

SAFETY DATA SHEET

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

| 1.1. Product identifier | | | | |
|--|---|---|--|---|
| Trade name or designation of the mixture | IMURAN TABLE | TS | | |
| Registration number | - | | | |
| Synonyms | | TABLETS * IMURAN THIOPRINE, FORMU | 50 MG TABLETS * IMUREK FILMTABI LATED PRODUCT | LETTEN * IMUREL |
| Issue date | 06-November-20 |)14 | | |
| Version number | 13 | | | |
| Revision date | 06-November-20 |)14 | | |
| 1.2. Relevant identified uses of | the substance or | mixture and uses ad | vised against | |
| Identified uses | Medicinal Produc | ct. | | |
| | handling this forr to medicinal use information/pack safety informatio | mulated product in the of the product. In this age insert/product lab | vide health, safety and environmental in workplace. It is not intended to provide instance patients should consult prescr el or consult their pharmacist or physicis ients used during manufacturing, refer to | information relevant ibing an. For health and |
| Uses advised against | No other uses ar | e advised. | | |
| 1.3. Details of the supplier of the | ne safety data shee | et | | |
| | | Road esex TW8 9GS UK | ess hours): +44-20-8047-5000 | |
| 1.4. Emergency telephone number | | | | |
| | | ll call: | +(44)-870-8200418 +1 703 527 3887 e response | |
| SECTION 2: Hazards ider | ntification | | | |
| 2.1. Classification of the substa | ance or mixture | | | |

Classification according to Directive 67/548/EEC or 1999/45/EC as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

Classification according to Regulation (EC) No 1272/2008 as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

2.2. Label elements

Label according to Regulation (EC) No. 1272/2008 as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

2.3. Other hazards Caution - Pharmaceutical agent. See section 11 for additional information on health hazards.

SECTION 3: Composition/information on ingredients

3.2. Mixtures

General information

| Chemical name | | % | CAS-No | / FC No | REACH Registration No. | | Notes |
|--|---|--|--------------------------------|---------------------------|---|--|---------------------------|
| | | /0 | 0/10-110. | , <u> </u> | | | 110163 |
| AZATHIOPRINE | | 31.4 - < 31.8 | | -86-6 175-4 | - | - | |
| Classification: | DSD: | Carc. Cat. 1;R4 Xn;R22-48, Xi;F | | | Muta. Cat. 2;R46, Repr. Cat. | 2;R61, | |
| | CLP: | Acute Tox. 4;H3 STOT SE 3;H3 1;H372 | 302, Skin Ir 35, Muta. 1 | rit. 2;H315 B;H340, C | , Skin Sens. 1;H317, Eye Irr arc. 1A;H350, Repr. 1B;H36 | it. 2;H319, 0, STOT RE | |
| Starch | | 10 - < 20 | | -25-8 679-6 | - | - | |
| Classification: | DSD: | - | | | | | |
| | CLP: | - | | | | | |
| HYDROXYPROPYL MET CELLULOSE | THYL | 1 - < 3 | 9004 | -65-3 | - | - | |
| Classification: | DSD: | - | | | | | |
| | CLP: | - | | | | | |
| | | | | | | | |
| MAGNESIUM STEARAT | | < 1 | | -04-0 150-3 | - | - | |
| Classification: | DSD: | | | | | | |
| | CLP: | - | | | | | |
| Titanium dioxide | | < 0.2 | | 3-67-7 675-5 | - | - | |
| Classification: | DSD: | - | | | | | |
| | CLP: | - | | | | | |
| Other components below | reporta | ble levels 40 - | < 50 | | | | |
| List of abbreviations and sy CLP: Regulation No. 127 DSD: Directive 67/548/EF M: M-factor vPvB: very persistent and PBT: persistent, bioaccur #: This substance has be | 2/2008. EC. d very bi nulative | ioaccumulative s and toxic substa | ubstance. ance. | exposure | limit(s). | | |
| Composition comments | | | - | - | displayed in section 16. | | |
| SECTION 4: First aid n | neasu | res | | | | | |
| General information | v | | Ensure that | medical p | well, seek medical advice im ersonnel are aware of the m | | |
| I.1. Description of first aid r | neasur | es | | | | | |
| Inhalation | s | | p or persist | . Under no | t, trained personnel should g ormal conditions of intended | | |
| Skin contact | | mmediately flush Get medical atten | | | ater. Take off contaminated our. | clothing and wash | ı before reus |
| Eye contact | | | | | or at least 15 minutes and co | | |
| Ingestion | а | f swallowed, rinse mount does occu advice from poiso | ur, call a po | ison contr | nly if the person is conscious ol centre immediately. Do no | s). If ingestion of a tinduce vomiting | a large without |
| .2. Most important sympto ind effects, both acute and lelayed | ms A h | Accidental exposi lives, itching, and | ure or conta I difficulty b | act might p reathing), | roduce: symptoms of hypers nausea, vomiting, increased | ensitivity (such a susceptibility to i | s skin rash, nfection. |
| 4.3. Indication of any mmediate medical attention and special treatment neede | n a | | e, refer to | | d. Treat according to locally a t prescribing information or to | | |

SECTION 5: Firefighting measures

| General fire hazards | No unusual fire or explosion hazards noted. |
|---|---|
| 5.1. Extinguishing media Suitable extinguishing media | Water. Foam. Dry chemical powder. Carbon dioxide (CO2). |
| Unsuitable extinguishing media | None known. |
| 5.2. Special hazards arising from the substance or mixture | During fire, gases hazardous to health may be formed. |
| 5.3. Advice for firefighters | |
| Special protective equipment for firefighters | Self-contained breathing apparatus and full protective clothing must be worn in case of fire. |
| Special fire fighting procedures | Move containers from fire area if you can do so without risk. |
| Specific methods | Use standard firefighting procedures and consider the hazards of other involved materials. |

SECTION 6: Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

| For non-emergency personnel | Keep unnecessary personnel away. Keep people away from and upwind of spill/leak. Keep out of low areas. Wear appropriate protective equipment and clothing during clean-up. Do not touch damaged containers or spilled material unless wearing appropriate protective clothing. Ensure adequate ventilation. Local authorities should be advised if significant spillages cannot be contained. For personal protection, see section 8. |
|--|--|
| For emergency responders | Keep unnecessary personnel away. Use personal protection recommended in Section 8 of the SDS. |
| 6.2. Environmental precautions | Avoid discharge into drains, water courses or onto the ground. |
| 6.3. Methods and material for containment and cleaning up | Stop the flow of material, if this is without risk. Prevent entry into waterways, sewer, basements or confined areas. Following product recovery, flush area with water. |
| 6.4. Reference to other sections | For personal protection, see section 8. For waste disposal, see section 13 of the SDS. |

SECTION 7: Handling and storage

| 7.1. Precautions for safe handling | Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Avoid contact with eyes, skin, and clothing. Avoid prolonged exposure. Do not taste or swallow. When using, do not eat, drink or smoke. Pregnant or breastfeeding women must not handle this product. Should be handled in closed systems, if possible. Provide adequate ventilation. Wear appropriate personal protective equipment. Wash hands thoroughly after handling. Observe good industrial hygiene practices. |
|---|---|
| 7.2. Conditions for safe storage, including any incompatibilities | Store locked up. Store in original tightly closed container. Store away from incompatible materials (see Section 10 of the SDS). |
| 7.3. Specific end use(s) | Medicinal Product. |

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

Occupational exposure limits

| GSK | | | |
|--|---------------|----------|---------------------|
| Components | Туре | Value | Note |
| AZATHIOPRINE (CAS 446-86-6) | 8 HR TWA | 3 mcg/m3 | |
| , | OHC | 4 | Carcinogen |
| | | 4 | Reproductive hazard |
| | | 4 | SKIN SENSITISER |
| HYDROXYPROPYL METHYL CELLULOSE (CAS 9004-65-3) | OHC | 1 | |
| UK. EH40 Workplace Exposure L | .imits (WELs) | | |
| Components | Туре | Value | Form |
| Starch (CAS 9005-25-8) | TWA | 4 mg/m3 | Respirable. |
| | | 10 mg/m3 | Inhalable |
| Titanium dioxide (CAS 13463-67-7) | TWA | 4 mg/m3 | Respirable. |

| UK. EH40 Workplace Expos Components | Туре | Value | Form | |
|--|--|--|---|--|
| | | 10 mg/m3 | Inhalable | |
| Biological limit values | No biological exposure limits noted for the in | ngredient(s). | | |
| Recommended monitoring procedures | Follow standard monitoring procedures. | | | |
| Derived no-effect level (DNEL) | Not available. | | | |
| Predicted no effect concentrations (PNECs) | Not available. | | | |
| Exposure guidelines | | | | |
| 3.2. Exposure controls | | | | |
| Appropriate engineering controls | General ventilation normally adequate. An E operations involving this material based upo outcome of a site- or operation-specific risk a | n the OEL/Occupational | | |
| ndividual protection measures, | such as personal protective equipment | | | |
| General information | Personal protection equipment should be ch discussion with the supplier of the personal personal protective equipment (PPE) is used | protective equipment. For | | |
| Eye/face protection | If contact is likely, safety glasses with side shields are recommended. (e.g. EN 166). Not normally needed. | | | |
| Skin protection | | | | |
| - Hand protection | For prolonged or repeated skin contact use resistant protective gloves (EN 374) with a p normally needed. | | | |
| - Other | Wear suitable protective clothing as protection splashes, EN ISO 13982 for dust). Not norm | | contamination. (EN 14605 for | |
| Respiratory protection | When workers are facing concentrations abore certified respirators. Where breathable aeros gases/vapours of organic, inorganic, acid inc EN 14387). No personal respiratory protection | sols/dust are formed, us organic, alkaline compou | e suitable combination filter founds and toxic particles (eg. | |
| Thermal hazards | Wear appropriate thermal protective clothing | g, when necessary. | | |
| Hygiene measures | Always observe good personal hygiene mea and before eating, drinking, and/or smoking. equipment to remove contaminants. For adv from a qualified environment, health and saf | Routinely wash work c | lothing and protective | |
| Environmental exposure control | ls | | | |
| Hazard guidance and control recommendations | Environmental manager must be informed o | f all major releases. | | |

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

| Appearance | | | | |
|--|----------------|--|--|--|
| Physical state | Solid. | | | |
| Form | Tablet. | | | |
| Colour | Not available. | | | |
| Odour | Not available. | | | |
| Odour threshold | Not available. | | | |
| рН | Not available. | | | |
| Melting point/freezing point | Not available. | | | |
| Initial boiling point and boiling range | Not available. | | | |
| Flash point | Not available. | | | |
| Evaporation rate | Not available. | | | |
| Flammability (solid, gas) | Not available. | | | |
| Upper/lower flammability or explosive limits | | | | |
| Flammability limit - lower (%) | Not available. | | | |
| Flammability limit - upper (%) | Not available. | | | |

| Vapour pressure Not available. |
|---|
| Vapour density Not available. |
| Relative density Not available. |
| Solubility(ies) |
| Solubility (water) Not available. |
| Solubility (other) Not available. |
| Partition coefficientNot available.(n-octanol/water) |
| Auto-ignition temperature Not available. |
| Decomposition temperature Not available. |
| Viscosity Not available. |
| Explosive properties Not available. |
| Oxidizing properties Not available. |
| 9.2. Other information No relevant additional information available. |

SECTION 10: Stability and reactivity

| 10.1. Reactivity | The product is stable and non-reactive under normal conditions of use, storage and transport. |
|---|--|
| 10.2. Chemical stability | Material is stable under normal conditions. |
| 10.3. Possibility of hazardous reactions | No dangerous reaction known under conditions of normal use. |
| 10.4. Conditions to avoid | Contact with incompatible materials. |
| 10.5. Incompatible materials | Strong oxidising agents. |
| 10.6. Hazardous decomposition products | None known. Irritating and/or toxic fumes and gases may be emitted upon the product's decomposition. |

SECTION 11: Toxicological information

Occupational exposure to the substance or mixture may cause adverse effects.

Information on likely routes of exposure

General information

Acute toxicity

| Inhalation | Under normal conditions of intended use, this material is not expected to be an inhalation hazard. |
|--------------|--|
| Skin contact | Health injuries are not known or expected under normal use. |
| Eye contact | Health injuries are not known or expected under normal use. |
| Ingestion | Health injuries are not known or expected under normal use. May be harmful if swallowed. |
| Symptoms | Accidental exposure or contact might produce: nausea, vomiting, symptoms of hypersensitivity (such as skin rash, hives, itching, and difficulty breathing), increased susceptibility to infection. |

11.1. Information on toxicological effects

Expected to be a low hazard for usual industrial or commercial handling by trained personnel. May be harmful if swallowed.

| Components | Species | Test results |
|----------------------------|-------------------------------|--------------|
| AZATHIOPRINE (CAS 446 | -86-6) | |
| Acute | | |
| Oral | | |
| LD50 | Rat | 400 mg/kg |
| HYDROXYPROPYL METH | IYL CELLULOSE (CAS 9004-65-3) | |
| Acute | | |
| Oral | | |
| LD50 | Rat | > 2000 mg/kg |
| MAGNESIUM STEARATE | (CAS 557-04-0) | |
| Acute | | |
| Oral | | |
| LD50 | Rat | > 2000 mg/kg |
| Titanium dioxide (CAS 134 | 63-67-7) | |
| Acute | | |
| Inhalation | | |
| LC50 | Rat | 6820 mcg/m3 |
| Oral | | |
| LD50 | Rat | > 24 g/kg |
| Material name: IMLIRAN TAB | LETS | 2D2 |

| Components | Species | Test results |
|--|-------------------------------|--|
| Chronic | | |
| Inhalation | | |
| LOEC | Rat | 8.6 mg/m3, 1 years TiO2 accumulated in interstitial macrophages, aggregated interstitial cells and particle laden macrophrages in lymphoid tissue. |
| NOAEC | Rat | 250 mg/m3, 2 years Highest dose |
| | | 5 mg/m3, 24 months |
| Subacute | | |
| Inhalation | | |
| LOEL | Rat | 0.1 - 35 mg/m3, 4 weeks Mild macrophage hyperplasia, no change in bronchio-alveolar lavage fluid. |
| NOAEC | Guinea pig | 26 mg/m3, 3 weeks No evidence of significant inflammation in respiratory tract |
| Oral | | |
| NOAEL | Rat | 100000 ppm, 14 Day Dietary study, highes dose tested. |
| Subchronic | | |
| Inhalation | | |
| LOEC | Rat | 3.2 - 20 mg/m3, 8 min Accumulation of TiO2 in macrophages and evidence of pulmonary inflammation. |
| * Estimates for product may | be based on additional compon | ent data not shown. |
| Skin corrosion/irritation | Health injuries are not know | n or expected under normal use. |
| Irritation Corrosion - Skin | | |
| TITANIUM DIOXIDE | | 0, Literature data Result: Non-irritant Species: Guinea pig 0, Literature data Result: Non-irritant Species: Human Acute dermal irritation; OECD 404, Literature data Result: Non-irritant Species: Rabbit |
| Irritation Corrosion - Skin: MAGNESIUM STEARA | | 0 |
| Serious eye damage/eye irritation | Health injuries are not know | n or expected under normal use. |
| Еуе | | |
| TITANIUM DIOXIDE | | OECD 405, Literature data Result: Mild irritant Species: Rabbit |
| Eye / Kay and Calandra cla | | |
| MAGNESIUM STEARATI | TE | 4 Decevery Deried: 2 deve |
| | No studies have been condu | Recovery Period: 2 days |
| Respiratory sensitisation Skin sensitisation | May cause an allergic skin r | |
| | | |
| Maximisation assay (Magn HYDROXYPROPYL ME | | Result: negative Species: Guinea pig |
| Sensitisation TITANIUM DIOXIDE | | 5 % Optimisation Test, Literature data - Vehicle: petrolatum Result: negative Species: Guinea pig Test Duration: 48 hour exposure |
| AZATHIOPRINE | | Occupational exposure, Literature data Result: Low incidence of contact hypersensitivity. |
| TITANIUM DIOXIDE | | Species: Human Patch test, Literature data Result: negative Species: Human |
| Germ cell mutagenicity | May cause genetic defects. | · |

| Mutagenicity | | | |
|---|--|--|--|
| AZATHIOPRINE | | Ames Assay, GLP assay; Literature data Result: positive | |
| TITANIUM DIOXIDE | | Ames, Literature data Result: negative | |
| | | Micronucleus Assay in vitro, CHO cells, Literature data | |
| | | Result: negative Micronucleus Assay in vitro, cultured human peripheral lymphocytes, Literature data Result: positive | |
| AZATHIOPRINE | | Micronucleus Test, GLP assay; Literature data | |
| TITANIUM DIOXIDE | | Result: positive Syrian Hamster Embryo (SHE) cell transformation assay | |
| | | Result: negative WIL2-NS HPRT/ t-Thioguanidine - Human B-Cell lymphoblastoid, Literature data Result: positive | |
| Carcinogenicity | Contains a material (Azathiop | rine) classified as a carcinogen by external agencies. | |
| TITANIUM DIOXIDE | | 0.5 mg/m3, Literature data Result: negative | |
| | | Species: Rat Test Duration: 24 months | |
| | | 0.72 - 14.8 mg/m3, Literature data | |
| | | Result: negative Species: Mouse | |
| | | 10 - 250 mg/m3, Dietary study - Literature data. | |
| | | Result: Inflammation at all doses with alveolar/bronchiolar | |
| | | adenoma at the highest concentration. Species: Rat | |
| | | Test Duration: 24 months | |
| | | 25000 - 50000 ppm, Dietary study Result: negative | |
| | | Species: Mouse | |
| | | 25000 - 50000 ppm, Dietary study - Literature data. Result: negative Species: Rat | |
| | | 7.2 - 14.8 mg/m3, Literature data | |
| | | Result: Lung tumour Species: Rat | |
| | | Test Duration: 24 months | |
| AZATHIOPRINE | | Literature search Result: positive | |
| IARC Monographs. Overall I | Evaluation of Carcinogenicity | | |
| AZATHIOPRINE (CAS 44 | , | 1 Carcinogenic to humans. | |
| Titanium dioxide (CAS 13 Reproductive toxicity | May damage fertility or the un | 2B Possibly carcinogenic to humans. | |
| Reproductivity | may damage lentility of the dif | born child. | |
| AZATHIOPRINE | | Literature search Result: Positive for teratogenicity, fertiltiy effects and may affect the quality of breast milk. | |
| Specific target organ toxicity - single exposure | Not available. | | |
| Specific target organ toxicity - repeated exposure | Causes damage to organs thr | ough prolonged or repeated exposure. | |
| AZATHIOPRINE | | Literature search Organ: Immune system, Bone marrow and formation of blood cells, Liver, Kidney | |
| Aspiration hazard | Not likely, due to the form of the | he product. | |
| Mixture versus substance information | No information available. | | |
| Other information | Caution - Pharmaceutical agent. Occupational exposure to the substance or mixture may cause adverse effects. | | |
| SECTION 12: Ecological in | nformation | | |
| 12.1. Toxicity | Not expected to be harmful to | aquatic organisms. | |
| | | | |

| components | | Species | | Test results |
|--|--|--|---|---|
| ZATHIOPRINE (CAS 446-8 | 36-6) | | | |
| Acute | 1050 | A | | |
| | IC50 | Activated slude | ge | > 1000 mg/l, 3 hours |
| Aquatic | | | | |
| Acute | | Croop algae (| Coopedoomuo | > 100 mg/L 72 hours Static test |
| Algae | EC50 | Green algae (Scenedesmus subspicatus) | | > 100 mg/l, 72 hours Static test |
| | NOEC | Green algae (Scenedesmus subspicatus) | | 100 mg/l, 72 hours Static test |
| Crustacea | EC50 | Water flea (Daphnia magna) | | > 100 mg/l, 48 hours Static test |
| | NOEC Water flea (Daphnia | | phnia magna) | > 100 mg/l, 48 hours Static test |
| YDROXYPROPYL METHY | L CELLULOSE (CAS | S 9004-65-3) | | |
| Aquatic | , | , | | |
| <i>Acute</i> Fish | EC50 | Fish | | > 100 mg/l, 96 hours |
| - | | FISH | | |
| IAGNESIUM STEARATE (0 Aquatic | JAS 557-04-0) | | | |
| Acute | | _ | | |
| Fish | EC50 | Orange-red kill latipes) | lfish (Adult Oryzias | 130 mg/l, 96 hours |
| itanium dioxide (CAS 13463 | 3-67-7) | | | |
| Aquatic | | | | |
| Acute | EC50 | Water flea (Da | | > 1000 mg/l, 48 hours Static test |
| Crustacea * Estimates for product r 2.2. Persistence and egradability Photolysis | nay be based on ado | ditional componer | nt data not shown. | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA | is-atmospheric) ARATE | ditional componer | nt data not shown. 17 Hours Estimated | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi | is-atmospheric) ARATE m wavelength | ditional componer | | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability | is-atmospheric) ARATE m wavelength ARATE | | 17 Hours Estimated 210 nm | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio | is-atmospheric) ARATE m wavelength | | 17 Hours Estimated 210 nm nt) | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability | is-atmospheric) ARATE m wavelength ARATE | | 17 Hours Estimated 210 nm nt) 4 %, 28 days Modifier | d Zahn-Wellens, DOC removal., |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio | is-atmospheric) ARATE m wavelength ARATE | | 17 Hours Estimated 210 nm nt) 4 %, 28 days Modified Activated sludge 4 %, 28 days Modified | d Zahn-Wellens, primary |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodeg) | | 17 Hours Estimated 210 nm ht) 4 %, 28 days Modifier Activated sludge 4 %, 28 days Modifier biodegradation, loss of | |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodegi | radation-inheren | 17 Hours Estimated 210 nm nt) 4 %, 28 days Modified Activated sludge 4 %, 28 days Modified | d Zahn-Wellens, primary |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodeg) | radation-inheren | 17 Hours Estimated 210 nm ht) 4 %, 28 days Modifier Activated sludge 4 %, 28 days Modifier biodegradation, loss of | d Zahn-Wellens, primary of parent., Activated sludge |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE MAGNESIUM STEA Percent degradatio AZATHIOPRINE MAGNESIUM STEA | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodegi ARATE on (Aerobic biodegi | radation-inheren | 17 Hours Estimated 210 nm 1t) 4 %, 28 days Modifier Activated sludge 4 %, 28 days Modifier biodegradation, loss of 77 %, 28 days BOD | d Zahn-Wellens, primary of parent., Activated sludge ed Sturm test. |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE MAGNESIUM STEA Percent degradatio AZATHIOPRINE MAGNESIUM STEA | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodegi ARATE on (Aerobic biodegi ARATE on (Aerobic biodegi | radation-inheren | 17 Hours Estimated 210 nm 4 %, 28 days Modified Activated sludge 4 %, 28 days Modified biodegradation, loss of 77 %, 28 days BOD 11 %, 28 days Modified 95 %, 22 days Sturm | d Zahn-Wellens, primary of parent., Activated sludge ed Sturm test. |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE MAGNESIUM STEA Percent degradatio MAGNESIUM STEA | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodege ARATE on (Aerobic biodege ARATE on (Aerobic biodege ARATE | radation-inheren | 17 Hours Estimated 210 nm 4 %, 28 days Modifier Activated sludge 4 %, 28 days Modifier biodegradation, loss of 77 %, 28 days BOD 11 %, 28 days Modifier | d Zahn-Wellens, primary of parent., Activated sludge ed Sturm test. |
| * Estimates for product r 2.2. Persistence and egradability Photolysis Half-life (Photolysi MAGNESIUM STEA UV/visible spectru MAGNESIUM STEA Biodegradability Percent degradatio AZATHIOPRINE MAGNESIUM STEA Percent degradatio AZATHIOPRINE MAGNESIUM STEA Percent degradatio MAGNESIUM STEA | is-atmospheric) ARATE m wavelength ARATE on (Aerobic biodege ARATE on (Aerobic biodege ARATE on (Aerobic biodege ARATE | radation-inheren | 17 Hours Estimated 210 nm 4 %, 28 days Modified Activated sludge 4 %, 28 days Modified biodegradation, loss of 77 %, 28 days BOD 11 %, 28 days Modified 95 %, 22 days Sturm | d Zahn-Wellens, primary of parent., Activated sludge ed Sturm test. |
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Volatility Henry's law AZATHIOPRINE HYDROXYPROPYL METHYL CELLULOSE

12.5. Results of PBT Not available. and vPvB assessment

12.6. Other adverse effects Not available.

SECTION 13: Disposal considerations

13.1. Waste treatment methods Residual waste Dispose of in accordance with local regulations. Empty containers or liners may retain some product residues. This material and its container must be disposed of in a safe manner (see: Disposal instructions). Avoid discharge into water courses or onto the ground. Empty containers should be taken to an approved waste handling site for recycling or disposal. Contaminated packaging Since emptied containers may retain product residue, follow label warnings even after container is emptied. The Waste code should be assigned in discussion between the user, the producer and the waste EU waste code disposal company. **Disposal methods/information** Collect and reclaim or dispose in sealed containers at licensed waste disposal site. Do not discharge into drains, water courses or onto the ground. Dispose in accordance with all applicable regulations. **Special precautions** Dispose in accordance with all applicable regulations.

0 atm m³/mol. 25 C Estimated

0 atm m3/mol Estimated

SECTION 14: Transport information

ADR

Not regulated as dangerous goods.

ΙΑΤΑ

Not regulated as dangerous goods.

IMDG

Not regulated as dangerous goods.

14.7. Transport in bulkMARPOL Annex II applies to liquids used in a ship's operation that pose a threat to the marine
environment. These materials may not be transported in bulk.**MARPOL73/78 and the IBC Code**

SECTION 15: Regulatory information

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

EU regulations

Regulation (EC) No. 1005/2009 on substances that deplete the ozone layer, Annex I Not listed.

Regulation (EC) No. 1005/2009 on substances that deplete the ozone layer, Annex II Not listed.

Regulation (EC) No. 850/2004 On persistent organic pollutants, Annex I as amended

Not listed

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 1 as amended Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 2 as amended Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 3 as amended Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex V as amended Not listed.

Regulation (EC) No. 166/2006 Annex II Pollutant Release and Transfer Registry

Not listed.

Regulation (EC) No. 1907/2006, REACH Article 59(10) Candidate List as currently published by ECHA Not listed.

Authorisations

Regulation (EC) No. 1907/2006, REACH Annex XIV Substances subject to authorization, as amended Not listed.

Restrictions on use

Regulation (EC) No. 1907/2006, REACH Annex XVII Substances subject to restriction on marketing and use as amended Not listed.

Directive 2004/37/EC: on the protection of workers from the risks related to exposure to carcinogens and mutagens at work

Not listed.

Directive 92/85/EEC: on the safety and health of pregnant workers and workers who have recently given birth or are breastfeeding

Not listed.

Other EU regulations

Directive 96/82/EC (Seveso II) on the control of major-accident hazards involving dangerous substances Not listed.

Directive 98/24/EC on the protection of the health and safety of workers from the risks related to chemical agents at work Not listed.

Directive 94/33/EC on the protection of young people at work

Not listed.

Other regulations

The product is classified and labelled in accordance with EC directives or respective national laws. This Safety Data Sheet complies with the requirements of Regulation (EC) No 1907/2006.

| National regulations | Young people under 18 years old are not allowed to work with this product according to the EU Directive 94/33/EC on the protection of young people at work. Follow national regulation for work with chemical agents. |
|-----------------------|---|
| 15.2. Chemical safety | No Chemical Safety Assessment has been carried out. |

assessment

SECTION 16: Other information

| List of abbreviations | Not available. |
|--|---|
| References | GSK Hazard Determination |
| Information on evaluation method leading to the classification of mixture | The classification for health and environmental hazards is derived by a combination of calculation methods and test data, if available. |
| Full text of any statements or R-phrases and H-statements under Sections 2 to 15 | R22 Harmful if swallowed. R36 Irritating to eyes. R37 Irritating to respiratory system. R38 Irritating to skin. R43 May cause sensitization by skin contact. R45 May cause cancer. R46 May cause heritable genetic damage. R48 Danger of serious damage to health by prolonged exposure. R61 May cause harm to the unborn child. H302 Harmful if swallowed. H315 Causes skin irritation. H317 May cause an allergic skin reaction. H319 Causes serious eye irritation. H335 May cause genetic defects. H360 May cause cancer. H360 May damage fortility or the unborn child. H372 Causes damage to organs through prolonged or repeated exposure. |
| Revision information | Product and Company Identification: Product and Company Identification Composition / Information on Ingredients: Ingredients Exposure Controls / Personal Protection: Physical & Chemical Properties: Ecological Information: Ecotoxicity Transport Information: Agency Name and Packaging Type/Transport Mode Selection Regulatory Information: Risk Phrases - Class. GHS: Classification |
| Training information | Follow training instructions when handling this material. |
| Disclaimer | The information and recommendations in this safety data sheet are, to the best of our knowledge, accurate as of the date of issue. Nothing herein shall be deemed to create any warranty, express or implied. It is the responsibility of the user to determine the applicability of this information and the suitability of the material or product for any particular purpose. |