GUIDANCE ON CONTROL OF EXPOSURES RELATED TO USE OF CARBON TETRACHLORIDE SOLVENT SUBSTITUTES IN THE OXYGEN MANUFACTURING AND REFILLING SECTOR











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The reader is advised to confirm the product specifications and related health/environmental hazards prior to purchase or use of any of the solvents profiled. No claim is made here for the absolute suitability of any solvent as a substitute for CTC in any application. Suitability of a product or method of cleaning for a particular application would need to be verified through trials prior to any larger-scale application with due consideration of health and safety aspects.

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PREFACE

Carbon Tetrachloride (CTC) is widely used as a solvent in many industrial sectors in India. It is an ozone depleting substance (ODS) similar to chlorofluorocarbons (CFCs). The UV-B and UV-C radiation coming from the sun interacts with CTC molecules that drift into the stratosphere and release their chlorine atoms. Each chlorine atom can destroy as many as 100,000 ozone molecules over a period of nearly 100 years. Thus, even a small amount of CTC released into the environment can cause tremendous damage to the ozone layer. Further, the global warming potential (GWP) of CTC has been estimated to be about 1,400 times higher than that of carbon dioxide (CO₂), the principal greenhouse gas. It is also hazardous to health via all routes of exposure viz., inhalation, ingestion and skin absorption. There is sufficient evidence of carcinogenicity in experimental animals with an increasing body of evidence for being a human carcinogen.

To protect the ozone layer, India is one of the 196 signatories to the Montreal Protocol for phasing out the production and consumption of ozone depleting substances. Under this agreement India has committed to phase-out the use of CTC solvent completely by 31st December 2009.

As the phase-out is progressing, CTC supplies in the market are dwindling rapidly. Beyond 31st December 2009 CTC will not be available for use as a solvent. Given the reduction of supply, the price of CTC has risen substantially making it costlier today, than most of its alternatives.

Within the framework of the Multilateral Fund of the Montreal Protocol, the Governments of Germany and France have mandated GTZ-Proklima to provide technical assistance to CTC consuming industries in India. In addition, World Bank, UNIDO and UNDP (on behalf of the Government of Japan) are assisting India, to address this issue specifically in industry sectors with large volumes of CTC use. These activities are coordinated under the National CTC Phase-out Plan by the World Bank as the leading implementing agency and the Ozone Cell of the Ministry of Environment and Forests, Government of India.

GTZ-Proklima offers technical assistance to industries using up to 10 metric tons of CTC per year. In close interaction with these industries, GTZ-Proklima aims to provide guidance in identifying CTC substitutes by addressing environmental, health and safety concerns without compromising on quality and cost effectiveness.

The Department of Environmental Health Engineering (DEHE), Sri Ramachandra University (SRU), Chennai (India) prepared a list of nearly 500 potentially hazardous substances that could be present in proprietary or non proprietary cleaning agents and provided the same to the GTZ to enable comparisons across potential CTC substitutes. Moreover, initial assessments at different workplaces indicated the potential for occupational exposures to solvent vapours (from products currently being used as CTC substitutes). Building on these earlier efforts, the present exercise was aimed at conducting a systematic evaluation of exposures and identification of risks associated with selected CTC substitutes across a spectrum of few *Oxygen Manufacturing and Refilling* industries in order to provide specific recommendations for control against workplace exposures to the solvents used.

The approaches adopted for risk management in the project have a dual focus on "risk prevention" through substitution and "risk mitigation" through installation of workplace engineering controls and improving general work practices.

Recommendations presented in this manual have been largely based on walkthrough visits and analytical results of workplace exposure measurements. Participatory discussions involving employers and professional organizations were used to generate feasible toolkits that can be readily implemented. Since improved handling also minimizes emissions and exposures, this manual is expected to contribute not only in improving occupational health and safety at work, but also facilitate the implementation of the Montreal Protocol with added environmental protection benefits as well.

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ABBREVIATIONS

ACGIH American Conference of Governmental Industrial Hygienist

CAS Chemical Abstract Services

CFCs Chlorofluorocarbons

CIM Centre for International Migration

CTC Carbon tetrachloride

EHS Environment Health and Safety

GG Spectacles/Goggles

GL Gloves

GTZ German Technical Cooperation

GWP Global Warming Potential

LC50 Lethal Concentration

LD50 Lethal Dose

MSDS Material Safety Data Sheet

NIOSH National Institute for Occupational Safety and Health

ODS Ozone Depleting Substance
OEL Occupational Exposure Limits

OSHA Occupational Safety and Health Administration

PCE Perchloroethylene

PEL Permissible Exposure Limits
PFTs Pulmonary Function Tests

PLE Permissible Limits of Exposure
PPEs Personal Protective Equipments

R Respirator

SRU Sri Ramachandra University
STEL Short Term Exposure Limit

TCE Trichloroethylene

TLV Threshold Limit Value
TWA Time Weighted Average

UNDP United Nation Development Programme

UNIDO United Nation Industrial Development Organization

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1. INTRODUCTION

Oxygen is manufactured from atmospheric air involving processes that use special solvents for cleaning contaminants from the machines and accessories during the manufacturing and refilling process. The removal of contaminants and the use of solvents in an unsafe manner can cause health and safety risk at the workplace. While oxygen is essential to sustain life on earth under normal range of concentrations, (at around 21 percent in air) deficient and enriched oxygen environments can be harmful and dangerous to human life and ecosystem. Oxygen is not flammable by itself but it supports combustion processes and can cause accidental fire and explosion with the contaminated hydrocarbons trapped in the machineries and accessories used in certain industrial processes. The higher the oxygen content and/or pressure in a system, the more vigorous will be the combustion and can reach quickly to explosive levels.

Because of the inherent flammable and combustible properties of oxygen in presence of hydrocarbons, it is critical to select a solvent with great care for cleaning the contaminants in oxygen manufacturing industries. In many of the oxygen manufacturing industries, Carbon tetrachloride (CTC) is being used for cleaning of valves (commonly known as nozzles), cylinders, compressor casings and heat exchangers because of its cleaning efficiency and nonflammable properties. The most common contaminants encountered by the oxygen service industries are: machining oils (including residual oil film), hydrocarbon-based grease and lubricants (including compressor oil) and organic compounds such as human skin oil and body fluids including insects and insect body parts.

With the most widely used solvent, CTC being an Ozone depleting substance (ODS), there is now a legal obligation to use alternative cleaning agents. This manual has been prepared to provide guidance on "safe use" of potential CTC substitutes on the basis of results of workplace exposure measurements, work practice assessments and the opinion and outcome of concerned stakeholder discussions. The suggested interventions thus are likely to be both feasible and effective on a sector-wide basis to reduce occupational health risks. However, occupational health risks covered in this manual need to be matched with environmental aspects as well as cleaning efficiency and costs acceptable to the industry before adopting a product "substitute". Separate guidance on environmental and safety aspects that are general to all sectors using CTC substitutes is available at www.ctc-phaseout.org to enable the same.

2. WORK PRACTICES RELATED TO USE OF SOLVENTS IN THE OXYGEN MANUFACTURING / REFILLING SECTOR

Hydrocarbon-based grease and lubricants trapped in the cylinder and its accessories can create an accidental fire or explosion in the presence of oxygen rich environment. Hence, contaminants such as machine oils, hydrocarbon based grease and lubricants, soaps, detergents, human skin oil and body fluids, insects and its body parts and carbon dust that are present on surfaces of components requires to be inspected (**Figure 1**) and cleaned. This requires a non flammable solvent with good cleaning efficiency that will not leave any residual contaminants.



Figure 1: Oils on cylinder surface and nozzle.

Most of the industries use Trichloroethylene (TCE), CTC and detergents as cleaning agents, while few industries use acetone as a cleaning agent for selected components. The common work practices employed in this sector are described below:

2.1. Cleaning of compressor

During regular operation, oil and dust get accumulated on the compressor body and it requires periodical cleaning. This is usually carried out by "wiping method". A rag or sponge is dipped into the solvent and wiped (**Figure 2**) to remove oily or greasy dirt that has been accumulated on the compressor and its components. The cleaning is often done with bare hands and the rags are usually disposed in a manner that allows the solvent to evaporate directly into the atmosphere. Compressor casing and other parts are also cleaned during preventive maintenance and break down repairs.



Figure 2: Cleaning of compressor

2.2. Cleaning of cylinders

When empty cylinders are returned to the filling station, these are checked for oil stains and contaminants on the nozzle and surface of cylinders. When oils are identified on cylinders that have no positive pressure, the interior of the cylinders are subjected to physical inspection using incandescent bulb. Then, the contaminants are cleaned by filling the cylinder with solvent and are rinsed by rolling the cylinder. Usually, the nozzles and cylinder surfaces are cleaned by wiping method (**Figure 3**).



Figure 3: Cleaning of cylinder surface

2.3. Testing of cylinders

Cylinders are subjected to periodical hydro stretch tests at regular intervals. During this process (**Figure 4**) cylinders are rinsed with CTC/TCE followed by nitrogen flushing.

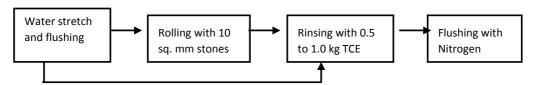


Figure 4: The Hydro Stretch Test Process

2.4. Cleaning of machines and components

Many components require routine cleaning (**Figures 5** to **8**). These include large components such as surface cleaning of machine by wiping method and small components such as nozzles and hose pipes by immersion method in conjunction with process such as agitation, scrubbing, wiping, spraying or flushing. It is important to ensure that no residual solvent or contaminant remains on the parts after the cleaning process.



Figure 5: Oils on surface of the machine



Figure 6: Oils on nozzles of machine



Figure 7:Air drying of nozzle



Figure 8: Cleaning of small components

3. HAZARD IDENTIFICATION AND RISK ASSESSMENT

The exposure concentration likely to be achieved with individual solvents primarily depends on physical, chemical and toxic properties, specific nature of cleaning procedures, work posture, work and hygiene practices, maintenance of machines and the workplace environmental conditions such as temperature, humidity and ventilation.

A key step in risk assessment is to recognize the job-specific exposure hazards and the associated risks. This can be done by shop floor visits/walk-through assessments, review of documents, interview with workers and supervisors and qualitative / quantitative assessments of workplace and environmental conditions. Based on a combination of such assessment methods, the following **Table 1** identifies the job-hazard-risk profile for key processes in the oxygen manufacturing and refilling sector.

Table 1: Job-hazard-risk profile

Job	Process/Hazard	Risk
Identification of contaminants	Identification of oil and grease contamination by physical and visual inspection.	Absorption through skin
Cleaning of components	Cleaning of nozzle by immersion	 Absorption through skin during immersion Exposure via lungs due to solvent evaporation from open container.
	Removal of residual solvent by spraying with inert gas.	 Exposure via skin, lungs and eyes due to solvent splashing and vaporization. Enhanced skin absorption due to spraying at high pressure with no/inappropriate gloves
Cylinder cleaning	Removal of surface contamination of the cylinder by wiping	Exposure through lungs, skin and eyes

Job	Process/Hazard	Risk
Compressor cleaning	Cleaning of compressor by wiping	 Exposure via lungs and eyes due to solvent splashing and vaporization. Enhanced skin absorption by not using gloves and absorption through open wounds or skin ruptures that are caused by friction with sharp edges.
Cylinder Testing	 Cleaning by rinsing and agitation Spillage of solvent during filling of solvent into the cylinder 	 Exposure via lungs during filling and draining. Skin absorption through hand and feet due to improper or no personal protection Contamination of water and soil through disposal of solvent waste.
Storage of cylinders Fire Hazard in the presence of solvent	Storage of cylinders near open/loose electrical connections	Risk of fire and explosion.
Disposal of cotton waste	Disposing cotton waste in workplace	 Exposure via lungs due to solvent evaporation from used cotton waste when stored within the work place. Risk of fire at the cotton waste disposal location.

4. EXPOSURE ASSESSMENT AND RISK CHARACTERIZATION

Exposure assessments that measure levels of solvent exposure under specific workplace conditions allow comparisons to be made with reference to the exposure standards. In order to capture alternative exposure conditions or wherever measurements are difficult to perform, exposures can also be simulated under laboratory conditions to estimate potential exposures that may be encountered in the workplaces. Such exercises in this sector have shown that while TCE levels consistently exceed the permissible exposure limits in most of the cleaning processes, PCE levels are usually lower than the permissible limits, with acetone levels intermediate between PCE and TCE. By using required quantity of solvents for specific cleaning applications, exposure concentrations are reduced significantly. **Box 1** emphasizes the level of reduction in exposure concentration achievable through use of optimal quantity of solvents.

Box 1: Optimum use of solvent and percentage reduction in exposure concentration.

Around 20 to 30 percent reductions in exposure levels are achieved through optimum use of solvents.

Results of hazard recognition and exposure assessments made at individual work locations in this sector were used to characterize risks and provide job specific recommendations for prevention and control of exposures as described in the sections below.

5. CONTROL GUIDANCE

Risk management involves application of a sequence of control measures to minimize the exposures. A typical hierarchy of controls involves hazard (i.e. solvent) substitution, engineering, administrative and personal protective measures. While exposures are central in arriving at a control strategy, often exposure reduction goals have to be matched with feasibilities based on worker preference, skills and attitudes of workers/management, workplace environmental conditions and cost of control measures.

The following sections provide general and process/work practice specific guidelines for the oxygen manufacturing and refilling sector.

5.1. General Work Practice Guidelines

It is the employer or the managements' responsibility to provide guidance and instructions to workers on environment, health and safety (EHS) management. At the same time it is employees' responsibilities to follow the instruction laid by the employer to create a healthier working environment. Although managements may have a policy that allocates resources for EHS management, it has to identify a responsible person to implement and supervise the EHS management programme. Simple generic guidelines are, therefore, separately provided for the supervisors/managers and workers to maintain minimum EHS standards. These guidelines are provided in **Tables 2** and **3**.

Table 2: General Guidelines for Managers and Supervisors

Storage and Handling

- Layout of storage facility should be planned at an early stage (Refer Figure 22 in the Part-I manual).
- Storage protocols should be prepared depending on the compatibility (Refer Figure 23 in the Part-I manual).
- MSDSs must be referred for chemical compatibility.
- In the event MSDSs are not available, information must be secured from the manufacturer/distributor.
- Adequate safety measures (First-aid, Firefighting equipments, emergency showers, PPEs etc.) must be provided in the storage facility.
- Safety equipment must be periodically checked for fitness and performance.
- Safety sign boards specific to hazards must be provided and placed at appropriate locations.
- Instructions on handling procedures must be provided to the workers.
- Expired chemicals must be returned to the manufacturer or disposed according to standard protocol.
- Empty containers must be returned immediately to the supplier or the manufacturer or disposed in a proper manner.

Labeling Personal hygiene Cleanup Labels to be prepared in local Routine implementation of clean Practice of good up schedule and procedures. language in accordance to MSDS. hygiene practice Label must contain key Identification of a responsible amongst the information (such as and trained person for cleanup workers. Provision of hand Flammability, Health Risks, etc.) activities. Hazard symbols must be provided Planning and implementation washing facility on the label. with adequate of periodical maintenance of soaps and Caution words and statements equipments and accessories. (Danger, Toxic, Irritant, Corrosive Deployment of spill control detergents. etc.) for hazardous chemicals Provision of a procedures to prevent health separate lunch need to be provided. and environmental risks. room. Damaged or worn out labels must Provision of appropriate PPEs Provision of be replaced. during cleanup of spills. adequate work Embossed labels can be requested Provisions of proper waste cloths. from the manufacturer. disposal protocols. Provision of shower stations.

Provision of training on

- The availability and importance of MSDS.
- Obtaining MSDS from the manufacturer or competent sources (Government or Safety Agencies) in case of non availability.
- Informing the workers on the importance of MSDS.
- Procedures and protocol development for safe handling of chemicals.
- Managing emergency situations.
- Risk assessment and implementing control measures.
- Evaluation of the training programs conducted for workers.
- Selection of appropriate PPEs.
- Importance of medical fitness for wearing PPEs and selecting workers for providing PPE.

Table 3: General Guidelines for Workers

Storage and Handling

- Follow storage and handling procedures.
- If safe handling procedures are not available, request your supervisor.
- Observe safety sign boards for recognizing the hazards.
- Handle any new chemicals in the presence of Manager/Supervisor at initial stage.
- Learn to use safety equipments.
- Use protective device while handling hazardous chemicals.
- Request for PPE if not available.
- Report any damage or inconsistency of safety equipments to the Manager/supervisor.
- Report if chemicals are used beyond expiry dates.
- Place back containers at designated location after use.
- Transport the containers in closed condition.
- Close the containers with air tight fittings to prevent spoiling of chemicals.
- Store drinking water separately away from the chemical store.

Labeling Personal hygiene Cleanup Request for label and Remove dust, oils and Wash hands before eating. label information in local dirt regularly from the Use soap for washing the hands. language machines and floors. Avoid smoking and eating at Read the label before Regularly clean up at workplaces. using the chemicals. end of the shift or end Avoid placing your fingers into of the day. Care should be taken to mouth, ears and nose while handling prevent label damage Report immediately chemicals. while transferring, about the spills to Trim the nails periodically to prevent transporting and managers/supervisors. chemical accumulation. handling. Request for Dress open wounds before handling Request to your MSDS/instructions in the chemicals. manager/ supervisor for local language for spill Wear clean work clothes. replacing the damaged cleanup. Change work clothes before leaving label. Immediately clean the the workplace. spills according to the Have a shower at the end of the shift instructions and (to reduce the dermal exposures and protocol. minimize the transportation of the Wear appropriate PPEs contaminants to home) during cleaning the Clearly mark the drinking water spills. bottles and always keep them a clean place.

Provision of training on

- Recognizing and understanding the hazardous nature and risk of using chemicals.
- Importance of material safety data sheet (MSDS)
- Good housekeeping procedures (Storage, handling, labeling and cleanup of chemicals).
- Handling the chemical spills and proper disposal methods.
- Managing emergency situations using first-aid, handling fire extinguishers, communications (persons to be contacted, contact phone number) etc.
- Checking the functioning of pollution control systems.
- Incident or failure reporting systems.
- PPE use, storage, cleaning and maintenance.

5.2. Process Specific Guidelines

Although generic guidelines provide an overall framework to design and implement an EHS policy, often supervisors and managers require customized guidance that is specific for a particular work situation and process, which is provided in **Table 4** and **5**. Each main process concerned with solvent use is provided with a set of engineering, administrative and personal protective controls applicable specifically for the nature of hazards and risks associated with the job. The guidance sheet may thus be used as a ready reckoner for implementing facility-wise or location specific controls. Instruction for using the guidance sheet is provided in **Box 2**.

Box 2: Instruction for reading the guidance sheet

For identifying job specific hazard, risk and control measures, read the guidance sheet from left to right across the row.

It must be emphasized that the guidance sheets only provide information on "safe use" of specific solvents being used in each process. Selection amongst solvent substitutes is governed by considerations that often go beyond occupational exposures. On the basis of a limited set of exposure measurements, it has been observed that in general PCE concentrations are the lowest owning to its higher boiling point as compared to acetone while, TCE concentrations consistently exceed prescribed exposure limits. In non-ventilated spaces, PCE concentrations may build up over time creating more risks of exposure. However, these two are reported as suspected carcinogens while the health effects of TCE are more evident than the effects of PCE.

Therefore, facilities are encouraged to prioritize the selected solvent against conditions required for safe use in a larger EHS (environment, health and safety) framework in order to provide best work practice in the process or sector. The information related to safe use of the solvents is provided in **Tables 6** to **8**.

Table 4: Job Specific Control Guidance Sheet

				Control Measures	
Job	Hazards/ Solvents	Risk/ Exposure Situation	Engineering	Administrative	PPES Refer Table 5 for PPE codes
Cylinder Cleaning The state of	 Acetone Trichloroethy lene (TCE) Perchloroethylene (PCE) 	 Inhalation, dermal (skin) and eye contact due to cleaning with bare hand, naked eyes, spills. Additional Inhalation and dermal risk by using inappropriate respirator, gloves. Additional Inhalation risk due to solvent evaporation from open or partially closed containers. 	 Transfer the solvent into small container using funnel to avoid spillage and splash. Cleaning area should be well ventilated (natural ventilation). Select the work location near a window in accordance to the wind direction Use mechanical ventilation like local exhaust system to dilute solvent vapours, in addition to natural ventilation 	 Use optimum quantity of solvent. Close the container after use. Restrict the presence of coworkers nearby, during the cleaning process. Prohibit dipping the cotton directly into the container to avoid skin contact and solvent contamination. Treat solvent soaked cotton as hazardous waste. Remove the spills immediately. Keep solvents away from ignition sources. Prohibit smoking and place visual sign boards in the workplace to avoid fire hazard. Acetone is highly flammable and is suggested only for surface cleaning under good or open ventilation condition. 	Gloves: Acetone: GL 1 TCE:GL 2 PCE: GL 3 Goggles/ Spectacles: GG 2 Use solvent resistant goggles Use power corrected goggle if necessary Not required if respirator (R2) is used Respirator: R1 or R2 Use this in the absence of LES/LEV

				Control Measures	
Job	Hazards/Solvents	Risk/ Exposure Situation	Engineering	Administrative	PPEs Refer Table 5 for PPE codes
Nozzle/valves Hose Pipe and tube cleaning	 Acetone Trichloroethyle ne (TCE) Perchloroethyle ne (PCE) 	 Dermal (skin) contact due to immersion with bare hand, Lungs and eye contact due to vapourization from open container and spills. 	Immersion cleaning should be carried out in well ventilated environment.	 To avoid excess use of solvent and to reduce the exposure, immerse the nozzle/valves in solvent for few minutes followed by flushing with inert gas. Close the container during cleaning. For hosepipe and nozzle cleaning, flushing with inert gas (Nitrogen) is mandatory 	Gloves: Acetone: GL 1 TCE: GL 2 PCE: GL 3 Spectacles/Goggles: GG 1 Use power corrected spectacle if necessary Not required if R2 is used In the absence of natural or mechanical ventilation, use R1 or R2 while cleaning more than four times in a day.

			Control Measures						
Job	Hazards/Solvents	Risk/ Exposure Situation	Engineering	Administrative	PPEs Refer Table 5 for PPE codes				
Compressor part cleaning The state of the s	 Acetone Trichloroethylene (TCE) Perchloroethylene (PCE) 	 Skin contact due to cleaning with bare hands. Enhanced skin absorption through open wounds or skin ruptures. Lung and eye contact from vapourization of solvent. 	Cleaning should be carried out in well ventilated environment.	 Restrict the presence of coworkers nearby during the cleaning process. Collect the solvent soaked cotton wastes in a bin with lid. Then follow proper disposal procedures. 	Gloves: Acetone: GL 1 TCE: GL 2 PCE: GL 3 • Use proper gloves that are flexible for the job and use forceps while gloves are not suitable for the process. Spectacles/Goggles: GG1 Use power corrected spectacle if necessary				

Table 5: Codes and types of personal protective equipments

Sl. No	Code	Recommended protection	PPEs (Gloves)
1	GL 1	8 hr: Butyl or latex/rubber gloves	Butyl Latex or Rubber
2	GL 2	8 hr: Poly Vinyl alcohol	
3	GL 3	Nitrile latex gloves	
Sl. No	Code	Recommended protection	Spectacles / Goggles
1	GG 1	Solvent resistant spectacle	
2	GG2	Solvent resistant goggle	
Sl. No	Code	Recommended protection	Respirator
1	R1	Half face respirator	
2	R2	Full face respirator	

Source for selection of gloves: OSH Technical reference material, Department of Energy, USA and Chemical Protective Clothing, National Institute for Occupational Health and Safety (NIOSH), USA.

Table 6: Data Sheet for Acetone

			Т	Coxic, C	Chemical	and Physi	cal proper	ties			
Туре	CAS LD ₅₀ and LC ₅₀		I	Boilin	Vapour pressure	Specific gravity	Flash point	Flammability	Dipole moment	Solvency power (Hansen parameter)	
Ketone	(67-64-1	ORAL (LD50): Acute: 5800 mg/kg [Rat]. 3000 mg/kg [Mouse]. 5340 mg/kg [Rabbit]. VAPOR (LC50): Acute: 50100 mg/m3 - 8 hours [Rat]. 44000 mg/m3 - 4 hours [Mouse].			56 °C	180 mm Hg	0.79 g/cm3	-20 °C	-20 °C Highly flammable		20.0
			-	•	Hea	lth guidan	ce				
_	mmediate) ffects	Chronic (Delayed) effe	ects	Target organ affected		Permissib			st aid	Medical Surveillance	
the respir and may c on the cer system, liv	of eyes and ratory tract cause effects ntral nervou ver, kidneys ointestinal	s with skin may cause derm	atitis. ffects	Eyes, sł respira system central nervou system	atory 1, I us 1	1000* 750****	750* 1000** 250*** 500****	support Swallow: N attention is	ly wash ly Respiratory	at the end of the Testing expired the end of the s Pulmonary Ful	d air for acetone at shift nction Tests (PFTs) or acetone at the
						vironmen	tal guidanc	e			
Incomp chem	patible nicals	Storage condition		Type of extinguis			Spi	ill control		Was	ste disposal
Oxidizers, acids		Fireproof Separate from strong oxidants	resista	amounts	nol- 1, water in 5, carbon	contained inert abs wash awa plenty of	ay into sewer water (extra	maining lique emove to saf Then wash personal	id in sand or e place. Do NOT	accordance	t be disposed of in with federal, state nvironmental ulations

Table 7: Data Sheet for Perchloroethylene (PCE)

	Toxic, Chemical and Physical properties													
Туре	CAS numbe	er	LD ₅₀ an	d LC ₅₀	Boiling point	Vapo		Spec grav		Flas poin	Hammahility	_	oole nent	Solvency power (Hansen parameter)
Chlorinated hydrocarbon	(127-18-	Acute: 26 (LD): Acu -4) [Rabbit]. 34200 mg	34200 mg/m 8 hours [Rat]. VAPOR (LC50): Acute: 5200 ppm 4 hours			⁰ C 14 1.62 None None None		e None	0 Debye		20.3			
					Healt	h guida								
Acute (Ir	nmediate	e) effects	Chron	onic (Delayed) effects		Target Permissib organ of expo affected STEL		-	1	First aid	Med		lical Surveillance	
The substance irritates the eyes, the skin and the respiratory tract. Swallowing the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis. The substance may cause effects on the central nervous system. Exposure at high levels may result in unconsciousness			contact cause substate on the	ted or prolonged It with skin may dermatitis. The Ince may have effect liver and kidneys. Ubstance is probably ogenic to humans	syste s liver, kidne	ratory m, ys, al ous	1003	****	100* 25**	***	Eye: Irrigate immediately Skin: Water flush promptly Breathing: Respira support Swallow: Medical attention immedia	,	(PCE) exhal shift Testin tetrac	chloroethylene content in end- ed air in prior to ng chloroethylene nt in blood in prior
					and env	ronme	ntal g	uidan	ice					
Incompati chemical		Storage con	lition	Type of fire extinguisher		Spill control					Waste disposal			
Strong oxidizers chemically-activ metals such as l beryllium & bar caustic soda; so hydroxide; pota	ve ithium, rium; dium	Separate from a See chemical da food and feedst Keep in the dar Ventilate along floor	ingers uffs k.	In case of fire in the surroundings: all extinguishing agents allowed	Ventilation. Collect leaking and spilled liquid in sealable containers as far as possible. Absorb remaining liquid in sand or inert absorbent and remove to safe place. Do NOT let this chemical enter the environment. (Extra personal protection: filter respirator for organic gases and vapours) Waste must be disposed of in accordance with federal, state and local environmental control regulations									

Table 8: Data Sheet for Trichloroethylene (TCE)

	Toxic, Chemical and Physical properties												
Туре	CAS number	LD ₅₀ and LC ₅₀	LD ₅₀ and LC ₅₀		Vapour pressure	Specific gravity	Flash point	Flammat	mability Dipole moment		Solvency power (Hansen parameter)		
Chlorinated hydrocarbon	1 1/9-111-61 1 3 3, 31		o, o	87ºC	58 mm Hg	1.47 g/cm 3	None	None	9	0.8 Debye	19.0		
				Health	guidance	<u> </u>	•						
Acute (Imme	diate) effects	Chronic (Delayed) effects		et organ fected	_	ssible limits exposure TWA	Fin	st aid	Medical Surveillance				
The substance eyes and the since some substance may on the central system, result respiratory fai Exposure coullowering of co	kin. e liquid may on into the risk of monitis. The reause effects nervous ing in lure d cause	Repeated or prolonged contact with skin may cause dermatitis. The substance may have effects on the central nervous system, resulting in loss of memory. The substance may have effects on the liver kidneys (see notes). This substance is probably carcinogenic to humans		ory system, er, kidneys		50* 100** 10****	Skin: V flush p Breath Respin suppo Swallo Medic attent	diately Water bromptly hing: ratory rt bw:	Testing Trichloroacetic acid in urine in end of the shift at end of workweek Testing Trichlorethanol in blood in end of the shift at end of workweek Testing Trichloroethylene in blood in end of the shift at end of workweek Testing Trichloroethylene in endexhaled air in end of the shift at end of workweek Pulmonary Function Tests Blood Plasma		at end of workweek thanol in blood in end of workweek bethylene in blood in end of workweek bethylene in end- d of the shift at end of		
					ronmenta	guidance							
Incompatible	chemicals	Storage condition	-	pe of fire inguisher						Waste disposal			
Strong caustics & alkalis; chemically-active metals (such as barium, lithium, sodium, magnesium, titanium & beryllium) Separate from metals See chemical dangers food and feedstuffs Keep in the dark. Ventilate along the floor				of fire in t ndings: all uishing age d	he seala rema remo respi	Ventilation. Collect leaking and spilled l sealable containers as far as possible. A remaining liquid in sand or inert absorb remove to safe place. (Extra personal prespirator for organic gases and vapour this chemical enter the environment			possible. Absorb nert absorbent and personal protection: filter and vapours). Do NOT let disposed of in accordance with federal, state and local environmen				

^{*} Permissible Limits of Exposure (PLE) prescribed by Indian Factories Act, 1948

^{**} Permissible Exposure Limits (PEL) prescribed by Occupational Safety and Health Administration (OSHA), USA

*** Recommended Exposure Limit (REL) prescribed by National Institute for Occupational Safety and Health (NIOSH), USA

**** Threshold Limit Value (TLV) recommended by American Conference of Governmental Industrial Hygienists (ACGIH, 2008), USA