

# NS-5111-G

## Syn-Tech Ltd.

Version No: 1.1 Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

## **SECTION 1 Identification**

#### Product Identifier

| Product name                  | NS-5111-G     |
|-------------------------------|---------------|
| Synonyms                      | Not Available |
| Other means of identification | Not Available |

#### Recommended use of the chemical and restrictions on use

Relevant identified uses Lubricant

#### Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

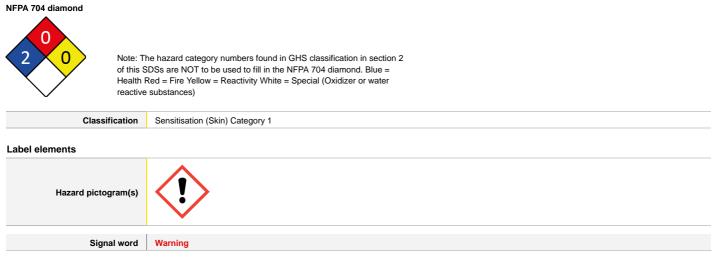
| Registered company name                           | Syn-Tech Ltd.   | Syn-Tech Ltd.                                |
|---|---|--|
| Address   | 1550 W Fullerton Ave, Unit F Illinois 60101 United States | 1550 W. Fullerton Ave Illinois United States |
| Telephone   | e 630-628-7290 630-628-7290                               |  |
| Fax   | x Not Available Not Available                             |  |
| Website www.syn-techlube.com www.syn-techlube.com |   | www.syn-techlube.com                         |
| Email   | msds@syn-techlube.com                                     | msds@syn-techlube.com                        |

#### Emergency phone number

| 5. 71                             |               |
|-----------------------------------|---------------|
| Association / Organisation        | Not Available |
| Emergency telephone<br>numbers    | Not Available |
| Other emergency telephone numbers | Not Available |

#### SECTION 2 Hazard(s) identification

# Classification of the substance or mixture



Chemwatch Hazard Alert Code: 2

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| Version | No <sup>.</sup> | 1.1 |
|---------|-----------------|-----|
| 101011  | 110.            |     |

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H317 May cause an allergic skin reaction.

#### Hazard(s) not otherwise classified

Not Applicable

### Precautionary statement(s) Prevention

| P280 | Wear protective gloves and protective clothing.                      |  |
|------|--|--|
| P261 | Avoid breathing dust/fumes.  |  |
| P272 | Contaminated work clothing must not be allowed out of the workplace. |  |

#### Precautionary statement(s) Response

| P302+P352 IF ON SKIN: Wash with plenty of water and soap. |  |
|---|--|
| P333+P313   | If skin irritation or rash occurs: Get medical advice/attention. |
| P362+P364   | Take off contaminated clothing and wash it before reuse.         |

#### Precautionary statement(s) Storage

Not Applicable

#### Precautionary statement(s) Disposal

| P501 | Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation. |
|------|--|
|      |  |

Not Applicable

## **SECTION 3 Composition / information on ingredients**

### Substances

See section below for composition of Mixtures

#### Mixtures

| CAS No      | %[weight] | Name  |  |
|-------------|-----------|---|--|
| 94270-86-7  | 0.3       | N-alkylated benzotriazole                                     |  |
| 125643-61-0 | 1         | C7-9 branched alkyl-3.5-di-tert-butyl-4-hydroxyhydrocinnamate |  |

#### **SECTION 4 First-aid measures**

#### Description of first aid measures

| Eye Contact  | Generally not applicable.   |  |
|--------------|---|--|
| Skin Contact | If skin contact occurs: <ul> <li>Immediately remove all contaminated clothing, including footwear.</li> <li>Flush skin and hair with running water (and soap if available).</li> <li>Seek medical attention in event of irritation.</li> <li>Generally not applicable.</li> </ul> |  |
| Inhalation   | <ul> <li>Generally not applicable.</li> </ul>   |  |
| Ingestion    | ► Generally not applicable.   |  |

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

See Section 11

Indication of any immediate medical attention and special treatment needed Treat symptomatically.

## **SECTION 5 Fire-fighting measures**

#### Extinguishing media

- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only.

#### Special hazards arising from the substrate or mixture

#### Fire Incompatibility

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result

#### Special protective equipment and precautions for fire-fighters

Fire Fighting

Alert Fire Brigade and tell them location and nature of hazard.
 Wear breathing apparatus plus protective gloves.

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|                 |                     | NS-5111-G                         |  | Print Date: 08/19/2022 |
|                 |                     |                                   |  |                        |
|                 | Provent by any mean | ns available, spillage from enter |  |                        |
|                 |                     | is available, spillage from enter |  |                        |

- DO NOT approach containers suspected to be hot.
  - DO NOT approach containers suspected to be not.
     Cool fire exposed containers with water spray from a protected location.
     If safe to do so, remove containers from path of fire.
     Equipment should be thoroughly decontaminated after use.
     Slight hazard when exposed to heat, flame and oxidisers.

|                       | Sign hazard when exposed to heat, name and oxidisers.  |
|-----------------------|--|
| Fire/Explosion Hazard | Combustible. Will burn if ignited.<br>Combustion products include:<br>carbon monoxide (CO)<br>carbon dioxide (CO2)<br>other pyrolysis products typical of burning organic material.<br>May emit corrosive fumes.<br>Articles and manufactured articles may constitute a fire hazard where polymers form their outer layers or where combustible packaging remains<br>in place.<br>Certain substances, found throughout their construction, may degrade or become volatile when heated to high temperatures. This may create a<br>secondary hazard. |

## **SECTION 6 Accidental release measures**

Personal precautions, protective equipment and emergency procedures See section 8

#### **Environmental precautions**

See section 12

## Methods and material for containment and cleaning up

| Minor Spills | <ul> <li>Clean up all spills immediately.</li> <li>Secure load if safe to do so.</li> <li>Bundle/collect recoverable product.</li> <li>Collect remaining material in containers with covers for disposal.</li> </ul>  |
|--------------|---|
| Major Spills | <ul> <li>Minor hazard.</li> <li>Clear area of personnel.</li> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> <li>Wear physical protective gloves e.g. Leather.</li> <li>Contain spill/secure load if safe to do so.</li> <li>Bundle/collect recoverable product and label for recycling.</li> <li>Collect remaining product and place in appropriate containers for disposal.</li> <li>Clean up/sweep up area.</li> <li>Water may be required.</li> </ul> |

Personal Protective Equipment advice is contained in Section 8 of the SDS.

## **SECTION 7 Handling and storage**

| Precautions for safe handling |   |
|-------------------------------|---|
| Safe handling                 | <ul> <li>Avoid all personal contact, including inhalation.</li> <li>Wear protective clothing when risk of exposure occurs.</li> <li>Use in a well-ventilated area.</li> <li>Prevent concentration in hollows and sumps.</li> <li>DO NOT enter confined spaces until atmosphere has been checked.</li> <li>DO NOT allow material to contact humans, exposed food or food utensils.</li> <li>Avoid contact with incompatible materials.</li> <li>When handling, DO NOT eat, drink or smoke.</li> <li>Keep containers securely sealed when not in use.</li> <li>Avoid physical damage to containers.</li> <li>Always wash hands with soap and water after handling.</li> <li>Work clothes should be laundered separately. Launder contaminated clothing before re-use.</li> <li>Use good occupational work practice.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> <li>Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.</li> </ul> |
| Other information             | Store away from incompatible materials.   |

#### Conditions for safe storage, including any incompatibilities

| Suitable container      | Generally packaging as originally supplied with the article or manufactured item is sufficient to protect against physical hazards.<br>If repackaging is required ensure the article is intact and does not show signs of wear. As far as is practicably possible, reuse the original<br>packaging or something providing a similar level of protection to both the article and the handler.   |
|-------------------------|--|
| Storage incompatibility | <ul> <li>Formaldehyde:</li> <li>is a strong reducing agent</li> <li>may polymerise in air unless properly inhibited (usually with methanol up to 15%) and stored at controlled temperatures</li> <li>will polymerize with active organic material such as phenol</li> <li>reacts violently with strong oxidisers, hydrogen peroxide, potassium permanganate, acrylonitrile, caustics (sodium hydroxide, yielding formic acid and flammable hydrogen), magnesium carbonate, nitromethane, nitrogen oxides (especially a elevated temperatures), peroxyformic acid</li> <li>is incompatible with strong acids (hydrochloric acid forms carcinogenic bis(chloromethyl)ether*), amines, ammonia, aniline, bisulfides,</li> </ul> |

|   | <ul> <li>gelatin, iodine, magnesite, phenol, some motorial catalysis can produce impurities: methy Aqueous solutions of formaldehyde:</li> <li>slowly oxidise in air to produce formic acid</li> <li>attack carbon steel</li> <li>Concentrated solutions containing formaldehyde</li> <li>unstable, both oxidising slowly to form form hydrate (methylene glycol) - the more conce (methanol and amine-containing compound</li> <li>readily subject to polymerisation, at room te formaldehyde), a solid mixture of linear poly also form</li> <li>Flammable and/or toxic gases are generated by strong reducing agents</li> <li>*The empirical equation may be used to determ log(BCME)ppb = -2.25 + 0.67• log(HCHO) ppm - Assume values for formaldehyde, in air, of 1 ppm</li> </ul>  | ylal, methyl formate<br>e are:<br>ic acid and polymerising<br>entrated the solution the<br>ls inhibit polymer formatii<br>emperature, in the preser<br>roxymethylene glycols co<br>y the combination of ald<br>ine the concentration of<br>+ 0.77• log(HCI)ppm | ; in dilute aqueous solutio<br>more polyoxymethylene ;<br>on)<br>nce of air and moisture, to<br>ntaining 90-99% formald<br>shydes with azo, diazo co<br>bis(chloromethyl)ether (B | glycol occurs as oligomers and polymers<br>of form paraformaldehyde (8-100 units of<br>ehyde; a cyclic trimer, trioxane (CH2O3), may<br>ompounds, dithiocarbamates, nitrides, and<br>CME) formed by reaction with HCI: |
|---|--|--|---|--|
| SECTION 8 Exposure contro   | bls / personal protection  |  |   |  |
| Control parameters  |  |  |   |  |
| Occupational Exposure Limits (O<br>INGREDIENT DATA<br>Not Available<br>Emergency Limits |  |  |   |  |
|   | TEEL-1   | TEEL-2   |   | TEEL-3   |
| NS-5111-G   | Not Available  | Not Available  |   | Not Available  |
| Ingredient  | Original IDLH  |  | Revised IDLH  |  |
| N-alkylated benzotriazole   | Not Available  |  | Not Available   |  |
| C7-9 branched alkyl-3,5-di-<br>tert-butyl-<br>4-hydroxyhydrocinnamate                   | Not Available  |  | Not Available   |  |
| Occupational Exposure Banding   |  |  |   |  |
| Ingredient  | Occupational Exposure Band Rating  |  | Occupational Expo   | sure Band Limit  |
| N-alkylated benzotriazole   | E  |  | ≤ 0.1 ppm   |  |
| Notes:  | Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the<br>adverse health outcomes associated with exposure. The output of this process is an occupational exposure band (OEB), which corresponds to a<br>range of exposure concentrations that are expected to protect worker health.   |  |   |  |
| Exposure controls   |  |  |   |  |
| Appropriate engineering<br>controls   | Articles or manufactured items, in their original of<br>Exceptions may arise following extensive use an<br>article, may be released to the environment.  |  |   |  |
| Personal protection   |  |  |   |  |
| Eye and face protection   | <ul> <li>Safety glasses.</li> <li>Safety glasses with side shields.</li> <li>Chemical goggles.</li> <li>Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]</li> </ul> |  |   |  |
| Skin protection   | See Hand protection below  |  |   |  |
| Hands/feet protection   | See Hand protection below Wear general protective gloves, eg. light weight rubber gloves. NOTE:  The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact. Contaminated leather items, such as shoes, belts and watch-bands should be removed and destroyed.  |  |   |  |
| Body protection   | See Other protection below   |  |   |  |
| Other protection  | <ul> <li>Overalls.</li> <li>P.V.C apron.</li> <li>Barrier cream.</li> <li>Skin cleansing cream.</li> <li>Eye wash unit.</li> </ul>   |  |   |  |

Continued...

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## **Respiratory protection**

Respiratory protection not normally required due to the physical form of the product.

## **SECTION 9** Physical and chemical properties

## Information on basic physical and chemical properties

| Appearance                                   | Cream colored grease, bland odor |   |                |
|--|----------------------------------|---|----------------|
| Physical state                               | Manufactured                     | Relative density (Water = 1)            | Not Available  |
| Odour  | Not Available                    | Partition coefficient n-octanol / water | Not Available  |
| Odour threshold                              | Not Available                    | Auto-ignition temperature (°C)          | Not Available  |
| pH (as supplied)                             | Not Available                    | Decomposition<br>temperature (°C)       | Not Available  |
| Melting point / freezing point<br>(°C)       | Not Available                    | Viscosity (cSt)                         | Not Available  |
| Initial boiling point and boiling range (°C) | Not Available                    | Molecular weight (g/mol)                | Not Available  |
| Flash point (°C)                             | Not Available                    | Taste                                   | Not Available  |
| Evaporation rate                             | Not Available                    | Explosive properties                    | Not Available  |
| Flammability                                 | Not Available                    | Oxidising properties                    | Not Available  |
| Upper Explosive Limit (%)                    | Not Available                    | Surface Tension (dyn/cm or<br>mN/m)     | Not Applicable |
| Lower Explosive Limit (%)                    | Not Available                    | Volatile Component (%vol)               | Not Available  |
| Vapour pressure (kPa)                        | Not Available                    | Gas group                               | Not Available  |
| Solubility in water                          | Immiscible                       | pH as a solution (Not<br>Available%)    | Not Available  |
| Vapour density (Air = 1)                     | Not Available                    | VOC g/L                                 | Not Available  |

## **SECTION 10 Stability and reactivity**

| Reactivity                          | See section 7   |
|-------------------------------------|---|
| Chemical stability                  | Product is considered stable and hazardous polymerisation will not occur. |
| Possibility of hazardous reactions  | See section 7   |
| Conditions to avoid                 | See section 7   |
| Incompatible materials              | See section 7   |
| Hazardous decomposition<br>products | See section 5   |

## **SECTION 11 Toxicological information**

#### Information on toxicological effects

| Inhaled      | The material is not thought to produce adverse health effects or irritation models). Nevertheless, good hygiene practice requires that exposure be occupational setting.  |   |  |
|--------------|---|---|--|
| Ingestion    | The material has <b>NOT</b> been classified by EC Directives or other classification corroborating animal or human evidence.  | ation systems as "harmful by ingestion". This is because of the lack of |  |
| Skin Contact | The material is not thought to produce adverse health effects or skin irritation following contact (as classified by EC Directives using animal models). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable gloves be used in an occupational setting. |   |  |
| Eye          | Although the material is not thought to be an irritant (as classified by EC Directives), direct contact with the eye may produce transient discomfort characterised by tearing or conjunctival redness (as with windburn).  |   |  |
| Chronic      | Skin contact with the material is more likely to cause a sensitisation reaction in some persons compared to the general population.   |   |  |
|              |   |   |  |
| NS-5111-G    | TOXICITY<br>Not Available   | IRRITATION<br>Not Available   |  |

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|  | ΤΟΧΙCΙΤΥ   | IRRITATION  |  |
|--|--|---|--|
| N-alkylated benzotriazole  | dermal (rat) LD50: >2000 mg/kg <sup>[2]</sup>  | Not Available   |  |
|  | Oral (Rat) LD50; 3300 mg/kg <sup>[2]</sup>   |   |  |
|  | ΤΟΧΙΟΙΤΥ   | IRRITATION  |  |
| C7-9 branched alkyl-3,5-di-<br>tert-butyl-   | dermal (rat) LD50: >2000 mg/kg <sup>[1]</sup>  | Eye (rabbit: non-ir   | ritating *   |
| 4-hydroxyhydrocinnamate  | Oral (Rat) LD50; >200 mg/kg <sup>[2]</sup>   | Skin (rat): non-irrit   | •  |
| Legend:  | 1. Value obtained from Europe ECHA Registered Substan<br>specified data extracted from RTECS - Register of Toxic E   | -   | ned from manufacturer's SDS. Unless otherwise  |
| N-ALKYLATED<br>BENZOTRIAZOLE   | *RT Vanderbilt MSDS Repeat dose toxicity: A combined test (OECD 422) revealed parental toxicity: A top mg/kg reduced thymus organ weight, and microscopic findings per day Genetic toxicity: The test compound did not cause analogue did not reveal any potential for clastogenic effects of benzotriazoles. There are several indications that the effects of phenolic reduced concentrations of testosterone, higher concentre As in these cases there are also indications for toxic effects are several benzotriazole UV stabilisers showed significant immunity, stem cell maintenance, and cellular differentia accumulate and exert potent physiological effects in hum stable and toxic ligands. The polycyclic aromatic hydroca induces its own metabolism and bioactivation to a toxic or Benzotriazole is the core structure present within the phr formation of 5- and 4-hydroxybenzotriazole (1.6 and 0.3) amount added) Oral acute studies in rats and mice yield mice and rats ranged from 400-1000 and 500-900 mg/kg values were =1000 mg/kg in rats and rabbits, and inhala short-term studies showed that oral administration to mic weight were observed in rats. Endocrine effects, normoor mg/kg. No effects on deaths and no clinical symptoms w Additionally, no dose-related effects on reproductive organst fed 12,100 ppm benzotriazole for 78 weeks. However effects could not be determined. Brain tumors occurred in increase was not observed in female mice fed 23,500 pp incidences varied from 0 to 7%. Genotoxicity studies ind TA100 in the presence or absence of S9, or Chinese har TA1535 in the absence of S9, but was mutagenic in the TA1537 and TA1538 and E. coll WP2 uvrA. It did not provinduced chromosomal aberrations in the presence of S9 in the mouse micronucleus assay at 800 mg/kg. Benzotr Benzotriazoles was identified as irritating to rabbit eyes are for phenolic benzotriazoles. Overall, oral exposure (either through gavage or in feed) liver weights were observed in several studies. Body we substances. Histopathological changes (e.g.,foci, hypert were | bw (clinical signs, reduced body w<br>in the thymus and spleen). The NC<br>se mutations in bacteria and in mar<br>acts in mammalian cells ** REACh I<br>benzotriazoles described in the lite<br>ations of CYP 450, or higher activit<br>acts on the liver reported, the effect<br>ambiguously as endocrine adverse<br>human aryl hydrocarbon receptor (.<br>tion A study indicated that certain b<br>nans, analogous to polycyclic arom<br>arbon the polycyclic aromatic hydro<br>metabolites.<br>enolic benzotriazole class. In vitro r<br>2% of the amount added, respectiv<br>ed LD50 values that ranged from 5<br>g, respectively. A mouse intravenou<br>tion LC50 values in rats were 1.5 n<br>ce produced minimal effects on boc<br>cytic anemia, and leukopenia were 1.<br>8 weeks (22%), but not in female rat.<br>8 weeks (22%), but not in female rat.<br>8 weeks (22%), but not in female rat.<br>8 weeks (22%), but not in female rat.<br>9 metabCaST fed 11,700 ppm benz<br>om benzotriazole for the same perior<br>icate that the compound was not m<br>mster ovary cells. Benzotriazole was<br>presence of S9.Conflicting results w<br>duce DNA damage in E. coli PQ37<br>and sister chromatid exchange in 1<br>riazole was identified as a non-sens<br>nd minimally irritating to rabbit and<br>0 of the tested chemicals to rats led<br>ight and body weight gain changes<br>rophy, and cytoplasmic vacuolizatic<br>enzotriazoles. Haematological effect | eight gains with lower food consumption, slightly<br>VAEL was considered to be 45 mg/kg body weight<br>nmalian cell culture Data obtained with a structural<br>Dossier<br>erature might be caused by endocrine disruption, e<br>y of ethoxyresorufin-O-deethylase (EROD-activity)<br>s might actually be only secondary effects. With the<br>effects of an equivalent level of concern.<br>AhR) ligand activity. The AhR has roles in regulating<br>renzotriazole UV stabilisers have the potential to<br>atic hydrocarbons and dioxins, which are known<br>carbon, benzo[a]pyrene (BaP), a ligand for AhR,<br>netabolism with rat liver microsomes yielded<br>ely).Overall metabolism was low (<5% of the total<br>60 to 909 mg/kg. Intraperitoneal LD50 values in<br>s LD50 of 238 mg/kg was identified. Dermal LD50<br>ug/L and 1.91 mg/L/3 hours). Subchronic and<br>ly weight while dose-dependent decreases in body<br>noted in rats dosed for 26 weeks. The TDLo was 10<br>inistered (in food) benzotriazole =78 weeks.<br>lastic liver nodules were observed in male Fischer<br>nees varied from 0 to 11% so the treatment-related<br>incidance of endometrial stromal polyps was<br>tts fed 12,100 ppm (16%). Significant increase in<br>otrizzole for 104 weeks. Comparatively, a.similar<br>od of time (6% increase). Historical laboratory contu<br>utagenic to S. typhimurium strains TA97, TA98, or<br>s also not mutagenic to S. typhimurium strains<br>i. In Chinese hamster ovary cells, benzotriazole<br>he absence of S9. Benzotriazole was not genotoxi<br>itizer in the guinea pig maximization test.<br>guinea pig skin<br>to liver effects. Increased absolute and/or relative<br>were observed after administration of several test<br>on) and altered liver enzyme content and activities |
| C7-9 BRANCHED ALKYL-3,5-DI-<br>TERT-BUTYL-   | Reproductive and teratology effects: The chemicals te<br>organ weights, but no direct studies in reproduction and<br>Genotoxicity None of the tested compounds were ident<br>vivo<br>Chemical Information Review Document for Phenolic Be<br>Toxicology Program October 2011<br>http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic<br>No significant acute toxicological data identified in literat<br>Non-sensitising to guinea pig skin * Everspring Chemica<br>Data show that acute toxicity following oral and topical u  | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.  | Some chemicals were shown to affect reproductiv<br>sence or presence of a metabolic system (S9) or i<br>on for Toxicological Evaluation by the National<br>odf   |
|  | Reproductive and teratology effects: The chemicals to organ weights, but no direct studies in reproduction and Genotoxicity None of the tested compounds were ident vivo           Chemical Information Review Document for Phenolic Be Toxicology Program October 2011           http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic No significant acute toxicological data identified in literat           Non-sensitising to guinea pig skin * Everspring Chemica Data show that acute toxicity following oral and topical u term use may affect the liver, thyroid, kidney and lymph           The following information refers to contact allergens as a Contact allergies quickly manifest themselves as contact ezzema involves a cell-mediated (T lymphocytes) immur involve antibody-mediated immune reactions. The signifi distribution of the substance and the opportunities for co distributed can be a more important allergen than one w   | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>I MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to a<br>t eczema, more rarely as urticaria o<br>ne reaction of the delayed type. Ott<br>icance of the contact allergen is no<br>intact with it are equally important. <i>i</i><br>ith stronger sensitising potential with   | Some chemicals were shown to affect reproductive<br>sence or presence of a metabolic system (S9) or in<br>on for Toxicological Evaluation by the National<br>odf<br>are not proven to cause mutations. However, long<br>ported.<br>this product.<br>or Quincke's oedema. The pathogenesis of contact<br>ther allergic skin reactions, e.g. contact urticaria,<br>simply determined by its sensitisation potential: th<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a   |
| TERT-BUTYL-<br>4-HYDROXYHYDROCINNAMATE<br>NS-5111-G & N-ALKYLATED  | Reproductive and teratology effects: The chemicals to organ weights, but no direct studies in reproduction and Genotoxicity None of the tested compounds were ident vivo           Chemical Information Review Document for Phenolic Be Toxicology Program October 2011           http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic No significant acute toxicological data identified in literat           Non-sensitising to guinea pig skin * Everspring Chemica Data show that acute toxicity following oral and topical u term use may affect the liver, thyroid, kidney and lymph           The following information refers to contact allergens as a Contact allergies quickly manifest themselves as contact eczema involves a cell-mediated (T lymphocytes) immuri involve antibody-mediated immune reactions. The signific distribution of the substance and the opportunities for con distributed can be a more important allergen than one w clinical point of view, substances are noteworthy if they provide the substance and the opportunity of   | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>I MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to a<br>t eczema, more rarely as urticaria o<br>ne reaction of the delayed type. Ott<br>icance of the contact allergen is no<br>intact with it are equally important. <i>i</i><br>ith stronger sensitising potential with   | Some chemicals were shown to affect reproductiv<br>sence or presence of a metabolic system (S9) or in<br>on for Toxicological Evaluation by the National<br>odf<br>are not proven to cause mutations. However, long<br>ported.<br>this product.<br>or Quincke's oedema. The pathogenesis of contact<br>her allergic skin reactions, e.g. contact urticaria,<br>a simply determined by its sensitisation potential: It<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a<br>more than 1% of the persons tested.  |
| TERT-BUTYL-<br>4-HYDROXYHYDROCINNAMATE<br>NS-5111-G & N-ALKYLATED<br>BENZOTRIAZOLE<br>Acute Toxicity   | Reproductive and teratology effects: The chemicals to<br>organ weights, but no direct studies in reproduction and<br>Genotoxicity None of the tested compounds were ident<br>vivo<br>Chemical Information Review Document for Phenolic Be<br>Toxicology Program October 2011<br>http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic<br>No significant acute toxicological data identified in literat<br>Non-sensitising to guinea pig skin * Everspring Chemica<br>Data show that acute toxicity following oral and topical u<br>term use may affect the liver, thyroid, kidney and lymph i<br>The following information refers to contact allergens as a<br>Contact allergies quickly manifest themselves as contac<br>eczema involves a cell-mediated (T lymphorytes) immur<br>involve antibody-mediated immune reactions. The signifi<br>distribution of the substance and the opportunities for co<br>distributed can be a more important allergen than one w<br>clinical point of view, substances are noteworthy if they p   | ested produced a variety of effects.<br>development were located.<br>lified as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>If MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to o<br>t eczema, more rarely as urticaria o<br>t eczema, more rarely as urticaria to<br>t is to of the delayed type. Oth<br>icance of the contact allergen is no<br>intact with it are equally important. A<br>ith stronger sensitising potential with<br>produce an allergic test reaction in   | Some chemicals were shown to affect reproductiv<br>sence or presence of a metabolic system (S9) or in<br>on for Toxicological Evaluation by the National<br>odf<br>ware not proven to cause mutations. However, long<br>ported.<br>this product.<br>or Quincke's oedema. The pathogenesis of contact<br>uer allergic skin reactions, e.g. contact urticaria,<br>simply determined by its sensitisation potential: th<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a<br>more than 1% of the persons tested.   |
| TERT-BUTYL-<br>4-HYDROXYHYDROCINNAMATE<br>NS-5111-G & N-ALKYLATED<br>BENZOTRIAZOLE<br>Acute Toxicity<br>Skin Irritation/Corrosion                                  | Reproductive and teratology effects: The chemicals to<br>organ weights, but no direct studies in reproduction and<br>Genotoxicity None of the tested compounds were ident<br>vivo<br>Chemical Information Review Document for Phenolic Be<br>Toxicology Program October 2011<br>http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic<br>No significant acute toxicological data identified in literat<br>Non-sensitising to guinea pig skin * Everspring Chemica<br>Data show that acute toxicity following oral and topical u<br>term use may affect the liver, thyroid, kidney and lymph i<br>The following information refers to contact allergens as a<br>Contact allergies quickly manifest themselves as contac<br>eczema involves a cell-mediated (T lymphocytes) immur<br>involve antibody-mediated immune reactions. The signifi<br>distribution of the substance and the opportunities for co<br>distributed can be a more important allergen than one w<br>clinical point of view, substances are noteworthy if they p   | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>I MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to i<br>t eczema, more rarely as urticaria of<br>t eczema, more rarely as urticaria of<br>ne reaction of the delayed type. Oth<br>icance of the contact allergen is no<br>intact with it are equally important. A<br>ith stronger sensitising potential with<br>produce an allergic test reaction in the<br>Carcinogenicity<br>Reproductivity   | Some chemicals were shown to affect reproductive<br>sence or presence of a metabolic system (S9) or it<br>on for Toxicological Evaluation by the National<br>odf<br>ware not proven to cause mutations. However, long<br>ported.<br>this product.<br>or Quincke's oedema. The pathogenesis of contact<br>were allergic skin reactions, e.g. contact urticaria,<br>a simply determined by its sensitisation potential: th<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a<br>more than 1% of the persons tested.   |
| TERT-BUTYL-<br>4-HYDROXYHYDROCINNAMATE<br>NS-5111-G & N-ALKYLATED<br>BENZOTRIAZOLE<br>Acute Toxicity<br>Skin Irritation/Corrosion<br>Serious Eye Damage/Irritation | Reproductive and teratology effects: The chemicals to<br>organ weights, but no direct studies in reproduction and<br>Genotoxicity None of the tested compounds were ident<br>vivo<br>Chemical Information Review Document for Phenolic Be<br>Toxicology Program October 2011<br>http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic<br>No significant acute toxicological data identified in literat<br>Non-sensitising to guinea pig skin * Everspring Chemica<br>Data show that acute toxicity following oral and topical u<br>term use may affect the liver, thyroid, kidney and lymph i<br>The following information refers to contact allergens as a<br>Contact allergies quickly manifest themselves as contact<br>eczema involves a cell-mediated (T lymphocytes) immur<br>involve antibody-mediated immune reactions. The signifi<br>distribution of the substance and the opportunities for co<br>distributed can be a more important allergen than one w<br>clinical point of view, substances are noteworthy if they p  | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>If MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to a<br>t eczema, more rarely as urticaria of<br>ne reaction of the delayed type. Ott<br>icance of the contact allergen is no<br>intact with it are equally important. <i>i</i><br>ith stronger sensitising potential with<br>produce an allergic test reaction in in<br>Carcinogenicity<br>Reproductivity<br>STOT - Single Exposure   | Some chemicals were shown to affect reproductive<br>sence or presence of a metabolic system (S9) or it<br>on for Toxicological Evaluation by the National<br>odf<br>ware not proven to cause mutations. However, long<br>ported.<br>This product.<br>In Quincke's oedema. The pathogenesis of contact<br>urricaria, e.g. contact urticaria,<br>er allergic skin reactions, e.g. contact urticaria,<br>simply determined by its sensitisation potential: th<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a<br>more than 1% of the persons tested.   |
| TERT-BUTYL-<br>4-HYDROXYHYDROCINNAMATE<br>NS-5111-G & N-ALKYLATED<br>BENZOTRIAZOLE<br>Acute Toxicity<br>Skin Irritation/Corrosion                                  | Reproductive and teratology effects: The chemicals to<br>organ weights, but no direct studies in reproduction and<br>Genotoxicity None of the tested compounds were ident<br>vivo<br>Chemical Information Review Document for Phenolic Be<br>Toxicology Program October 2011<br>http://ntp.niehs.nih.gov/ntp/noms/support_docs/phenolic<br>No significant acute toxicological data identified in literat<br>Non-sensitising to guinea pig skin * Everspring Chemica<br>Data show that acute toxicity following oral and topical u<br>term use may affect the liver, thyroid, kidney and lymph i<br>The following information refers to contact allergens as a<br>Contact allergies quickly manifest themselves as contac<br>eczema involves a cell-mediated (T lymphocytes) immur<br>involve antibody-mediated immune reactions. The signifi<br>distribution of the substance and the opportunities for co<br>distributed can be a more important allergen than one w<br>clinical point of view, substances are noteworthy if they p   | ested produced a variety of effects.<br>development were located.<br>iffied as mutagenic in vitro in the ab<br>enzotriazoles: Supporting Nominatic<br>benzotriazoles_cird_oct2011_508.p<br>ure search.<br>I MSDS<br>se of hindered phenols is low. They<br>nodes. Liver tumours have been re<br>a group and may not be specific to i<br>t eczema, more rarely as urticaria of<br>t eczema, more rarely as urticaria of<br>ne reaction of the delayed type. Oth<br>icance of the contact allergen is no<br>intact with it are equally important. A<br>ith stronger sensitising potential with<br>produce an allergic test reaction in the<br>Carcinogenicity<br>Reproductivity   | Some chemicals were shown to affect reproductive<br>sence or presence of a metabolic system (S9) or it<br>on for Toxicological Evaluation by the National<br>odf<br>ware not proven to cause mutations. However, long<br>ported.<br>this product.<br>or Quincke's oedema. The pathogenesis of contact<br>were allergic skin reactions, e.g. contact urticaria,<br>a simply determined by its sensitisation potential: th<br>A weakly sensitising substance which is widely<br>h which few individuals come into contact. From a<br>more than 1% of the persons tested.   |

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Continued...

Legena:

– Data either not available or does not till the criteria for classification

Data available to make classification

## **SECTION 12 Ecological information**

|  | Endpoint         | Test Duration (hr) | Species                       | Value            | Source         |
|--|------------------|--------------------|-------------------------------|------------------|----------------|
| NS-5111-G                                  | Not<br>Available | Not Available      | Not Available                 | Not<br>Available | Not<br>Availab |
|  | Endpoint         | Test Duration (hr) | Species                       | Value            | Source         |
| N-alkylated benzotriazole                  | EC50(ECx)        | 24h                | Crustacea                     | 1.4mg/l          | Not<br>Availab |
|  | LC50             | 96h                | Fish                          | 1.3mg/l          | Not<br>Availab |
|  | Endpoint         | Test Duration (hr) | Species                       | Value            | Source         |
|  | EC50             | 72h                | Algae or other aquatic plants | 3mg/l            | Not<br>Availab |
| C7-9 branched alkyl-3,5-di-<br>tert-butyl- | EC50             | 48h                | Crustacea                     | >0.008mg/l       | 2              |
| tert-butyi-<br>4-hydroxyhydrocinnamate     | EC50(ECx)        | 72h                | Algae or other aquatic plants | 3mg/l            | Not<br>Availab |
|  | LC50             | 96h                | Fish                          | >74mg/l          | Not<br>Availab |

٩qu DXICITY ment Data 6. NITE (Jap an) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

#### Persistence and degradability Ingredient Persistence: Water/Soil Persistence: Air No Data available for all ingredients No Data available for all ingredients

| Bioaccumulative potential |                                       |
|---------------------------|---------------------------------------|
| Ingredient                | Bioaccumulation                       |
|                           | No Data available for all ingredients |
| Mahilitain anil           |                                       |
| Mobility in soil          |                                       |
| Ingredient                | Mobility                              |
|                           | No Data available for all ingredients |

## **SECTION 13 Disposal considerations**

| Waste treatment methods      |  |  |
|------------------------------|--|--|
| Product / Packaging disposal | <ul> <li>Recycle wherever possible or consult manufacturer for recycling options.</li> <li>Consult State Land Waste Management Authority for disposal.</li> <li>Recycle wherever possible or consult manufacturer for recycling options.</li> <li>Consult State Land Waste Authority for disposal.</li> <li>Bury or incinerate residue at an approved site.</li> <li>Recycle containers if possible, or dispose of in an authorised landfill.</li> </ul> |  |

### **SECTION 14 Transport information**

| Labels Required  |    |
|------------------|----|
| Marine Pollutant | NO |

## Land transport (DOT): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

#### Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

## Sea transport (IMDG-Code / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Transport in bulk according to Annex II of MARPOL and the IBC code

#### Not Applicable

## Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

| Product name              | Group         |
|---------------------------|---------------|
| N-alkylated benzotriazole | Not Available |

| ersion No: 1.1  | Page 8 of 9                 |                            | Issue Date: 08/19/202       |   |
|---|-----------------------------|----------------------------|-----------------------------|---|
|   |                             | NS-5111-G                  | <u> </u>                    | Print Date: 08/19/202                         |
|   |                             |                            |                             |   |
| Product name  | Group                       |                            |                             |   |
| C7-9 branched alkyl-3,5-di-<br>tert-butyl-                            | Not Available               |                            |                             |   |
| 4-hydroxyhydrocinnamate   |                             |                            |                             |   |
| Transport in bulk in accordan   | ce with the ICG Code        |                            |                             |   |
| Product name  | Ship Type                   |                            |                             |   |
| N-alkylated benzotriazole   | Not Available               |                            |                             |   |
| C7-9 branched alkyl-3,5-di-<br>tert-butyl-<br>4-hydroxyhydrocinnamate | Not Available               |                            |                             |   |
| SECTION 15 Regulatory inf   | ormation                    |                            |                             |   |
| Safety, health and environme  | atal regulations / logislat | tion specific for the sub- | stanco or mixturo           |   |
|   |                             | •                          |                             |   |
| N-alkylated benzotriazole is fou<br>US Toxic Substances Control Act   |                             | •                          | LIS TSCA Chemical Substance | Inventory - Interim List of Active Substances |
|   |                             |                            |                             | inventory - Interim List of Active Substances |
| C7-9 branched alkyl-3,5-di-tert-                                      |                             |                            |                             |   |
| US Toxic Substances Control Act                                       | (TSCA) - Chemical Substance | e Inventory                | US ISCA Chemical Substance  | Inventory - Interim List of Active Substances |
| Federal Regulations   |                             |                            |                             |   |
| Superfund Amendments and  | Reauthorization Act of 1    | 986 (SARA)                 |                             |   |
| Section 311/312 hazard categor  | ies                         |                            |                             |   |
| Flammable (Gases, Aerosols, Liq                                       |                             |                            |                             | No  |
| Gas under pressure  | . ,                         |                            |                             | No  |
| Explosive   |                             |                            |                             | No  |
| Self-heating  |                             |                            |                             | No  |
| Pyrophoric (Liquid or Solid)  |                             |                            |                             | No  |
| Pyrophoric Gas  |                             |                            |                             | No  |
| Corrosive to metal  |                             |                            |                             | No  |
| Oxidizer (Liquid, Solid or Gas)                                       |                             |                            |                             | No  |
| Organic Peroxide  |                             |                            |                             | No  |
| Self-reactive   |                             |                            |                             | No  |
| In contact with water emits flamma                                    | able gas                    |                            |                             | No  |
| Combustible Dust  |                             |                            |                             | No  |
| Carcinogenicity   |                             |                            |                             | No  |
| Acute toxicity (any route of exposi                                   | ure)                        |                            |                             | No  |
| Reproductive toxicity   |                             |                            |                             | No  |
| Skin Corrosion or Irritation  |                             |                            |                             | No  |
| Respiratory or Skin Sensitization                                     |                             |                            |                             | Yes   |
| Serious eye damage or eye irritati                                    | on                          |                            |                             | No  |
| Specific target organ toxicity (sing                                  | le or repeated exposure)    |                            |                             | No  |
| Aspiration Hazard   |                             |                            |                             | No  |
| Germ cell mutagenicity  |                             |                            |                             | No  |
| Simple Asphyxiant   |                             |                            |                             | No  |
| Hazards Not Otherwise Classified                                      |                             |                            |                             | No  |
| US. EPA CERCLA Hazardous So<br>None Reported                          | ubstances and Reportable C  | Quantities (40 CFR 302.4)  |                             |   |
|   |                             |                            |                             |   |
| State Regulations   |                             |                            |                             |   |
| State Regulations US. California Proposition 65                       |                             |                            |                             |   |

## National Inventory Status

| National Inventory                                 | Status  |  |
|--|---|--|
| Australia - AIIC / Australia<br>Non-Industrial Use | Yes   |  |
| Canada - DSL                                       | Yes   |  |
| Canada - NDSL                                      | No (N-alkylated benzotriazole; C7-9 branched alkyl-3,5-di-tert-butyl-4-hydroxyhydrocinnamate) |  |
| China - IECSC                                      | Yes   |  |
| Europe - EINEC / ELINCS / NLP                      | No (N-alkylated benzotriazole; C7-9 branched alkyl-3,5-di-tert-butyl-4-hydroxyhydrocinnamate) |  |
| Japan - ENCS                                       | No (N-alkylated benzotriazole; C7-9 branched alkyl-3,5-di-tert-butyl-4-hydroxyhydrocinnamate) |  |



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| National Inventory  | Status  |  |  |
|---------------------|---|--|--|
| Korea - KECI        | Yes   |  |  |
| New Zealand - NZIoC | Yes   |  |  |
| Philippines - PICCS | Yes   |  |  |
| USA - TSCA          | Yes   |  |  |
| Taiwan - TCSI       | Yes   |  |  |
| Mexico - INSQ       | No (N-alkylated benzotriazole)  |  |  |
| Vietnam - NCI       | Yes   |  |  |
| Russia - FBEPH      | No (C7-9 branched alkyl-3,5-di-tert-butyl-4-hydroxyhydrocinnamate)  |  |  |
| Legend:             | Yes = All CAS declared ingredients are on the inventory<br>No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration. |  |  |

#### **SECTION 16 Other information**

| Revision Date | 08/19/2022 |
|---------------|------------|
| Initial Date  | 08/20/2022 |

#### Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chemwatch Classification committee using available literature references.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

#### **Definitions and abbreviations**

PC-TWA: Permissible Concentration-Time Weighted Average PC-STEL: Permissible Concentration-Short Term Exposure Limit IARC: International Agency for Research on Cancer ACGIH: American Conference of Governmental Industrial Hygienists STEL: Short Term Exposure Limit TEEL: Temporary Emergency Exposure Limit. IDLH: Immediately Dangerous to Life or Health Concentrations ES: Exposure Standard OSF: Odour Safety Factor NOAEL :No Observed Adverse Effect Level LOAEL: Lowest Observed Adverse Effect Level TLV: Threshold Limit Value LOD: Limit Of Detection OTV: Odour Threshold Value BCF: BioConcentration Factors BEI: Biological Exposure Index AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

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end of SDS