INVIRONMENTAL PROTECTION ACT 1990, Part I

The Environmental Protection (Prescribed Processes and Substances) Regulations 1991. The Environmental Protection (Applications, Appeals and Registers) Regulations 1991.

See Notes on pages 3 and 4 before completing this form.

APPLICATION FOR AUTHORISATION to carr	y out prescribed proces	S
under section 6 of the Environmental Protection Act 199	a i	

To(1) Huntingdonshire District	Council
1 Name and address of applicant (2) (in the case of a roffice) W R Grace Ltd	registered Company, name, number and registered
Northdale House	
Northcircular Road, London	Tel.No. 081/965/0611
2 Name and address of premises where process is or will	be carried on (not applicable to mobile processes)
See attached report	Tel.No.
3 In the case of mobile plant, name and address of the pri	ncipal place of business
	Tel.No.
4 Address for correspondence relating to the application	
See attached report	
C	
Contact name	Tel.No.
5 List of maps or plans enclosed with the application process is or will be carried on. TITLE	
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See attached report for item 6 - 17 inclusive

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10 Give details of the so process. (use a continuat		unt of current and	or anticipated emission	ons to air from th
11 Give the assessmer continuation sheet if nec	essary)			
	and the second section of the sectio	er e	en e	
12 What monitoring is o	r will be carried out of e	missions to air?		

7 When was the plant first installed?

13 What monitoring will be o	arried out of the environ	mental consequer	ices of emissions	to air?
14 How will you monitor the	techniques described in	the answer to que	estion 9?	
15 State how you will ensur				
1990 will be achieved and ho	ow the condition implied	by section 7(4) of	the Act will be co	omplied with.(6)
				
details. (use a continuation si	heet if necessary)	•		
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17 Give any other addition authority in considering you		ou would like to	be taken into a	ccount by the loca
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If you require any further in Council at the address show	information or assistant in below.	ce in completing	this form, please	e contact your loc

Please complete the final section of this form on page 4 overleaf.

Cheques should be made payable to:
I HEREBY CERTIFY that all the information contained in this application is correct to the best of my knowledge and belief [and that I am authorised to sign on behalf of the Company].
Signature
Official title
Date
Please complete and return this form together with FOUR copies of each of the plans listed in the reply t question 5 and the required fee to:

(8)

I enclose the fee of £

Tel.No.

NOTES

- 1 This is the local authority in whose area the prescribed process will be carried on, or in the case of mobile plant, the local authority in whose area the applicant has his principal place of business.
- 2 Please state the person/Company who is operating or will operate the process, not an agent who may be completing the application on the operator's behalf.
- 3 A list of prescribed processes for local authority control is given in Appendix A, which accompanies this form. Further advice can be obtained if necessary from the local authority.
- 4 A list of prescribed substances for release into the air is given in **Appendix 8**, which accompanies this form. "Harm" includes offence to the senses or harm to property.
- 5 Please list fully all pollution control measures for all stages of the process, from the receipt of raw materials to the despatch of wastes and finished products, including, for example, the height and location of any stacks or vents; the abatement technology; process control and operational data; arrangements for maintenance; the extent of supervision; the relevant qualifications and experience of the workforce; staff training; and contingency plans for breakdowns and emergencies.

All calculations should be shown, particularly for the chimney height(s). Justification for the selection of a particular abatement option should be given.

- 6 Section 7(2) and 7(4) of the Environmental Protection Act 1990 requires every operator of a prescrib process to use the best available techniques not entailing excessive cost for -
 - (i) preventing the release of prescribed substances, or where that is not practicable, for reducing the release of such substances to a minimum and rendering them harmless; and
 - (ii) rendering harmless any other substances which might be released.
- Much of the information contained in the application form will be included in a register which the local authority is required to keep for public examination in accordance with section 20 of the Environmental Protection Act 1990 and the Environmental Protection (Applications, Appeals and Registrars) Regulations 1991. Sections 21 and 22 provide for certain information (affecting national security, or commercial confidentiality) to be excluded from the register. Such information should be clerarly identified in this application form.
- £800 in the case of initial applications.
 £530 in the case of applications for a substantial change.
 £530 in the case of processes transferred from previous HMIP control.
 £100 for small waste oil burners.

Container Solvent Plant

WR Grace Ltd Cromwell Road, St Neots

Application for Authorisation under the Environmental Protection Act 1990

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APPLICATION FOR AUTHORISATION

SECTION 6: ENVIRONMENTAL PROTECTION ACT 1990

1.0 NAME AND ADDRESS OF APPLICANT

W R Grace Limited Northdale House North Circular Road London NW10 7UH

Telephone: 081 965 0611

2.0 NAME AND ADDRESS OF PREMISES WHERE PROCESS IS CARRIED ON

Container Solvent Plant WR Grace Ltd Cromwell Road St Neots Cambridgeshire PE19 2ER

3.0 NOT APPLICABLE

4.0 ADDRESS FOR CORRESPONDENCE

Container Solvent Plant WR Grace Ltd Cromwell Road St Neots

Contact Name: Dr John Measures, Plant Manager

Telephone: 0480 475061

5.0 LIST OF MAPS/PLANS

1. Ordnance Survey Map.

2. Site Plan.

6.0 DESCRIPTION OF PRESCRIBED PROCESS

The Environmental Protection (Prescribed Processes and Substances) Regulations 1991 requires processes involving the manufacture of dyestuffs, printing ink and coating materials to be authorised under the Environmental Protection Act 1990. The criteria for notification to the local authority is whether the process uses 100 tonnes or more of an organic solvent within a 12 month period. The Container Solvent Plant uses in excess of 100 tonnes and hence the application is being submitted as a Part B process for authorisation by the local authority.

The plant produces a dispersion of rubber in a solvent which is used extensively throughout the food industry as a sealant for containers. There are three basic rubbers used styrene butadiene rubber, neoprene rubber and nitrile rubber. The first part of the process involves the production of a rubber master batch which is prepared using a Banbury mixer where the constituents are mixed mechanically to produce a solid rubber mass. This is a coarse mixture which is then transferred to a 2 roll mill located immediately adjacent to the Banbury mixer where further mixing and compounding continues. The resultant compound is then removed from the 2 roll mill in the form of strips or sheets which are then fed to a chipper which cuts the material into small strips. These strips are then fed from the chipper up along the conveyor and then transferred across to the solvent plant from the rubber compounding room. Extraction systems are provided within the rubber compounding room at points 1, 2 and 3 which are shown on the site plan.

The rubber strips are then transferred by conveyor to 1 of 4 mixers shown on the site plan as SW, MM, SSW and EKATO. The rubber is generally dispersed in either hexane which consists of C5-C7 isomers with less than 5% n-hexane, which forms the bulk of the production or a mixture of acetone and propan-1-ol. Other solvents are also used which include isopropanol which is used for the manufacture of liquid fluxes and isopar G which is produced as a nozzle dousing fluid. The mixing process is carried on within closed mixers although they are vented to atmosphere via condensers.

The mixer is constantly agitated by means of a mechanical stirrer and the compound is pumped to bulk storage in one of the five bulk storage tanks shown on the site plan. The compound is then recirculated through a homogeniser to bulk storage. Figure 1 illustrates a mixer in diagrammatic form. Material Safety Data Sheets in respect of the three main products are contained at Appendices 3, 4 and 5.

The solvents namely hexane and acetone are all stored in bulk tanks located below ground which are vented to atmosphere via vent pipes each of which has a flame trap fitted. The location of the storage tanks is also shown on the site plan.

7.0 PLANT INSTALLATION

The plant was first installed in 1972 and since that date there have been a number of modifications, details of which are given below:-

1976

- second five tonne mixer installed (MM).

1986

- 3½ tonne acetone mixer installed (EKATO).

December 1986 - 25,000 litre interceptor pit installed.

March 1989

- Pipe bundle type condensers fitted to all bulk tanks and mixing vessels which are terminated exterior to the building via a flame trap. (No's 4-12, Appendix 2).

In addition increased pump capacity was provided to give better circulation and therefore better heat exchange and more efficient cooling.

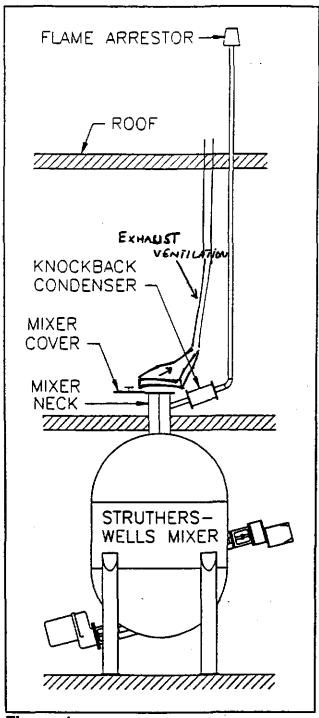


Figure 1 A Struthers Wells mixer,

April 1989

- A new honeycomb was fitted to the cooling tower to provide more effective cooling and therefore reduce potential solvent emissions.

1990

- The small 21/2 tonne Struthers mixing vessel was installed (SSW).

1992

- A 5th bulk storage tank was installed together with a 2 tonne liquid flux kettle (BT5 + LF tank).
- A 600 gallon (2 tonnes) isopropyl alcohol storage tank was also installed. (IPA tank).

August 1991

- The heat exchanger was thoroughly cleaned and descaled which produced a 10°C reduction on the temperature of the mix. This resulted in temperatures on the input side of the homogeniser of between 50-60°C and on the output side of the homogeniser of 30-40°C. The initial boiling point of hexane is 55°C.

8.0 THE LIST OF PRESCRIBED SUBSTANCES

Volatile organic compounds.

9.0 DESCRIPTION OF TECHNIQUES USED FOR PREVENTING RELEASES

The whole production strategy at the Container Solvent Plant is to reduce losses due to evaporation or other emissions, particularly of hexane and acetone both of which are expensive raw materials and account for something like £250 per tonne.

The first major improvement that was carried to the plant to reduce emissions was the fitting of pipe bundle condensers to each of the vents from each of the mixers and bulk tanks which has had a marked effect in reducing losses in 1989 of approximately 88 tonnes down to 28-48 tonnes which is the current loss rate which represents approximately 1-1½% of total production for hexane based materials. Additional improvements have also been made by way of increasing both the pump capacity to achieve better circulation and also to improve the efficiency of the cooling tower by fitting a new honeycomb and also descaling the heat exchangers. The combined effect of these improvements has resulted in the reduction of the temperature of the cooling water and thereby improved the cooling facilities at the homogeniser which has resulted in the output temperature from the homogeniser being reduced from the initial boiling point of hexane to below it thereby reducing still further the potential for emissions to atmosphere. Emissions to atmosphere from the rubber compounding room are controlled via dust extraction units of which there are 3 which are vented at points 1, 2 and 3 as shown on the site plan.

10.0 DETAILS OF SOURCE, NATURE AND AMOUNT OF EMISSIONS

The principal source of the emissions will be via solvent losses from the solvent plant itself from each of the vents to atmosphere as shown on the plan which are numbered 4-15. The principal losses are likely to occur from the mixers and the solvent blending tanks. There will however, be some fugitive emissions which will result from the filling out process and also from the charging of the rubber into the mixer. In the latter case this is achieved by displacing hexane vapour when the rubber master batch is added. Exhaust ventilation is provided around the charge opening of each of the vessels to control vapour principally for the purposes of ensuring that operators are not exposed to significant concentrations of hexane vapour which could endanger their health.

The actual losses which arise from the production facility have been calculated from accurate mass balance calculations which are set out below:

Para 10.1 shows the total losses which have been corrected for loss of moisture, scrap to compound as well as losses due to bag retention and also dust collection in the dust extractors. The corrected figures are shown in Para 10.2.

10.1 Hexane Process

(1) Total Losses

Period	Production	Total Loss	% Loss
November 1989/October 1990	4386001	106958(1)	2.44
November 1990/October 1992	4916827	48061(1)	0.98
November 1991/June 1992	3354068	55981(2)	1.67

Note (1) Total loss is calculated from all raw materials received less product produced corrected for opening and closing stocktake figures.

Stocktaking takes place at October month end each year.

Note (2) Eight month figure based on a stocktake of only solvent and semifinish and consequently inherently less accurate.

(2) Corrected Losses

The rubbers and fillers used in the process contain volatile components principally water. during the production of a rubstock in a banbury mixer the temperature reaches 130°C and moisture is driven off.

Additionally, during packing of the final product filtration occurs removing any undispersed material.

Filters are cleaned into drums and this waste product is disposed of as notifiable waste losses which can be attributed to these two sources are as follows:-

	November 1989/ October 1990	November 1990/ October 1991	November 1991/ June 1992
Total loss (kg) Moisture	106958 9052	48061 10167	55981 6963
Compound scrapped (2 per month x 170 kg)	4080	4080	4080
Less Bag losses	3702	4161	2855
Dustextractor Collections	<u>1150</u>	<u>1150</u>	<u>1150</u>
Corrected loss (kg)	<u>88974</u>	28503	40933

Corrections have been made for the following of 0.1 kg per bag.

Bag Ullage

Bags used November 1989/October 1990	$37028 \times 0.1 = 3702.8 \text{ kg}$
Bags used November 1990/October 1991	$41616 \times 0.1 = 4161.6$
Bags used November 1991/June 1992	$28556 \times 0.1 = 2855.6$

Dust Extractor Collections

Extraction system collects 25 kg/week = 1150 kg (46 week year).

(3) Comparison of Corrected Losses with Solvent Losses

Period	Solvent Required for Products Produced (1) (tonnes)	Solvent Purchased (2)	Difference	Calculated Corrected Loss
November 1989/90	2631	2683	52	88.9
November 1990/91	2949	2982	33	28.9
November 1991/ June 1992	2006	2038	32	40.9

Notes: 1. Using average total solids.

2. Receipts corrected for opening and closing balances.

10.2 Acetone Process

Total Losses

Period	Production (kg)	Total Loss (kg)	% Loss
November 1989/October 1990	288500	8197(1)	2.84
November 1990/October 1991	315970	4405(1)	1.39
November 1991/June 1992	163970	7593(2)	4.63

Note 1

Total loss calculated from all raw materials received less product produced corrected for opening and closing stocktake figures.

Stocktaking takes place at October month end each year.

Note 2

Eight month figure based on a stocktake of only solvent and semi-finish and consequently inherently less accurate.

10.3 Isopar G Process Losses

Period	Production (kg)	Total Loss (kg)	% Loss
November 1989/October 1990	46940	952	2.03
November 1990/October 1992	59940	940	1.57
November 1991/June 1992	34812	198	0.56

The total losses therefore from each of the 3 basic processes are as follows based on 1991/92 mass balance figures.

Hexane process - 40.9 tonnes Acetone process - 7.5 tonnes Isopar G process - 0.19 tonnes.

Total VOC loss - 48.95 tonnes per annum.

11.0 ASSESSMENT OF LIKELY ENVIRONMENTAL CONSEQUENCES

11.1 General

VOC's emitted to atmosphere have the ability to react with other pollutants in the atmosphere principally oxides of nitrogen to form both ozone and other chemicals known as photochemical oxidants. Visibility can be impaired by such particulates formed during the photochemical reaction and the oxidants and their precursors can be transported over long distances. The essential problem is that whilst ozone in the upper atmosphere is essential as a shield against UV radiation from the sun excess ozone near ground level can present a problem if it is not dispersed readily.

11.2 The actual level of VOC emission from the solvent plant amounts to less than 50 tonnes per annum equivalent to approximately 1-1½% of total production throughput. This represents a relatively small amount and hence the potential environmental impact is likely to be insignificant. Over the 20 years that the plant has been in operation there have been no complaints from residents within Cromwell Road even though the emission levels have been higher than those that exist at present.

12.0 DETAILS OF MONITORING

Monitoring within the plant has been carried out over a number of years to determine exposure of employed persons to both dust and fume and generally good control has been achieved. As a special exercise however, for the purposes of this application atmospheric monitoring was carried out around the perimeter of the plant to determine the concentration of hexane and in particular any presence of n-hexane. The results of this monitoring are shown in Tables 1, 2 and 3. Table 1 shows the results of the environmental sampling exercise exterior to the plant and Tables 2 and 3 confirm the results of sampling in respect of worker exposure to both dust and fume.

As a general comment there appears to be little cause for concern with very little if any presence being detected of either isohexane or n-hexane.

TABLE 1 WR Grace St Neots - Tables and Results

No 1 Pumped Tube Samplers for Environmental Solvent Emissions

Description of Test		Conc ⁿ of Solvent in Air (ppm)					
See Attached Plan for Locations		isohexane (all isomers)	n-hexane				
Large NIOSH carbon tubes, pumped at 200ml/min. Analysis by CS ₂ desorption and GC.							
Samples taken 23th June 1992 (approximately 07.30 to 12.00)							
A1	By entrance gate	ND	ND				
B1	On fence near roadway	Trace < 0.2	ND				
C1	On fence near boundary	ND	ND				
D1	On fence far corner	Trace, 1.4	ND				
E1	On fence railway side	Trace 1.2	ND				
F1	On lamp post 10 ft high	Trace < 0.2	ND				
G1	"Worst case" in plant	19	ND				
Samples taken 24.6.92 (approximately 11.45 to 16.30)							
A2	As above.	ND	ND				
B2	As above.	2.4	ND				
C2	As above.	ND	ND				
D2	As above.	ND	ND				
E2	As above.	ND	ND				
F2	As above.	Trace <1.8	ND				
G2	As above.	57	Trace < 0.2				

Notes:

1. "Trace" means a slight detection of the substance was found, but that the signal to noise ratio on GC resulted in a reporting error of greater than $\pm 50\%$.

Where the figures is quoted without "trace", error is within 95% confidence limit.

- 2. All results are time weighted averages over the sampling periods quoted.
- 3. The occupational exposure standards are as follows: isohexane 500 ppm, n-hexane 20 ppm (reference HSE Guidance Note EH 40/92). Samples G1 and G2 are the only relevant ones to compare directly with this standard.
- 4. As a "rule of thumb", a figure of one thirtieth of the o.e.s. may be taken as "reasonable" environmental standard as regards hazards to health to persons outside the plant. This gives provisional standards of 16.7 ppm and 0.7 ppm respectively for assessing samples A1-F1 and A2-F2 inclusive.

TABLE 2

No 2 Vapour Absorber Badges for Worker Exposure to Solvents

Description of Test	Conc ⁿ of Solvent in Air ppm		
3M Vapour Badges, exposed full shift, analysis by CO ₂ desorption and GC	isohexane (all isomers)	n-hexane	acetone
A. Samples Taken 23rd June 1992			
OU 4905 operator 1, charging hexane mixer (4 off) and acetone mixer (1 off).	73	Trace < 0.5	75
OU 5088 operator 2, packing hexane based products into drums.	21	None detected	
OU 5040 operator 3, general duties around plant.	1.7	None detected	-
OU 5066 Static "engineering" sample, on Homogenises, running all day.	166	Trace < 0.5	-
OU 5159 Static sample, mixer conveyor over mixer neck (hexane compound).	173	Trace < 0.5	-
B Samples Taken 24th June 1992			
OU 5059 operator 1, charging hexane mixer.	119	None detected	-
OU 4959 operator 2, packing hexane product and acetone product into drums.	49	None detected	35
OU 5042 operator 3, general duties	None detected	d -	-
OU 5063 Static sampler on homogeniser, hexane compound,	69	Trace < 0.5	•
OU 4986 Static sampler on mixer neck, hexane compound	165	Trace < 0.5	-

Notes

"Trace" means an indication on GC but below reliable reporting limits, approximately 1 ppm in air.

A dash in the 3rd column means analysis was not conducted.

TABLE 3

No 3 Rubber Dust and Fume Samplers

Description of Test

Concⁿ of Particulate in Air, mg/m³

To HSE MDHS 14 "Rubber Fume" methods with Soxhlet extraction using cyclohexane (See separate notes for production schedules) Total Resp Solvent Soluble						
A	Samples Taken 30th June 1992					
1.	Personal, operator 1, mill operator and bag splitting (Cyclone)	0.89	0.30	-		
2.	Personal, operator 2, Banbury operator (Cyclone).	1.42	0.32	-		
3.	Static "engineering" sample, on scales by Banbury mixer (open).	1.58	-	0.40		
4.	Static sample, by small transfer conveyor, on ground floor (open).	0.84	-	0.22		
В	Samples Taken 1st July 1992					
1A	as 1 above (Cyclone)	1.19	0.39	-		
2 A	as 2 above (Cyclone)	1.68	0.30	-		
3A	as 3 above (open)	1.33	-	0.32		
4A	as 4 above (open)	0.86	-	0.24		

Notes

- (1) The reference to "cyclone" above is a miniature Higgs and Dewall personal sampler for respirable and total dust. "Open" refers to 7 hole "UK" heads with pre weighed and solvent extracted GFA filters, for subsequent soxhlet extraction.
- (2) A dash in any column means that parameter was not analysed.
- (3) OES for rubber fume is 0.6 mg/m³ solvent soluble fraction.

13.0 MONITORING OF ENVIRONMENTAL CONSEQUENCES

It is not proposed to carry out any monitoring of the environmental consequences as the monitoring already carried indicates that the current emissions levels are considered to be insignificant in relation to their environmental impact.

14.0 MONITORING OF TECHNIQUES FOR PREVENTING RELEASES AND/OR EMISSIONS

There are very limited techniques that can be used for monitoring the effectiveness of the prevention techniques currently employed. The key areas for consideration are regular maintenance of condensing equipment and any plant or equipment associated with the heat exchangers. This will mean therefore a preventive maintenance regime for cleaning out the cooling tower, associated heat exchangers and also for ensuring that the condensers have the correct surface area available and that they are running with the correct level of cooling water flow. No further monitoring at the perimeter of the plant is proposed although monitoring of the workplace will continue for the purposes of the Control of Substances Hazardous to Health (COSHH) regulations.

15.0 COMPLIANCE WITH OBJECTIVES IN SECTION 7(2) OF THE ENVIRONMENTAL PROTECTION ACT

In considering the best available techniques available for controlling emissions to atmosphere of VOC's further considerations will include the evaluation of

- 1. Installation of heat exchangers on the infeed to the homogenisers to reduce the temperature of the compound and thereby reduce any potential emissions.
- 2. The installation of heat exchangers in the packing lines to reduce the temperature of compound during packing and thereby prevent further any losses etc due to fugative emissions.

16.0 POSSIBLE PROPOSALS FOR IMPROVEMENTS

A considerable amount has already been done prior to the introduction of the Environmental Protection Act 1990 to reduce emissions to atmosphere and hence there is very limited scope for further improvement that can be regarded as cost effective within the meaning of BATNEEC. There are possibilities however, in terms for controls and some evaluation may be possible of controlling mixer temperatures by automatic temperature control. There is also a possibility of considering tanker line balancing during off loading of raw materials.

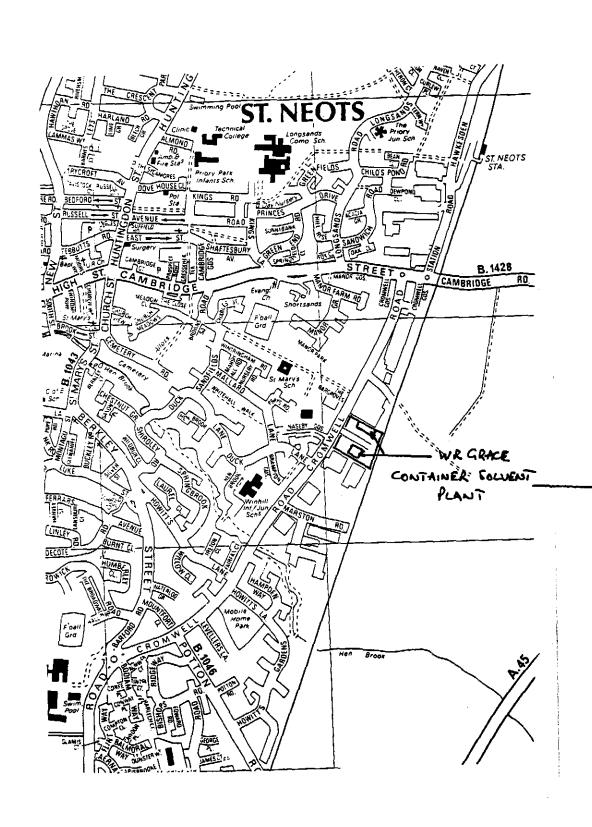
A the present time it is considered that the solvent plant currently operates to BATNEEC standards.

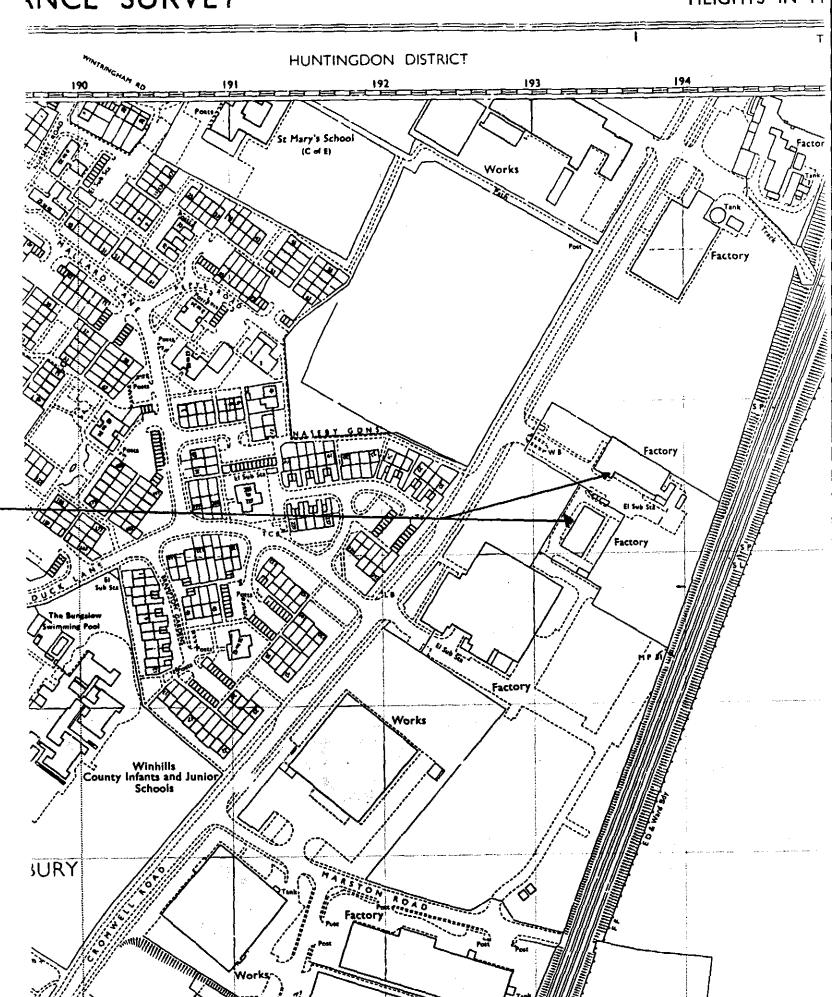
APPENDIX 1 ORDNANCE SURVEY MAP OF AREA

SIM (SUPPLY OF SURVEY

NCE SURVEY

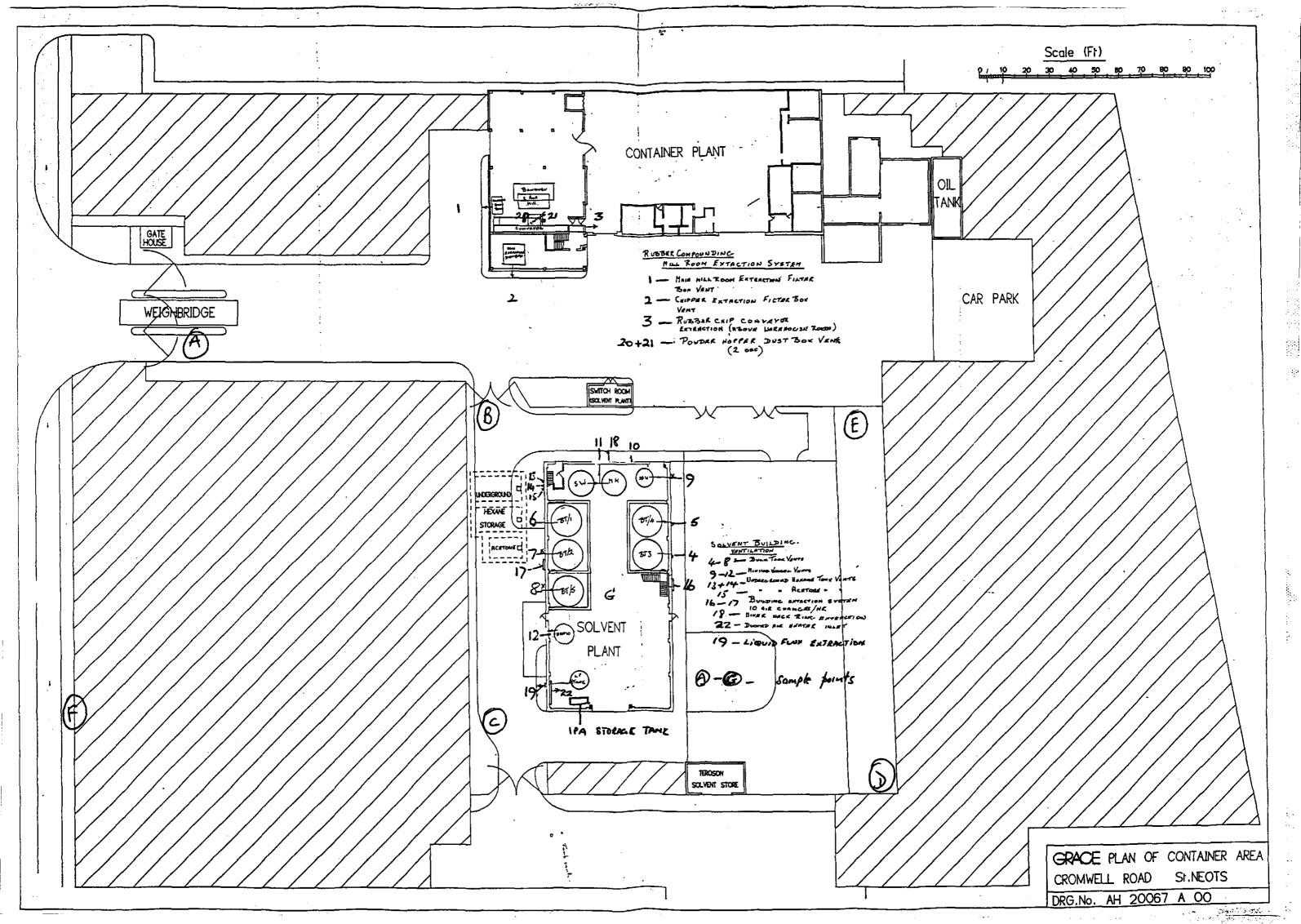
HEIGHTS IN M





APPENDIX 2

SITE PLAN



APPENDIX 3

MATERIAL SAFETY DATA SHEET DAREX 100A/39 LVE

TRADE_NAME:

DAREX SLC 100A/39LVE

CHEMICAL FAMILY:

Solvent dispersion of a rubber based composition.

INGREDIENTS:

Volatiles:

61% w/w

C₅ - C₇ fraction from petroleum

(< 0.1% benzene, <5.0% n-hexane)

Non-volatiles:

39% w/w

rubber

resin

inert mineral fillers

pigments

antioxidants

PROPERTIES:

Viscous grey dispersion with odour of petroleum.

Boiling point Vapour pressure at 38°C Specific gravity

65-95°C

330mm Hg

0.82

FIRE AND EXPLOSION DATA:

(C₅-C₇ fraction)

Highly flammable Flash Point Lower explosive limit

<-30°C

0.9% v/v

FIRE EXTINGUISHING MEDIA:

Use CO_2 or dry chemical on small fires; foam or water spray on large fires.

Issue 11 July 1992

Page 1 of 3

DAREX SLC 100A/39LVE

SPECIAL FIRE-FIGHTING PRECAUTION:

Fire-fighters should be equipped with self-contained breathing apparatus.

PRINCIPAL HEALTH HAZARDS:

In high concentrations well in excess of the OES⁽¹⁾ the solvent vapours are irritating to the eyes, nose and throat and cause dizziness, nausea, lack of co-ordination and, ultimately, unconsciousness.

PRECAUTIONS IN NORMAL USE:

Avoid sparks and open flames.

Avoid inhaling vapour.

Applicable standards: OES(1)

n-hexane $100 \text{ppm} = 360 \text{mg/m}^3$ cyclohexane $300 \text{ppm} = 1050 \text{mg/m}^3$ n-heptane $400 \text{ppm} = 1600 \text{mg/m}^3$ pentane (all isomers) $600 \text{ppm} = 1800 \text{mg/m}^3$ others $500 \text{ppm} = 1800 \text{mg/m}^3$

For the mixture of hydrocarbon isomers in the solvent system present in SLC 100A/39LVE, an OES of $400 \mathrm{ppm} = 1400 \mathrm{mg/m^3}$ is recommended.

Avoid contact with skin and eyes.

Wear impervious gloves, goggles and protective clothing.

DAREX SLC 100A/39LVE

IN EVENT OF SPILLAGE:

Absorb onto sand or earth and sweep up. Dispose of in accordance with local authority regulations.

WASTE DISPOSAL:

This material may be disposed of at official landfill tips under approved conditions in compliance with the requirements of the Control of Pollution Act 1974 and the Control of Pollution (Special Waste) Regulations 1980.

Advice on disposal should always be sought from the Local Authority Waste Disposal Officer; unofficial dumping should be avoided at all times.

Care should be exercised in disposing of 'empty' drums of this material as they are likely to contain solvent vapours in concentrations greater than their lower explosive limit.

FIRST AID PROCEDURE:

Inhalation Remove from exposure and obtain medical attention.

Eye contact Immediately flush with water for 15 minutes. Obtain

medical attention.

Skin contact Wash immediately with soap and water.

Ingestion Do not induce vomiting. Obtain immediate medical

attention.

REFERENCE:

(1) Occupational Exposure Standard, Recommended Limit in Guidance Note EH 40/89 from the Health and Safety Executive (UK).

Issue 11 July 1992

Page 3 of 3

APPENDIX 4

MATERIAL SAFETY DATA SHEET DAREX SLC 239

TRADE NAME:

DAREX SLC 239

CHEMICAL FAMILY:

Acetone dispersion of a rubber based composition.

INGREDIENTS:

Volatiles:

acetone propan-1-ol 53.5 3.0 56.5% w/w

Non-volatiles:

43.5%

rubber inert mineral fillers pigments

antioxidants

PROPERTIES:

Viscous grey dispersion with odour of acetone.

Boiling point (acetone) Specific gravity at 20°C

56°C 1.03

FIRE AND EXPLOSION DATA:

(Acetone)

Highly flammable Flash Point CC Lower explosive limit

0°C

2.6% v/v

FIRE EXTINGUISHING MEDIA:

Use CO₂ or dry chemical on small fires; foam or water spray on large fires.

Issue 9, July 1992

Page 1 of 3

DAREX SLC 239

SPECIAL FIRE-FIGHTING PRECAUTION:

Fire-fighters should be equipped with self-contained breathing apparatus.

PRINCIPAL HEALTH HAZARDS:

Acetone vapours is irritating to the eyes, nose and throat. Propan-1-ol vapours are harmful by inhalation and irritating in the eye.

PRECAUTIONS IN NORMAL USE:

Avoid sparks and open flames.

Avoid inhaling vapour.

Applicable standards: OES (1)

acetone propan-1-ol $1000 \text{ppm} = 2400 \text{mg/m}^3$ $200 \text{ppm} = 500 \text{mg/m}^3 \text{ (sk)}$

Avoid contact with skin and eyes.

Wear impervious gloves, goggles and protective clothing.

IN EVENT OF SPILLAGE:

Flush away with large volumes of water into sewer in accordance with local authority regulations.

DAREX SLC 239

WASTE DISPOSAL:

This material may be disposed of at official landfill tips under approved conditions in compliance with the requirements of the Control of Pollution Act 1974 and the Control of Pollution (Special Waste) Regulations 1980.

Advice on disposal should always be sought from the Local Authority Waste Disposal Officer; unofficial dumping should be avoided at all times.

Care should be exercised in disposing of 'empty' drums of this material as they are likely to contain solvent vapours in concentrations greater than their lower explosive limit.

FIRST AID PROCEDURE:

Inhalation Remove from exposure and obtain medical attention.

Eye contact Immediately flush with water for 15 minutes. Obtain

medical attention.

Skin contact Wash immediately with soap and water.

Ingestion Do not induce vomiting. Obtain immediate medical

attention.

REFERENCE:

(1) Occupational Exposure Standard, Recommended Limit in Guidance Note EH 40/89 from the Health and Safety Executive (UK).

APPENDIX 5

MATERIAL SAFETY DATA SHEET DAREX SLC 9291E/E

TRADE NAME:

DAREX SLC 9291E/E

CHEMICAL FAMILY:

Solvent dispersion of a rubber based composition.

INGREDIENTS:

Volatiles:

57.0% w/w

C₅ - C₇ fraction from petroleum

(< 0.1% benzene, <5.0% n-hexane)

Non-volatiles:

43.0% w/w

rubber resin

inert mineral fillers

pigments antioxidants

PROPERTIES:

Viscous grey dispersion with odour of petroleum.

Boiling point Vapour pressure at 38°C Specific gravity 65-70°C 330mm Hg

0.85

FIRE AND EXPLOSION DATA:

 $(C_5 - C_7)$

Highly flammable Flash Point Lower explosive limit

<-30°C 0.9% v/v

FIRE EXTINGUISHING MEDIA:

Use CO₂ or dry chemical on small fires; foam or water spray on large fires.

Issue 9, July 1992

Page 1 of 3

DAREX SLC 9291E/E

SPECIAL FIRE-FIGHTING PRECAUTION:

Fire-fighters should be equipped with self-contained breathing apparatus.

PRINCIPAL HEALTH HAZARDS:

In high concentrations well in excess of the OES⁽¹⁾ the solvent vapours are irritating to the eyes, nose and throat and cause dizziness, nausea, lack of co-ordination and, ultimately, unconsciousness.

PRECAUTIONS IN NORMAL USE:

Avoid sparks and open flames.

Avoid inhaling vapour.

Applicable standards: OES (1)

For the mixture of hydrocarbon isomers in the solvent system present in SLC 9291E/E, an OES of $400 \text{ppm} = 1400 \text{mg/m}^3$ is recommended.

Avoid contact with skin and eyes.

Wear impervious gloves, goggles and protective clothing.

DAREX SLC 9291E/E

IN EVENT OF SPILLAGE:

Absorb onto sand or earth and sweep up. Dispose of in accordance with local authority regulations.

WASTE DISPOSAL:

This material may be disposed of at official landfill tips under approved conditions in compliance with the requirements of the Control of Pollution Act 1974 and the Control of Pollution (Special Waste) Regulations 1980.

Advice on disposal should always be sought from the Local Authority Waste Disposal Officer; unofficial dumping should be avoided at all times.

Care should be exercised in disposing of 'empty' drums of this material as they are likely to contain solvent vapours in concentrations greater than their lower explosive limit.

FIRST AID PROCEDURE:

Inhalation Remove from exposure and obtain medical attention.

Eye contact Immediately flush with water for 15 minutes. Obtain

medical attention.

Skin contact Wash immediately with soap and water.

Ingestion Do not induce vomiting. Obtain immediate medical

attention.

REFERENCE:

(1) Occupational Exposure Standard, Recommended Limit in Guidance Note EH 40/89 from the Health and Safety Executive (UK).

Issue 9, July 1992

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Public Notices

ENVIRONMENTAL PROTECTION ACT 1990. PART 1.

APPLICATION FOR AUTHORISATION

W R Grace limited has applied for an authorisation from Huntingdonshire District Council to operate a prescribed process, namely the manufacture of a solvent based coating material at Container Solvent Plant, Cromwell Road, St. Neots, Cambs PE19 2ER,

A copy of the application is available for public inspection, free of charge, at Pathfinder House, St. Marys Street, Huntingdon, Cambs PE18 6TN.

Written representations about the application may be. sent to Huntingdonshire District Council at the above

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19-Heart to Heart

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19-Heart to Heart 🛣

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MALE 17 Warped sense of sensitive female, looks reply Box No.P, 4007 (see

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4036 (see panel) .Two previous owners. Needs TLC. Smokes a little. Photo please. Please reply Box No.P 4008 (see

ROMANTIC Female 45 recently moved would like to meet friends, male/female to help her find happiness and company. Please reply Box No.P 4009 (see panel)

22-Public & Legal Notices

ENVIRONMENTAL PROTECTION ACT 1990, PART 1 🗱 🔠 🐔 APPLICATION FOR AUTHORISATION

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Written representations about the application may be sent to Huntingdonshire District Council at the above address within 28

KATHLEEN ANNIE HIBBERD DECEASED ***

PURSUANT to the Trustee Act 1925 ALC PERSONS having claims against or claiming to be beneficially interested in the estate of Kathleen Annie Hibberd late of 28 Main Street, Hartford, Huntingdon, Cambridgeshire who died on the 28th day of July 1992 are required to send particulars thereof in writing to the undersigned on or before the 19th November 1992 next after which the Executor will proceed to distribute the assets having regard only to the claims of which he has then had notice. Dated this 17th day of September 1992.

: Leed Day Solicitors to the said Executor

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THE COUNTY OF CAMBRIDGESHIRE (40mph SPEED LIMIT) (No7) ORDER 1992 IMPOSITION OF 40mph SPEED LIMIT AT BROUGHTON

NOTICE is hereby given that on the 8th September 1992, the County Council made an Order under Sections 84(1) and 84(2) of the Road Traffic Regulation Act 1984, the effect of which is to impose a 40mph speed limit in Broughton as follows:

School Road - From its junction with Bridge Road and Causeway Road in a north westerly direction for a distance of 450 metres. 💯

Bridge Road - From its junction with School Road and Causeway Road in a south

westerly and south easterly direction for a distance of 650 metres. Causeway Road - From its junction with School Road and Bridge Road in a north

easterly and south easterly direction for a distance of 435 metres. A copy of the Order, which comes into operation on the 21st September 1992, together with a map showing the roads and lengths of road concerned, may be examined at the office of the undersigned, and at the offices of The Chief Executive, Huntingdonshire District Council, Pathfinder House, St. Mary's Street, Huntingdon

Any person wishing to question the validity of the Order or of any provision contained in it on the grounds that it is not within the powers conferred by the Road Traffic Regulation Act 1984 as amended, on the grounds that any requirement of that Act or of any instrument made under it has not been complied with in relation to the Order may, within six weeks from the 8th September 1992, apply to the High Court for

J.M. SHARPE, Director of Transportation, Shire Hall, Castle Hill, Cambridge CB3 0AP. 16th September 1992.

CAMBRIDGESHIRE COUNTY COUNCIL NOTICE OF CONFIRMATION OF RECLASSIFICATION ORDER WILDLIFE AND COUNTRYSIDE ACT 1981

Definitive Map and Statement of Public Rights of Way for the former County of Huntingdon dated 9th September 1953 as reviewed 14th December 1961.

The County of Cambridgeshire (RUPP No.3 Waresley) Definitive Map Reclassification

On 26th August 1992, following a local public inquiry, an inspector appointed for the purpose by the Secretary of State for the Environment confirmed the above named order. The effect of the order, as confirmed, is to modify the Definitive Map and Statement for the area by reclassifying as a Byway Open to All Traffic the Road Used as a Public Path No.3 Waresley commencing on the St. Neots Road by Wildmere Plantation and running in a easterly direction to its termination on the St. Ives Road,

Copies of the order and the map contained in it have been deposited at the offices of the Chief Executive, Huntingdonshire District Council, Pathlinder House, St. Mary's Street, Huntingdon and of the undersigned and may be inspected free of charge during normal office hours or purchased at a cost of £2 from the address below.

The order became operative as from 26th August 1992, but if any person aggrieved by the order desires to question the validity thereof, or of any provision contained therein on the ground that it is not within the powers of the Wildlife and Countryside Act 1981, or on the ground that requirement of the Act, or of any regulation made thereunder, has not been complied with in relation to the order, they may before Thursday 29th October 1992 make any application of the purpose to the High Court.

Dated this Thursday 17th September 1992. J.M. SHARPE, Director of Transportation, Castle Court, Shire Hall, Cambridge CB3 OAP



DEPARTMENT OF TRANSPORT ROAD TRAFFIC REGULATION ACT 1984 - SECTION ·14(1)

THE AT TRUNK ROAD (STIBBINGTON TO WATER **NEWTON AND MORBORNE TO NORMAN CROSS** ROUNDABOUT, CAMBRIDGESHIRE) (TEMPORARY PROHIBITION AND RESTRICTION OF

TRAFFIC) ORDER 1992 Notice is hereby given that the Secretary of State for Transport proposes to make an Order on the A1 Trunk Road in Cambridge-

The Order, which is being made to enable resurfacing work to be carried out on the southbound carriageway of the A1 Trunk Road between Stibbington and Water Newton and between Morborne turn and Norman Cross Roundabout will be in force when the appropriate signs are erected from 21 September 1992 until the works are completed. The work is expected to start on 5 October 1992 and will last approximately 3 weeks.

The effect of the Order will be to prohibit vehicles commencing at 1900 hours for a period of 11 hours, from -

a) proceeding at a speed exceeding 30 miles per hour in the southbound carriageway,

b) using the layby located at: 300 metres north of the Nene Valley Railway, 600 metres north of the Nene Valley Railway, 1300 metres north of the Nene Valley Railway, or 710 metres south

Access to Manor House 400 metres south of the Nene Valley Railway and the Bungalow 1030 metres south to Morborne turn will be

Any vehicle being used in connection with the said works or for police, fire brigade or ambulance purposes will be exempt from the restrictions imposed during the works.

MRS T NASH

A Higher Executive Officer In the Department of Transport

832 SE

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