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1. Introduction

1.1. HARRPA

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HARRPA (Hydrocarbon Resins, Rosin Resins and Pine Chemicals Producers Association) is located in Brussels. HARRPA is a CEFIC (European Chemical Industry Council) sector group.

HARRPA represents the Europe based, hydrocarbon resins, rosin resins and pine chemicals producers. HARRPA has strong world wide connections to associations, like PCA (Pine Chemicals Association) in the USA, ANIPIN (Associacão Nacional dos Industriais do Pinus) in South America, and with Asia-Pacific producers.

HARRPA represents the producers of resins based on natural raw materials (tall oil rosin and gum rosin) coming from pine trees, and resins derived from hydrocarbon feedstock from the petrochemical industry.

HARRPA's mission is to create a forum for industry discussion, to share best practices with all stake holders, and to represent our industry by interacting with the authorities.

1.2. Products applications

After processing, the raw materials become resins. Depending on the starting materials and the nature of the reaction, the resins can range in colour from water white to dark yellow/brown and from liquid to solid at ambient temperature.

These resins find numerous applications amongst others paints, coatings, adhesives, hot melt adhesives, sealants, printing inks, rubber, tyres, paper, road construction, and also chewing gum, beverages including wine, cosmetics, fragrances and perfumes, etc...

To find out more about HARRPA, please visit our website www.harrpa.org.

1.3. Scope of the brochure

This brochure is designed to share information about risks associated with the handling of solid resins. (For definitions see § 2.1.)

The goal of this brochure is:

- to identify and collate the best practices in the industry
- to share those practices with safety professionals and others
- to reduce the risks related to handling of solid resins

This document is written on behalf of HARRPA.

It is not intended to give answers to all safety aspects related to the handling of these solid resins. This document shares best practices and ideas from the companies that helped compile it.

It should be noted that circumstances may differ, and one should always study the local rules and legislation and take the appropriate measures related to that particular situation.

It is the responsibility of local management to make sure that all precautions are taken to safeguard people and the environment.

The information and advice in this brochure is given in good faith, but it is up to the user to check the suitability of equipment and operating procedures for their own particular situation. Neither HARRPA nor Cefic shall be liable for any loss, damage or injury which may arise from the use of these guidelines, whether caused by negligence or otherwise.

2. Products covered

2.1. Definitions

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In this brochure, the word **resin** is not limited to hydrocarbon resins, or rosin resins but includes: tall oil, wood, gum rosins and their derivatives.

The handling of these resins might create dusts. These dusts have certain explosion characteristics. Under adverse conditions explosions may occur.

Solid resins are delivered as prills, flakes, pellets, or in drums (cast solid). They are brittle materials prone to create dust during handling. These combustible dusts present a fire or explosion hazard when dispersed in air or other gaseous oxidiser. This may lead to violent explosions if ignited.

2.2. Clusters of rosin resins and rosin derivatives

- Cluster R1 Rosin disproportionated, dehydrogenated, hydrogenated, dimerized, decarboxylated rosin
- Cluster R2 Glycerol pentaerythritol esters from Cluster R1
- Cluster R3 Rosin treated with maleic/fumaric acid or maleic anhydride
- Cluster R4 Glycerol, pentaerythritol esters from Cluster R3 and their salts like Li, Na, K, Mg, Ca, Zn, including aliphatic amine salts and ammonium salts
- Cluster R5 Rosin, condensed with formaldehyde, phenol, alkyl phenols, aryl phenols and if applicable esterified with polyvalent alcohols and if applicable their metal salts and if applicable modified with hydrocarbon resins on basis C5-C10 and if applicable fatty acids (like tall oil fatty acids, soy bean oil, coconut oil, linseed oil, castor oil) and if applicable their esters
- Cluster R6 Rosin, treated with maleic/fumaric acid or maleic anhydride, condensed with formaldehyde, phenol, alkyl phenols, aryl phenols and if applicable esterified with polyvalent alcohols and if applicable their metal salts and if applicable modified with hydrocarbon resins on basis C5-C10 and if applicable fatty acids (like tall oil fatty acids, soy bean oil, coconut oil, linseed oil, castor oil) and if applicable their esters

Cluster R7 Rosin acids and resin acids condensed with formaldehyde

2.3. Clusters of hydrocarbon resins and derivatives

- Cluster H1 Aliphatic, alicyclic resins Aliphatic unsaturated (C5 dienes, C4-C6 olefins), aliphatic cyclic (DCPD, terpene resins)
- Cluster H2 Terpene resins α -pinene, β -pinene, α/β -pinene, d-limonene, camphene, dipentene / β -phellandrene
- Cluster H3 Aromatic containing resins Aliphatic/aromatic (pure monomers), aromatic (pure monomers), aromatic/cyclic, C9 aromatic, aliphatic/C9 aromatic
- Cluster H4 Hydrogenated resins Aliphatic, aliphatic/aromatic, aliphatic/aromatic/cyclic (Terpene) Aromatic: DCPD/indene/styrene/α-methyl styrene/vinyl toluene/isobutylene, α-methyl styrene/styrene, α-methyl styrene/vinyl toluene, styrene, others
- Cluster H5 Polycyclopentadiene resins
- Cluster H6 Coumarone-indene resins Coumarone indene, modified coumarone indene

2.4. Dust explosion characteristics

Typical range of measured dust characteristics of solid resins based on the fraction of $\leq 63 \mu m$. (All the explosion characteristics are given and should be measured in accordance with International Standard Methods).

Minimum Ignition Energy	≅ 1 - 3 mJ
Lower Explosion Limit (in air)	≅ 15 g/m³
K _{st} (explosion index)	≅ 200 - 315 bar.m.s ⁻¹
Electrical Resistivity	> 10 ¹⁴ Ohms.m
Dust Auto Ignition Temperature	≅ 360 °C
Maximum Explosion Pressure	≅ 8.8 bar
Hazard Class	= St -2 or St -3
Layer Ignition Temperature	= Not measurable (product melts)

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3. Hazards and precautionary measures

3.1. General remarks

Combustible dust when finely divided or suspended in air presents a fire or explosion hazard when ignited.

Minimize airborne dust and eliminate all ignition sources. Keep away from heat, hot surfaces, sparks, and flame. Control sources of static electricity at conveyors, ducts, piping, and mechanical and manual resin handling operations by installing and maintaining proper electrical bonding and grounding systems.

Establish good housekeeping practices. According to NFPA standard 654, dust layers 0.8 mm thick can be sufficient to warrant immediate cleaning of the area.

Do not use air hoses for cleaning. Minimize dry sweeping to avoid generation of dust clouds. Vacuum dust-accumulating surfaces and remove to a chemical disposal area. Vacuums with explosion-proof motors should be used.

Handle in accordance with good industrial hygiene and safety practices. These practices include avoiding unnecessary exposure to and removal of the material from eyes, skin and clothing. Wash thoroughly after handling.

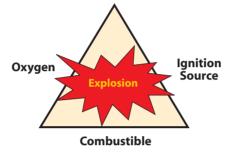
Use continuous suction at points of dust generation to capture and minimize the accumulation of dust clouds. Always use appropriate Personal Protective Equipment (PPE).

Train workers in the recognition and prevention of hazards associated with combustible dust in the plant.

Always read and understand the SDS (Safety Data Sheet) and label information before using the products.

3.2. Conditions leading to an explosion

Uncontrolled combustion may lead to an explosion and can take place provided three components are present simultaneously and in the right amount. The components are: combustible material (fuel), oxygen, source of ignition. This can be illustrated by the well known fire triangle.



For combustion/explosion, the fuel/oxygen ratio must be within a specific range. Also, the ignition energy must be sufficiently high to initiate combustion. Resins are flammable and the minimum ignition energy for resin dust is low. (See table §2.4.).

Prevention of fires and explosions is generally accomplished by removing one or better two sides of the fire triangle.

The recommended preventive measures are based on that principle.

3.3. Summary table

The table hereafter is a summary of the main hazards and related safety measures.

HAZARD	PRECAUTIONARY MEASURES
Ignitions sources	
• Flames:	> Working methods:
welding flame, cutting flame,	orders, procedures, work permit
combustion engine, cigarettes,	
Electric spark:	> Equipment selection and maintenance
Motors, switches, fixed and portable	ATEX regulation (area classification / adequate
instruments, cellular phone	equipment selection)
 Mechanic spark / hot spot: 	> Equipment selection and maintenance
Hammer, sandblasting, frictions,	ATEX regulation (area classification / adequate
	equipment selection)
Electrostatic spark:	> Grounding / bonding of conductive pieces,
Charges generated by friction, contact	selection of anti-static materials (PE bags, operator
(Pouring product, product flow, machinery)	clothes), hygrometry measurement and control.
Auto-ignition:	
Hot surfaces (steam line, machinery,),	> Equipment and facilities design, dust
self - heating due to product oxidation	accumulation prevention (design, housekeeping)

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Flammable material		
Flammable dust in cloud	 Minimize dust generation (design, equipment selection, oil/water pulverisation), avoid accumulation (dust collection system, housekeeping) 	
Oxygen		
	> Inerting	
Explosion		
	> Explosion mitigation: explosion vent,	
	explosion suppression, secondary explosion risk	
	prevention, explosion propagation prevention	
Other risk: Health risk		
Suspended dust presents respiratory risks	> Anti-dust mask	
for the personnel exposed to it		

3.4. Equipment and facilities design

A few simple rules should be observed when designing a plant:

- Avoid the creation of dust.
- Use where possible equipment, which does not induce the formation of dust clouds.
- Cleaning: only clean by suctioning in order not to disperse the dusts.
- Install dust collection equipment at the debagging station or at the product processing station.
- Do ensure that dust concentrations are outside the explosive range.
- Create an inert (or oxygen-depleted) atmosphere in the equipment presenting risks when the presence of a dust cloud is inevitable, bearing in mind the process involved.
- Protect the equipment using discharge devices such as (e.g.):
 - explosion hatches
 - safety panels / rupture discs
 - automatic expansion devices
- Connect the vents to a safe location outside of the plant.
- Ensure proper electrical grounding of all equipment.
- Choose mechanical and electrical equipment complying with local regulation and not susceptible to generating heat or sparks (e.g. power break) leading to the ignition of the dust.
- Forklift trucks: use of specific types.

3.4.1. Electrical equipment

This equipment must necessarily be adapted to the ATEX classification of the zone concerned, and be designed to be operated in a dust-laden atmosphere.

Please note: If the equipment is dismantled while still live, or if protective casings are opened, safety can no longer be guaranteed.

3.4.2. Explosion vent

The conditions of use (pressure) should be stipulated for all new equipment; bearing in mind the K_{st} value of resin powder, and the maximum pressure reached during the combustion of a dust cloud, devices for relieving the overpressure should be calculated correctly.

The following dimension guides are generally used in this regard:

- NFPA-68: Guide for venting deflagration (National Fire Protection Association USA)
- VDI 3673: Pressure relief of dust explosion (Verlag Düsseldorf Germany).



The vents should be ducted outside the plant using a duct with suitable diameter and sturdiness.

Exhaust pipes should not have any sharp bends or be too long.

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3.4.3. Explosion suppression

Another protection method consists of introducing into the enclosure, as soon as an unusual rise in pressure has been detected, a pressurised extinguishing agent using a pyrotechnical device.

3.4.4. Grounding/bounding

As a general rule: do remember that static electricity is always **present**. It is generated in varying degrees by the friction of the products against the walls of the equipment and pipes (pneumatic transport).

Sparks, resulting from a potential difference, can even appear between an operator moving closer to a piece of equipment and the equipment itself. The pulse energy in this specific case is of the order of 30 - 50 mJ.

The mere flow of solid particles down the slopes of a heap inside a silo causes a build-up of charge.

Therefore make sure that:

- Electrical continuity connections are present, and ducts and pipes connected to the various
 apparatus are earthed correctly.
- The foot-racks on metal platforms and the internal metal parts of the equipment are earthed.
- Pneumatic conveyor lines and sieves are earthed.
- Debagging hoppers and pipes connected to them are earthed.

Electrical continuity and equipotent bounding must be checked on a regular basis in accordance with the regulations in force in the country.

Regular cleaning of the metal floors also contributes to their proper electrical continuity.

Insulating materials which can impair proper electrical continuity must be banned.

Equipment that is liable to contain dust must not be allowed to act like a lightning rod.

Please note: personnel must wear safety shoes with conductive soles.

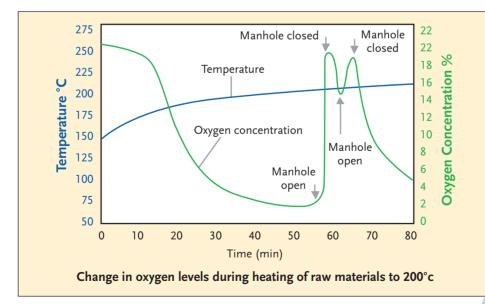
3.4.5. Inerting

Inerting is the process of converting from flammable or reactive conditions into a safe nonflammable state by the addition of an inert gas (usually nitrogen, but also carbon dioxide, or others can be used).

An inert gas such as nitrogen can be used to lower the oxygen concentration inside the vessel. According to experts opinion, the maximum oxygen content should be as low as possible and, as a rule of thumb, should not exceed 4 %, but should be checked in any case.

The inerting system must be designed to prevent air ingress when loading the vessel. When equipment has been purged with inert gas, particular attention must be paid to the risk of asphyxiation for personnel operating the plant or entering the enclosures concerned for maintenance purposes. In such cases, access permits must be required in addition to work permits.

Basic understanding of the chemistry of the manufacturing process and good engineering practices are necessary to determine the risks and the appropriate design of the process equipment. A study by T.J. Snee (T.J. Snee, 1985) shows that all steps in the production process should be considered. In this study T.J. Snee has shown that opening an inerted vessel increases the oxygen concentration dramatically during warming up of the content of the vessel (see graph hereafter). The oxygen concentration increases to 21 % in a short time. The presence of an ignition source or sufficient heat may have serious consequences.



ARRPA

Pure nitrogen or carbon dioxide, or a mixture of air and nitrogen may be used for purging purposes.

In the latter case, adjusting the mixture can be achieved in a simple way by using an oxygen analyser after mixing at a sufficient distance from the nitrogen inlet point.

Operating outside the flammability limits is under all circumstances essential and inerting can help to achieve this. Suppliers of inert gases can help with design and delivery of the right equipment and gas for the specific production process.

Explosive atmospheres are not only created by resin dusts, but may include dusts from other raw materials leading to completely different explosion conditions.

Apart from dust explosions, be aware of other explosive atmospheres created by gas or liquids.

When working with combustible or flammable materials, run risk analysis.

An article of G. Raykovitz, (Safety first for maximum benefits – Adhesive Technology , March 2001), describes the T.J. Snee study and the challenges created by relying on inerting alone.

3.4.6. Secondary explosion risk

A primary explosion can spread outside the zone in which it first occurred. An ensuing pressure wave can then scatter dust deposits into suspension (steel structures, pipes, equipment, etc.), and, through the energy released by the primary ignition, produce a secondary explosion.

Example: ignition of coal-gas in a mine causing a secondary dust explosion.

4. Product handling

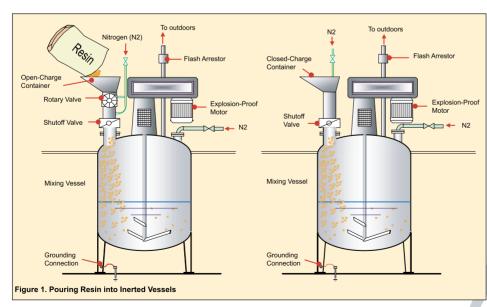
4.1. Pouring product

4.1.1. Static charge generated when pouring resins

In the past, recognised institutes completed tests to determine the magnitude of static electricity generated when emptying bags of resins. Plain, plastic-lined, and plastic-coated kraft paper bags, as well as polyethylene plastic bags, were used in the tests. Although the testing was performed primarily for resins, finely divided polypropylene flake and sand were included for comparison. All were found to generate similar amounts of static electricity. Consequently, we conclude that the potential to generate static electricity exists when dumping resins from any of these bags.

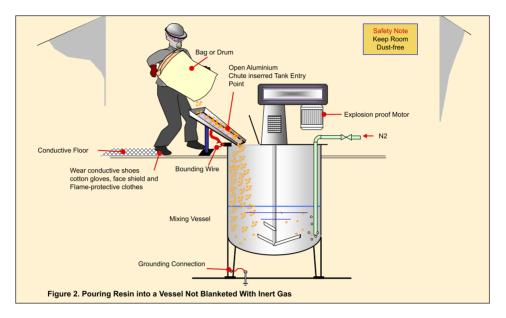
The static electricity generated (on the operator, bag, and equipment) while dumping the bags was, in several cases, found to have the potential to ignite flammable vapours, and some combustible dusts exhibiting low ignition energies. The information that follows will help to prevent electrostatic charge build up.

Since the build up of electrostatic electricity depends on many factors, (floor and operator shoes conductivity, hygrometry during operations, presence of dust, etc.) there is no guarantee our guidelines will prevent static charge build up. However, implementation of the following guidelines should minimize the potential for fire when pouring resins.



Resin dust explosion risks

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4.1.2. Suggestion for reducing dust when pouring bags

The following practice for minimizing dust has been found effective. However, it should not preclude the use of a metal chute or implementation of the aforementioned safety measures.



- Step 1: Make a 10 to 15 cm slit at top rear of bag. (The slit will function as a vent and minimize turbulence to provide free - less dusty - flow.)
- Step 2: Suspend end of bag over chute. Reach under bag and cut open to make a flap as indicated by arrows.





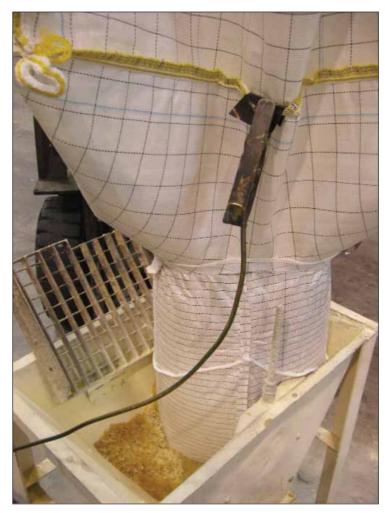
Step 3: Empty bag slowly.

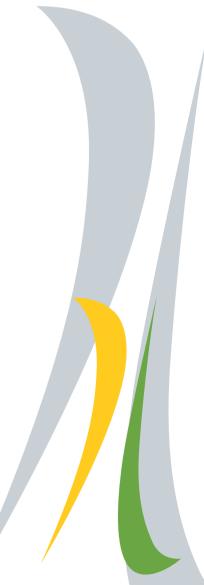
4.1.3. Use of flexible intermediate bulk containers (FIBCs)

When using FIBCs extra precautions should be taken. When emptying these bags large quantities of resin pour out. The warning label on the FIBC should be read before handling.

This flow of material can generate large electrostatic charge build-up, depending on flow rate, type of material, type of FIBC and equipment used. In general the discharge speed should not exceed the maximum of 2 kg/s. Consult the proper experts for further detail on maximum discharge speed and proper discharge equipment.

Avoid sparks by grounding all equipment and the FIBC (grounding cable must be connected to the bag ground connection).





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5. Working methods

Four main points must be observed:

• A clean and tidy plant: elimination of all secondary explosion risks.

This must be ensured to prevent all risks of dust becoming airborne. Regular cleaning of the plant must be carried out with a vacuum cleaner complying with ATEX regulations based on the classification of the plant's zone. As a general rule, ensure good house keeping. The pictures below illustrate exemples of dust accumulation which may lead to explosion risks.







• Elimination of hot spots:

• Open flames and welding:

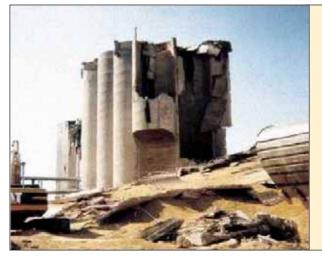
Operations generating flames (welding / cutting / « de-solidification » using a welding torch, etc.) should not, where possible, be carried out in a plant in operation. In all cases, a work permit and a fire permit must be issued beforehand.

• Defective equipment or electrical short circuit:

Repair defective equipment immediately, or replace if necessary.

• Mechanical friction and impacts:

- i. Tackifying powders tend to produce crusting which can induce stress in mechanical parts, and, as a result, overheating.
- ii. Ban the use of impact tools such as hammer, chisel, impact wrench, etc., not suitable for the work atmosphere.



Illustrative example:

Explosion at Blaye (France) – 1997

Explosion of wheat dust in a confined environment, purportedly due to overheating in a dust-extraction fan

11 fatalities – 1 injured

• Maintenance of equipment

Have preventive maintenance in place. Be aware of inerted equipment, and spreading of dust by opening of equipment. Use the right explosion proof tools.

- Training of personnel:
 - Properties and hazards of the products
 - Operating instructions
 - Failure alerts
 - Emergency plans

Notice:

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Provide instructions for contractors and subcontractors on safety rules and work permits.

6. Health risks

Apart from the ignition/explosion hazards inherent to dusts, they also present genuine risks to human health through inhalation.

It is important to recall that particles with a size ranging from 5 μ m to 30 μ m are stopped by the upper respiratory tract (nose, mouth, pharynx). If the concentration of these particles is too high, they line the mucous membranes and cause real discomfort. This discomfort can be exacerbated by the tackifying nature of the product.

The smallest particles (1 μ m and under) enter the lower respiratory system: from the trachea right down to the air sacs where they will be destroyed by macrophage. Any presence of powder exceeding the macrophage processing capacity will cause a build-up and, as a result, inflammation.

It is therefore important to protect the people working in an atmosphere laden with suspended fine powder in order to eliminate health risks.

Personal protective equipment:

Non-valve dust respirators and valve dust respirators (for high dust concentrations) are typical personal protective equipment, which should not replace collective protective equipment.

They can be worn during short periods of time and in particularly dust-laden conditions.

It should be noted that standard EN149 dated 2001 provides for 3 effectiveness classes (FFP1...FFP3) measured on particles of 0.01 to 1 $\mu m.$

The appropriate class should be chosen on the basis of dust levels:

FFP1: stops 77% of particles FFP2: stops 92% of particles FFP3: stops 98% of particles.

Operators should wear breathing masks marked EN149:2001.

Summary of the Personal Protection Equipment (PPE) which might be used

The following table is only given as a recommendation.

The choice of the PPE must be result of a formal analysis including risks assessment, hazard assessment \ldots

	HARRPA	PPE	
	'Normal work conditions Low Dusts concentration	'Unsual work conditions (*) High Dusts concentration	
Helmet	~	~	
Gloves	~	~	
'Working Suit anti-static	~	~	
' Safety Shoes (with conductive soles)	~	~	3
Safety Glasses	~		
'Dusts-proof Goggles		✓	40
' Special non-wooven Suit Dusts proof (anti-static)		~	
Hair cap		✓	
Anti-Dusts Mask FFP3		~	

(*) ie.: Special maintenance works

or less in diameter (i.e., material passing through a U No.40 standard sieve), that presents a fire or explos hazard when dispersed and ignited in air or ot gaseous oxidizer.DustDustAny finely divided solid material, 500 µm (425 µm U.S.) or less in diameter (i.e., material passing throug U.S. No.40 standard sieve).ECElectrical ConductivityThe reciprocal of resistivity expressed as Pico Siem per meter. picoS = 1 x 10 ⁻¹² dP/dtMaximum Rate of Explosion PressureAn indicator of explosion severity.IGInert GasA non-flammable, non-reactive gas that reduces the a lable oxygen level in air to the point where a flamma mixture is no longer possible.KatExplosion Indices (St stands for Staub)The maximum value for the material specific const "K _{mm} " that describes the rate of pressure rise, dP/dt a function of vessel volume.LEL/MECLower Explosion Limit Minimum Explosion ConcentrationThe lowest concentration of a combustible dust in expressed in grams per cubic meter, which will propa te a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant).LITLayer Ignition Temperature (smoulder temperature)The smoulder temperature of a heated, free-standing su as the lowest temperature of a heated, free-standing su					
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Image: second	Dust		U.S.) or less in dia U.S. No.40 standa	meter (i.e., materia rd sieve).	al passing through a
Explosion PressureA non-flammable, non-reactive gas that reduces the a lable oxygen level in air to the point where a flamma mixture is no longer possible.K#Explosion Indices (St stands for Staub)The maximum value for the material specific const "Kmax" that describes the rate of pressure rise, dP/dt a function of vessel volume.Hazard ClassKst function of vessel volume.Hazard ClassKst 	EC	Electrical Conductivity	The reciprocal of resistivity expressed as Pico Siemens per meter. $picoS = 1 \times 10^{-12}$		
Iable oxygen level in air to the point where a flammar mixture is no longer possible.KmExplosion Indices (St stands for Staub)The maximum value for the material specific const "Kmmx" that describes the rate of pressure rise, dP/dt a function of vessel volume.Hazard ClassKst bar.m.s'Pmmx (bar)St-120010St-2201-30010St-3>30012LEL/MECLower Explosion Limit Minimum Explosion ConcentrationThe lowest concentration of a combustible dust in expressed in grams per cubic meter, which will propa- te a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant).LITLayer Ignition Temperature (smoulder temperature)The smoulder temperature of a heated, free-standing su ce which is capable of igniting a 5 mm thick dust layerMIEMinimum Ignition EnergyLowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust	dP/dt		An indicator of explosion severity.		
KatExplosion Indices (St stands for Staub)The maximum value for the material specific const "Kmax" that describes the rate of pressure rise, dP/dt a function of vessel volume.Hazard ClassKst function of vessel volume.ItelLower Explosion Limit Minimum Explosion ConcentrationThe lowest concentration of a combustible dust in expressed in grams per cubic meter, which will propa te a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant).LITLayer Ignition Temperature (smoulder temperature)The smoulder temperature describes the flammab behaviour of a flat dust layer on a hot surface. It is defin as the lowest temperature of a heated, free-standing su ce which is capable of igniting a 5 mm thick dust layerMIEMinimum Ignition EnergyLowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust	IG	Inert Gas	lable oxygen level	in air to the point	
St-3 >300 12 LEL/MEC Lower Explosion Limit Minimum Explosion Concentration The lowest concentration of a combustible dust in expressed in grams per cubic meter, which will propa- te a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant). LIT Layer Ignition Temperature (smoulder temperature) The smoulder temperature describes the flammab behaviour of a flat dust layer on a hot surface. It is defin as the lowest temperature of a heated, free-standing su ce which is capable of igniting a 5 mm thick dust layer MIE Minimum Ignition Energy Lowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust	Kst		"K _{max} " that describ function of vessel Hazard Class	es the rate of press volume. K _{st} bar.m.s ⁻¹	sure rise, dP/dt as a Pmax (bar)
LEL/MEC Lower Explosion Limit The lowest concentration of a combustible dust in expressed in grams per cubic meter, which will propative a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant). LIT Layer Ignition The smoulder temperature describes the flammab behaviour of a flat dust layer on a hot surface. It is defined (smoulder temperature) MIE Minimum Ignition Lowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust			St-2	201-300	10
Minimum Explosion Concentrationexpressed in grams per cubic meter, which will propa te a flame. (Specific to the PSD of the tested sample a is not an intrinsic material constant).LITLayer Ignition Temperature (smoulder temperature)The smoulder temperature describes the flammab behaviour of a flat dust layer on a hot surface. It is define as the lowest temperature of a heated, free-standing su ce which is capable of igniting a 5 mm thick dust layerMIEMinimum Ignition EnergyLowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust			St-3	>300	12
Temperature (smoulder temperature)behaviour of a flat dust layer on a hot surface. It is define as the lowest temperature of a heated, free-standing su ce which is capable of igniting a 5 mm thick dust layerMIEMinimum Ignition EnergyLowest energy required, which upon discharge is su cient to ignite the most ignitable mixture of air and dust		Minimum Explosion Concentration			
Energy cient to ignite the most ignitable mixture of air and de		Temperature (smoulder temperature)	behaviour of a flat dust layer on a hot surface. It is defined as the lowest temperature of a heated, free-standing surfa- ce which is capable of igniting a 5 mm thick dust layer.		
	MIE	-	cient to ignite the	most ignitable mix	-

7. Definitions (glossary)

MIT	Minimum Ignition	The minimum temperature at which a dust cloud can be
	Temperature	ignited.
MOC	Minimum Oxygen	The lowest molar (or volume) gas-phase concentration
	Concentration	of oxidant (usually oxygen) at which a specified fuel
		(dust) can propagate a flame. (For most organic dusts
		<12 vol % O ₂ , paraformaldehyde <4 vol %)
MSIE	Minimum Spark	Lowest electrical energy stored in a capacitor, which
	Ignition Energy	upon discharge is sufficient to ignite the most ignitable
		mixture of air and dust, under specific test conditions.
PSD	Particle Size	Explosive properties are usually measured at a dust frac-
	Distribution	tion of ≤63 µm.
SA	Surface area	
	Zone 20	A place in which an explosive atmosphere in the form of
		a cloud of combustible dust in air is present continuous-
		ly, or for long periods or frequency.
	Zone 21	A place in which an explosive atmosphere in the form of
		a cloud of combustible dust in air is likely to occur in nor-
		mal operation occasionally.
	Zone 22	A place in which an explosive atmosphere in the form of
		a cloud of combustible dust in air is not likely to occur in
		normal operation but, if it does occur, will persist for a
		short period only.

8. References

Documentation on risks of flammable mixtures, measures and engineering practices can be found on the HSE UK website where a brochure titled "Guidance and Codes of Practice relating to inerting" is published.

HS(G)140 Safe use and handling of flammable liquids, HSE, 1996. Paragraph 36 gives guidance on the transfer of flammable liquids using gas pressure.

HS(G)158 Flame arresters : preventing the spread of fires and explosions in equipment that contains flammable gases and vapors, HSE, 1996.

Paragraph 35 gives guidance on the use of inert gas blanket within vessels as alternative to flame arresters, where explosions may occur in emergency vents.

HS(G)103 Safe handling of combustible dusts : precautions against explosions, HSE, 1994. Paragraphs 35 to 36 give guidance on the use inerting to prevent dust explosions. GS5 Entry into confined spaces, HSE, 1994. Gives guidance on vessel entry where inert gases are used.

BS 5908 : 1990 Code of Practice For Fire Precautions in the Chemical and Allied Industries, British Standards Institution. General. Section 10, Paragraph 58.5 provides guidance on the flammability of powders and measures to protect against ignition including inerting.

F.P. Lees, 'Loss Prevention in the Process Industries: Hazard Identification, Assessment and Control', Second Edition, 1996.

G. Raykovitz, Safety first for maximum benefit, Adhesive Technology, March 2001.

T.J. Snee, An instrument for measuring the concentration of combustible airborne material, Ann. occup. Hyg. Vol. 29, No. 1, pp. 81-90, 1985, © 1985 British Occupational Hygiene Society, Published by Oxford University Press.

ATEX Directive 1999/92/EC.

U.S. Chemical Safety and Hazard Investigation Board, Investigation Report, Combustible Dust Hazard Study. Report No. 2006-H-1 November 2006 HARRPA

NFPA 654 Standard for the prevention of fire and dust explosions from the manufacturing, processing, and handling of combustible particulate solids 2006 edition.

L138 HSE UK Dangerous substances and explosive atmospheres regulation 2002.

Guidelines for safe manufacture of resins with particular reference to fire and explosion, European Resin Manufacturers Association, 1995, ISBN 0903809 17 6.





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