17		2.7	1.8	12.2	None	Men
, l31	STAPLEX KA, monomeric ester type												-	Dos

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 ≈ 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.

Michael a: Ware

MAW:bjw Attach.

ESTIMATED

OVERALL COMPLIANCE SCHEDULE

DAVID M COMPANY VOC CONTROL PROJECT

PHASE

Data Accumulation

Spreader Enclosure - (1) For Temperature Evaluation

Process Modifications and Development

Methods Comparison/Analysis - Thermal Incineration Vs. Recovery

Final Specification Development

Final Specification/Quotations and Analysis

Appropriations Request Development and Approval

Oven Enclosure and Exhaust System Installation

Solvent Recovery Delivery and Installation

Operational Shakedown, Debugging and Compliance Testing

January 1, 1984

January 15, 1984

April 1, 1984

May 15, 1984

July 1, 1984

September 1, 1984

December 1, 1984

March 1, 1985

August 1, 1985

December 1, 1985



SUTCLIFFE SPEAKMAN INC

SUITE 200, HEAVER PLAZA, 1301 YORK ROAD, LUTHERVILLE, MARYLAND 21093

TELEPHONE: 301-337-2800 TELEX: 908020 240189 ARYLAI

BUPY

YOUR REF

OUR REF: . E 0379

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, Fiorida

SOLVENT RECOVERY PLANT

ACTIVATED CARBON ADSORPTION UNIT

AUTOMATICALLY CONTROLLED

BY

GAS ANALYSER

PRELIMINARY PROPOSAL



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 indlecate overloading from excessive steam flow or shortage or failure of the cooling water supply.

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

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.

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 stalnless 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 26¹¹ 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

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" dlameter 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 solld 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-hazard-ous 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 piant items.

David M. Company

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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 pslg.

17. DRAWINGS AND INSTRUCTIONS

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

Foundation drawing Fiowsheet 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 2.4.0189

YOUR REF:

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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

DE SPATCH

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

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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.

SUTCL IFFE SPEAKMAN INCORPORATED

William C. Moses Vice President

E. & O. E.

David M. Company

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.

STEAM1190 lb/	hr, dry	saturated	quality
at 40 ps	ig.	,	

COMPRESSED AIR............20 scfh.

SPACE

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

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 plling 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 Unasion Inhibitor is? Chromoly

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 work-manlike 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.

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.

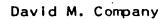
10. INSURANCE

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.



Proposal No. S 7212

BUDGETARY PRICE

(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. 5 7212.



SOLVENT RECOVERY

WITH

ACTIVE CARBON

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

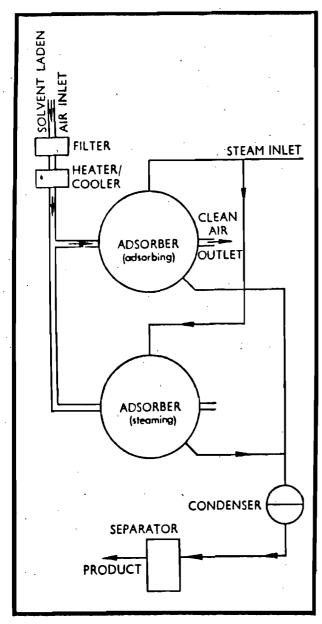
A general improvement in working and safety conditions often results from the effective collection system required to achieve high overall efficiency.

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.

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 LOSS VALUE LBS/HOUR/1000 SCFM PER ANNUM X1000 CONC. % V/V STERLING WEIGHT Jo 40. 60 70. 100 0.7 125-0.5 150 175 0.4 200 225 0.5% CON. METHYL ETHYL KETONE -M.W. 72-1 LOSS 56-5 LBS/HOUR/1000 SCFM OF AIR VALUE PER ANNUM (OF 8000 HOURS) AT 10 PENCE/LB IS £18.800 VALUE PER ANNUM (OF 8000 HOURS) AT 11-5 CENTS/LB IS \$52,000

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	В		A	В
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)	118.18	2.4	Methylens Chlorida	84.93	Non-flam
Butyl Alcohol (Normel)	74.12	1.45	Nonane	128.25	0.74
Butyl Alcohel (Iso)	74.12	1.68	Octone	114.23	0.95
Cerbon Bisulphide	78.13	1.0	Pentane (Normal)	72,15	1.3
Carbon Tetrachloride	153.84	Non-flem.	Pentana (Iso)	72.09	1.3
Chloroform	119.39	Non-flam.	Perchlorethylene	165.85	Non-flam
Cyclobexane	84.18	1.35	Propyl Acetate (Normal)	102.3	2.0
Decane (Normal)	142.28	0.7	Propyl Acetate (Iso)	102.3	2.0
Dichloroethylane	97	9.7	Propyl Alcohol (Normal)	60.09	2.15
Ether (Diethyl)	74.12	1.65	Propyl Alcohol (Iso)	60.03	2.02
Ethyl Acetate	88.10	2.25	Tatrachlorethane	187.66	Non-flam
Ethyl Alcohol	46.87	3.3 -	Tatrahydrofuran	72.10	1.84
Ethylene Dickloride	96.97	8.2	Talvene	92.13	1.3
Forferal	96.08	2.1	Trichlorethylene	131.4	Non-flam
Heptane (Normal)	100.2	1.0	Xylene	106.16	1.0
Herane (Normal)	86.17	1.25			

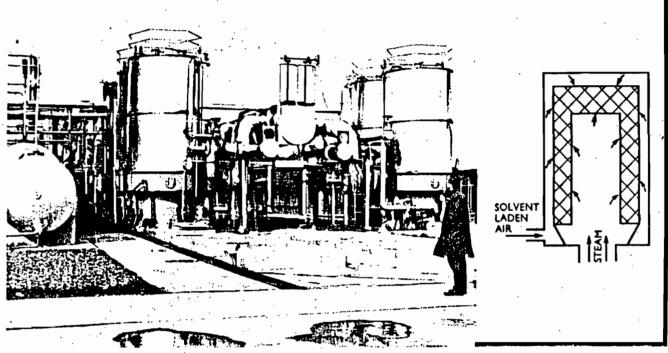
A Mol. Wt.

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.

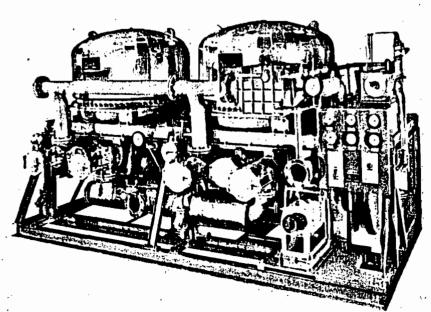
STATIC BED PLANT

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.

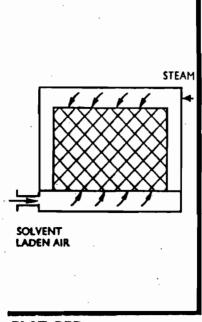


Adsorption section of a large recovery complex at an Acetate Fibre Factory in Belgium. The plant is arranged for future extension.

ANNULAR BED



Packaged type fully automatic plant of cylindrical flat had design



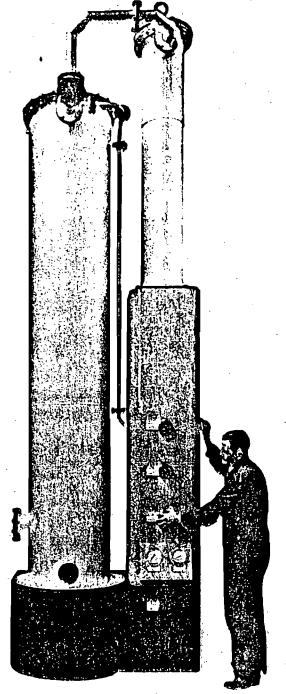
FIAT RED

LIQUID PHASE ADSORPTION

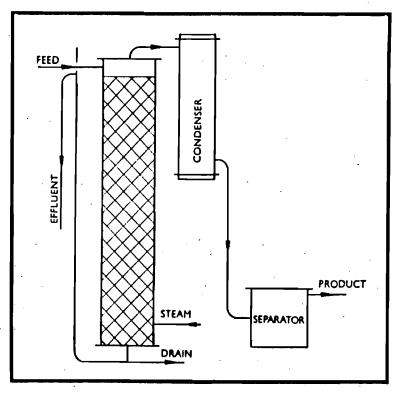
This system uses active carbon for the adsorption of small amounts of certain water-miscible solvents from solutions with water. The weak solvent/water liquor is passed through an active carbon bed and, when the carbon has adsorbed the designed charge of solvent, the solvent is distilled off by our patented 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.

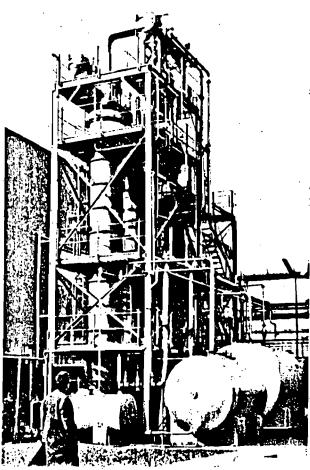
A particular feature of this system is the extreme simplicity of operation and control which can be manual or automatic.



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

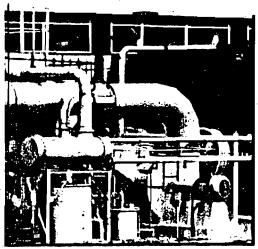
SUITE 200, HEAVER PLAZA, 1301 YORK RO, LUTHERVILLE, MARYLANO 21093 TELEPHONE: 301-337-2800

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	Concen- tration (%V/V)	*Initial adsorp- tion (%W/W)	§Cyclic adsorp- tion (%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	58			
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	B6.7			
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 3, 3, 5-trimethyl cyclohexyl	0.3	35.6	11.9	3.7	153-193			
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 en economical steem consumption.

*Steam ratio refers to the total steam used to desorb the solvent from the edsorbent end is en overall figure inclusive of condensete and moisture condensed on the carbon bed.

SENSOR CONTROLS

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:

(a) Inlet concentration 1000 ppm Outlet concentration 35 ppm

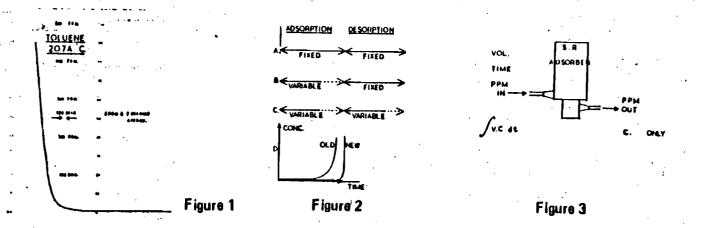
$$\mathbf{n} = \frac{1000 - 35}{1000} \times 100\% = 96.5\%$$

(b) Inlet concentration 2000 ppm Outlet concentration 35 ppm

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

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.



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

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

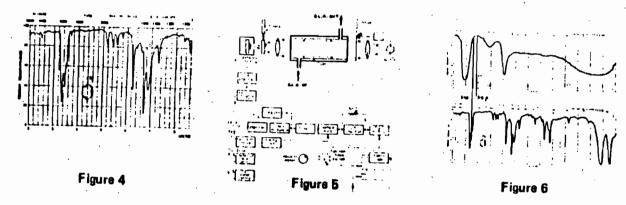
as solvent laden air enters the adsorber we can measure solvent concentration and flow rate, and compute, by integration with respect to time, the total quantity of solvent passed to the adsorber. When a predetermined amount of solvent has been passed a signal is generated which is used to change over 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 inditate 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



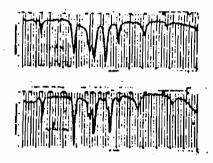
being sensitive to background absorptions caused by fouling of the sample cell windows, broad band absorptions in the sample gas (such as that caused by water vapor), and temperature differences between sample and reference beams. However, most of the disadvantages of the two beam system can be eliminated 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 mlcrons 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





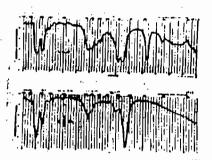


Figure 7

Figure 8

Figure 9

To illustrate just how complex the 1-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 stronly, Figure 7 shows the 1-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 absorbtion 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 absorbtion bands.



Figure 10

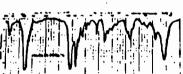


Figure 10 shows the I-R spectra for ethyl acetate and hexane. Here we have a good illustration of the relative weakness of the CH band at 3.5 microns for ethyl acetate and the corresponding strength of the CH band for hexane at the same point in the spectrum.

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.

- (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.



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YOUR REF:

OUR REF:

DATE: 4 August 1983

DESTON FOR EXPANDINGLIETY L

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 gailons 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 sultable hoods for the spreading machines and a collection system when you so require.

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