

# **PINE ROSIN FLAKES**

#### **SAFETY DATA SHEET**

This Safety Data Sheet contains information concerning the potential risks to those involved in handling, transporting and working with the material, as well as describing potential risks to the consumer and the environment. This information must be made available to those who may come into contact with the material or are responsible for the use of the material.

SECTION 1: Identification of the substance/mixture and of the company/undertaking

#### 1.1 Product identifier

Product name: ROSIN

Other names: Colophony, Kolophonium, Pine Resin, Gum Rosin, Wood Rosin

EC number: 232-475-7

CAS number (EC inventory): 8050-09-7

REACH Registration Number: 01-2119480418-32-xxxx

1.2 Relevant identified uses of the substance or mixture and uses advised against

Friction Aid, Grip Aid, Solder Flux, Electrical Insulation, Non-Draining Compound, Coating, Emulsion, Adhesives, Depilatory Wax, Clay Pigeon, Polish, Sealants, Dental Cement, Pigment Manufacture

1.3 Details of the supplier of the safety data sheet

LiveMoor
Unit 9A Forresters Business Park
Estover Close
Plymouth
Devon
PL6 7PL

SECTION 2: Hazards identification

2.1 Classification of the substance

According to 1272/2008/EC Regulation

Skin sensitiser Category 1

2.2 Label elements

**Pictogram** 



Signal word: Warning



# Hazard Statement/Precautionary Statements

H317 – May cause an allergic skin reaction

P210 – Keep away from heat/sparks/open flames/hot surfaces. No smoking

**P261** – Avoid breathing dust/fumes

P280 – Wear protective gloves/protective clothing

P302 + P352 - If on skin: Wash with plenty of soap and water

P333 + P313 – If skin irritation or rash occurs: Get medical advice/attention

P363 – Wash contaminated clothing before reuse

P501 – Dispose contents/container according to the end user disposal procedure

### According to 67/548/EC Regulation



Xi - Irritant Risk Phrase

# Safety Phrases

S 24 - Avoid contact with skin

\$ 37 – Wear suitable gloves

# SECTION 3: Composition/information on ingredients

In accordance with Annex II of Regulation (EC) No. 1907/2006 (point3), the product contains:

Identification	Chemical name/classification			Concentration
CAS: 8050-09-7	Rosin	ATP CLP00		
EC: 232-475-7	Directive 67/548/EC	Xi: R43	×	100%
Index: 650-015-7	Regulation 1272/2008	Skin sens1: H317-Warning	<b>(1)</b>	

# SECTION 4: First aid measures

# 4.1 Description of first aid measures

EYE CONTACT: Immediately flood eyes with plenty of low-pressure water for at least 10 minutes, holding eye open. Remove contact lenses. Obtain medical assistance if redness develops or persists.

SKIN CONTACT: Wash skin thoroughly with soap and water. Obtain medical help if redness develops.

INGESTION: Wash mouth out with water. Do not induce vomiting. Obtain Medical attention if a large amount has been swallowed.

INHALATION: Remove from exposure, keep warm and at rest. Obtain medical attention if breathing difficulty occurs.

# 4.2 Most important symptoms and effects, both acute and delayed:

Acute and delayed effects are indicated in section 2 and 11.



**SECTION 5: Firefighting Measures** 

5.1 Extinguishing media: Carbon dioxide, dry chemical or water fog Unsuitable extinguishing media: Water

5.2 Special hazards arising from the substance or mixture:

Protective equipment: Self contained breathing apparatus needed for fires in enclosed areas.

Other information: Rosin dust is a severe explosion hazard in air, avoid dispersing dust in air Minimum exposable concentration: 15g/m<sup>3</sup>. Risk of spark ignition prevented by reducing oxygen concentration below 17% by dilution with carbon dioxide.

5.3 Advice for fire fighters - Do Not Use Water as extinguishing media. No special firefighting equipment required.

SECTION 6: Accidental Release Measures

- 6.1 Personal precautions: Avoid breathing dust. Wear dust respirator, gloves, chemical goggles and overalls.
- 6.2 Environmental precautions: Avoid dispersion of the product
- 6.3 Methods and materials for containment and clearing up: Sweep up or otherwise recover the product and remove to a safe environment

Additional Advice: Avoid propagating more dust than is absolutely unavoidable.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

Person Protection: Wear protective clothing and dust mask

Dust: Avoid processes that generate excessive dust

7.2 Conditions for safe storage

Suitable Storage Container: Galvanised iron drums or paper sacks internally coated with silicone or with a polypropylene liner.

Handling/Storage Precautions: Explosion proof ventilation adequate to meet any dust conditions at room temperature. Store in cool, well ventilated conditions.

Explosion proof electrical services should be used where dusty conditions prevail.

Prevent static sparks. Product can spontaneously heat during storage.

Exercise care: Avoid placing the material next to, or inContact with, oxidising agents.

7.3 Specific end use(s)

Except for the instructions alre4ady specified it is not necessary to provide special recommendation regarding the uses of this product.



### SECTION 8. Exposure controls/personal protection

#### 8.1 Control parameters - No data available

Substances whose occupational exposure limits have been monitored in the work environment Nuisance dust: Inhalable dust 10mg/m³ Respirable dust: 4mg/m³ DNEL workers

Acute/short-term exposure – systemic effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available Acute/short-term exposure – local effect

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Long-term exposure – systemic effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available

Long-term exposure – local effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available General population

Acute/short-term exposure – systemic effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available Oral DN(M)EL: No-threshold effect and/or no dose-response information available Acute/short-term exposure – local effect

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available

Long-term exposure – systemic effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available Oral DN(M)EL: No-threshold effect and/or no dose-response information available Long-term exposure – local effects

Dermal DN(M)EL: No-threshold effect and/or no dose-response information available Inhalation DN(M)EL: No-threshold effect and/or no dose-response information available Components to control with biological restricted values:

**PNEC** 

Freshwater: 0,0016mg/L

Freshwater sediments: 0,007mg/kg sediment dw

Marine water: 0,00016 mg/L

Marine sediments: 0,0007mg/kg sediment dw

Oral: No bioaccumulation potential

STP: 1000 mg/L

Soil: 0,0045 mg/kg soil dw



### 8.2 Exposure controls

Appropriated technical control measures: Dust extraction pointed to local of formation

Personal protective equipment

Respiratory protection: Wear respiratory protection against organic vapours

Hand protection: Wear appropriate gloves to prevent skin exposure Eye/face protection: Wear chemical splash goggles conforming to EN 166 Skin protection: Wear appropriate clothing to prevent skin exposure

Environmental exposure control

Organisational measure: Apply general preventative measures in chemical safe handling

# SECTION 9: Physical and chemical properties

#### 9.1 Information on basic physical and chemical properties

Appearance: Water-white to dark brown vitreous solid

Odour: Characteristic (slight turpentine odour

Odour threshold: No data available

pH: Not applicable Melting point: 65-90°C Flashpoint: >205°C

Auto ignition temperature: >390°C

Explosive Properties: Severe dust explosion hazard in air

Oxidising Properties: None Vapour Pressure: < 0.1 hPA Relative Density at 15°C: -1.08 Solubility – Water: Insoluble

Solubility – Solvent: Aromatic hydrocarbons, chlorinated hydrocarbons, alcohols, ketones

Other properties: Unsaponifiable matter 8% max

# SECTION 10: Stability and Reactivity

10.1 Reactivity: No hazardous reactions are expected

# 10.2 Chemical stability

Stability: Substance is stable at normal temperature and pressure.

Oxidation: Substance is oxidised by atmospheric oxygen

- 10.3 Possibility of hazardous reactions: Not if stored correctly
- 10.4 Conditions to avoid: Generation of dust in air
- 10.5 Incompatible materials: Oxidising agents
- 10.6 Hazardous decomposition products: Thermal oxidation may result in formation of formaldehyde and carbon monoxide



# SECTION 11: Toxicological information

### 11.1 Information on toxicological effects

**Oral Acute toxicity** 

Method: OECD Guideline 420 (Acute Oral Toxicity)

Species: rat (male and female)

Route of administration: oral, gavage

Dose: Single dose

Conclusion: LD50> 2000mg/kg similar substance

The classification criteria for oral acute toxicity or specific target organ toxicity by acute exposure were

not met, according to GHS, CLP and 67/548/EC. Analysis based on acquired data.

Inhalation: Based on available data, the classification was not met.

Dermal acute toxicity

Method: OECD Guideline 402 (Acute Dermal Toxicity)/EU Method B.3 (Acute Dermal Toxicity)

Species: rabbit (male and female)
Route of administration: dermal, topic

Doses: 2000mg/kg bw

Period of exposure: 24 hours

Conclusion: LD50>2000mg/kg (rabbit, male/female), similar substance. Classification criteria for dermal acute toxicity, irritation or skin corrosion of for single target organ toxicity by acute exposure, based on GHS, CLP and 67/548/EC, were met. Analysis based on acquired data for similar substance.

Irritation and skin corrosion: Classification not, determined due to lack of data Method: OECD Guideline 404 (acute Dermal Irritation/Corrosion)/Eu Method B.4

Species: rabbit

Route of administration: dermal, semi occlusive

Doses: 0.5g

Period of exposure: 4hrs

Conclusion: during tests no edema was observed. Classification criteria for dermal acute toxicity, irritation or skin corrosion or for single target organ toxicity by acute exposure, based on GHS, CLP and

67/548/EC, were not met. Analysis based on acquired data.

Eye irritation and damage

Method: OECD Guideline 405 (Acute Eye Irritation/Corrosion), equivalent method to

Species: rabbit

Route of administration cornea, topic

Doses: 100mg/eye/specimen
Period of exposure: single dose

Conclusion: slight reversable redness of the conjunctivae was observed on some animals. Classification criteria for eye irritation, based on GHS, CLP and 67/548/EC, were not met. Analysis

based on acquired data.

Skin sensitisation

Method: OECD Guideline 428 (LLNA), method equivalent to

Species: rat (female)

Conclusion: Based on study results, rosin is not a moderate or strong skin sensitiser to rats. Rosin did not show any potential to acts as skin sensitiser. The classification criteria for skin sensitising, based on GHS, CLP ans 67/548/EC, were not met. Analysis based on acquired data. However, Annex I in



Direction 67/548/EEC and table 3,1 in CLP Regulation present rosin in R43 – May cause sensitisation in contact with skin and H317: May cause skin allergic response, respectively.

Sensitising effects by inhalation: Based on data available, the classification requirement was not met.

# Mutagenicity

Method: OECD Guideline 471 (Bacterial Reverse Mutation Assay); EU Method B.13/14(Mutagenicity –

Reverse Mutation Test Using Bacteria)

Conclusion: Absence of genotoxic and mutagenic effects. Classification criteria were not met.

Analysis based on acquired data.

Carcinogenicity - Based on the data available, the classification was not met

### Toxicity for reproduction

Method: OECD Guideline 421 (Reproduction/Development Toxicity Screening Test)

Conclusion: NOAEL reproductive – 3000ppm (nominal) (male and female) Not determined on FI generation. Criteria were not met and the product is not classified for the Development or Reproductive Toxicity according to GHS, CLP and 67/548/EC. Analysis based on acquired data.

Long exposure effects: No reliable data available

STOT – single exposure: Based on available data, the classification was not met. STOT – repeated exposure: Based on available data, the classification was not met.

Respiratory risks: Based on available data, the classification was not met.

# SECTION 12: Ecological information

#### 12.1 Toxicity

Short term toxicity

Fish: 96h,LL50 < 10mg/L (nominal, based on mortality)

Brachydanio rerio, OECD 203

48, NOELr = 750mg/L (nominal, based on mortality) 48h, EL50 – 911mg/L (nominal, based on mortality)

Aquatic invertebrates: 48h, EL50 – 911mg/L (nomina Daphnia magna, OECD 202

72h, NOELr ≥ 1000mg/L (nominal, similar substance, based on growth

rate and biomass)

Algae and aquatic plants: 72h, LL50 = 1000mg/L (nominal, similar substance, based on growth

rate and biomass) Selenastrum capriconutum, OECD 201

Other organisms: Not determined

Long-term toxicity

Fish: Not determined Aquatic invertebrates: Not determined Algae and aquatic plants: Not determined Other organisms: Not determined

# 12.2 Persistence and degradability

Abiotic degradability: Not determined

Photochemical and physico-

Chemical elimination: Not determined

Biodegradation: Substance is readily biodegradable

70% degradation after 28 days



### OECD Guidelines 301 D (Ready Biodegradability: Closed Bottle Test)

# 12.3 Bioaccumulative potential

Partition coefficient n-octanol/water

Log (Kow): refer to section 9

Bioconcentration factor (BCF): 56,23 L/kg wet/wt (QSAR, regression-based m.)

Bioaccumulation factor (BAF): 694 000 (Arnot-Gobas upper trophic m.)

12.4 Soil Mobility

Partition coefficient soil/water (Koc) log Koc = 3,7289 (QSAR, estimated from log Kow)

12.5 PBT evaluation results: Product does not fulfil PBT or vPvB criteria

12.6 Other adverse effects/additional information: N/A

**SECTION 13: Disposal Considerations** 

### 13.1 Waste treatment methods

Recommendation: Dispose of waste to a licensed disposal site in accordance with the requirements of the local Waste Disposal Authority. Do not allow into drains, sewers and water courses.

Contaminated packaging: Used containers should be recycled after dust removal.

**SECTION 14: Transport Information** 

Not classified in accordance with ADR/RID, IMDG and IATA

SECTION 15: Regulatory Information

15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture

Classification acc. to:

Regulation (CE) n. 1005/2009: Not applicable

Regulation (CE) n. 1907/2006

- Annex XVII Not restricted substance:

Directive n. 2003/53/EC: Not applicable

Directive n. 2003/105/CE: Substance not classified in the scope in the directive

Regulation (CE) n. 850/2004: Not applicable

The German Federal Water

Management Act: Classe WGK 1



### SECTION 16: Other Information

Abbreviations and acronyms:

ADR: International Carriage of Goods by Road IMDG: International Maritime Dangerous Goods IATA: International Air Transport Association ICAO: International Civil Aviation Organisation

**BCF**: Bioconcentration Factor

LD50: Lethal Dose 50

LC50: Lethal Concentration 50

**KOC: Partition Coefficient of Organic Carbon** 

OECD: Organisation for Economic Cooperation Development

NOEL: No Observed effect level STOT: Specific Target Organ Toxicity

QSR: Quantitive Structure Activity Relationship

LLNA: Local Lymph Node Assay

The information in this safety data sheet is based on the properties of the material known to LiveMoor at the time the data sheet was issued. The safety data sheet is intended to provide information for a health and safety assessment of the material and the circumstances under which it is packaged, stored or applied in the workplace. This document is not intended for quality assurance purposes.