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

ZOFRAN TABLETS

Registration number

ZOFRAN TABLETS 4 MG * ZOFRAN TABLETS 8 MG * ONDANSETRON HYDROCHLORIDE **Synonyms**

TABLETS * ONDANSETRON HYDROCHLORIDE DIHYDRATE, FORMULATED PRODUCT

Issue date 11-August-2014

Version number

Revision date 11-August-2014

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

Identified uses Medicinal Product

> This safety data sheet is written to provide health, safety and environmental information for people handling this formulated product in the workplace. It is not intended to provide information relevant

to medicinal use of the product. In this instance patients should consult prescribing

information/package insert/product label or consult their pharmacist or physician. For health and safety information for individual ingredients used during manufacturing, refer to the appropriate

safety data sheet for each ingredient.

Uses advised against No other uses are advised.

1.3. Details of the supplier of the safety data sheet

GlaxoSmithKline UK 980 Great West Road

Brentford, Middlesex TW8 9GS UK

UK General Information (normal business hours): +44-20-8047-5000

Fmail Address: msds@gsk.com Website: www.qsk.com

1.4. Emergency telephone

number

TRANSPORT EMERGENCIES::

UK In-country toll call: +(44)-870-8200418 International toll call: +1 703 527 3887

available 24 hrs/7 days; multi-language response

SECTION 2: Hazards identification

2.1. Classification of the substance 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.

Supplemental label information None.

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

Material name: ZOFRAN TABLETS

General information

Chemical name % CAS-No. / EC No. REACH Registration No. INDEX No. Notes

MICROCRYSTALLINE CELLULOSE 20 - < 30 9004-34-6

232-674-9

Classification: DSD: -

CLP: -

OPASPRAY M-1-8429 5 - < 10 Unassigned -

Classification: DSD: R10, Xn;R68/20/21/22

CLP: Flam. Liq. 2;H225, STOT SE 2;H371

HYDROXYPROPYL METHYL CELLULOSE

3 - < 5 9004-65-3

Classification: DSD: -

CLP: -

ONDANSETRON HYDROCHLORIDE 3 - < 5 103639-04-9

DIHYDRATE

Classification: DSD: T;R25, Xi;R41, N;R50/53

CLP: Acute Tox. 3;H301, Eye Dam. 1;H318, Aquatic Acute 1;H400, Aquatic

Chronic 2;H411

Starch 3 - < 5 9005-25-8 - -

232-679-6

Classification: DSD: -

CLP: -

MAGNESIUM STEARATE < 1 557-04-0 -

209-150-3

Classification: DSD: -

CLP: -

Other components below reportable levels 50 - < 60

CLP: Regulation No. 1272/2008.

DSD: Directive 67/548/EEC.

M: M-factor

vPvB: very persistent and very bioaccumulative substance.

PBT: persistent, bioaccumulative and toxic substance.

#: This substance has been assigned Community workplace exposure limit(s).

Composition comments The full text for all R- and H-phrases is displayed in section 16.

SECTION 4: First aid measures

General information In the case of accident or if you feel unwell, seek medical advice immediately (show the label

where possible).

4.1. Description of first aid measures

Inhalation If breathing is difficult, trained personnel should give oxygen.

Skin contact Immediately flush skin with plenty of water. Take off contaminated clothing and wash before reuse.

Get medical attention if symptoms occur.

Eye contact Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.

Ingestion If swallowed, rinse mouth with water (only if the person is conscious). Do not induce vomiting

without medical advice. If ingestion of a large amount does occur, call a poison control centre immediately.

immediate

4.2. Most important symptoms and effects, both acute and delayed

Material name: ZOFRAN TABLETS

Direct contact with eyes may cause temporary irritation. Exposed may experience eye tearing,

redness, and discomfort.

4.3. Indication of any immediate medical attention and special treatment neededNo specific antidotes are recommended. Treat according to locally accepted protocols. For additional guidance, refer to the current prescribing information or to the local poison control information centre.

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. Use water spray to cool unopened

containers.

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. Wear appropriate personal protective equipment. 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

6.2. Environmental precautions

Avoid release to the environment. Contact local authorities in case of spillage to drain/aquatic environment. Prevent further leakage or spillage if safe to do so. Do not contaminate water. 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. Collect spillage. Prevent product from entering drains. Following product recovery, flush area with water.

6.4. Reference to other

For personal protection, see section 8. For waste disposal, see section 13.

sections

SECTION 7: Handling and storage

7.1. Precautions for safe

handling

Avoid prolonged exposure. Provide adequate ventilation. Wear appropriate personal protective equipment. Observe good industrial hygiene practices. Avoid release to the environment. Do not empty into drains.

7.2. Conditions for safe storage, including any incompatibilities

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 |
|-----|
| Com |

| Components | Туре | Value | |
|--|----------|-----------|--|
| HYDROXYPROPYL METHYL CELLULOSE (CAS 9004-65-3) | OHC | 1 | |
| MAGNESIUM STEARATE (CAS 557-04-0) | OHC | 1 | |
| MICROCRYSTALLINE CELLULOSE (CAS 9004-34-6) | OHC | 1 | |
| ONDANSETRON HYDROCHLORIDE DIHYDRATE (CAS 103639-04-9) | 8 HR TWA | 30 mcg/m3 | |
| • | OHC | 3 | |

Biological limit values

No biological exposure limits noted for the ingredient(s).

Recommended monitoring

procedures

Follow standard monitoring procedures.

Derived no-effect level (DNEL) Not available.

Material name: ZOFRAN TABLETS SDS MALTA 110606 Version No.: 11 Revision date: 11-August-2014 Issue date: 11-August-2014

Predicted no effect concentrations (PNECs) Not available.

8.2. Exposure controls

Appropriate engineering

controls

Good general ventilation (typically 10 air changes per hour) should be used. Ventilation rates should be matched to conditions. If applicable, use process enclosures, local exhaust ventilation, or other engineering controls to maintain airborne levels below recommended exposure limits. If exposure limits have not been established, maintain airborne levels to an acceptable level.

Individual protection measures, such as personal protective equipment

Follow all local regulations if personal protective equipment (PPE) is used in the workplace. **General information**

Personal protection equipment should be chosen according to the CEN standards and in

discussion with the supplier of the personal protective equipment.

If contact is likely, safety glasses with side shields are recommended. (eg. EN 166) Eye/face protection

Skin protection

- Hand protection For prolonged or repeated skin contact use suitable protective gloves. Select suitable chemical

resistant protective gloves (EN 374) with a protective index 6 (>480min permeation time).

Wear suitable protective clothing as protection against splashing or contamination. (EN 14605 for - Other

splashes, EN ISO 13982 for dust)

Where breathable aerosols/dust are formed, use suitable combination filter for gases/vapours of Respiratory protection organic, inorganic, acid inorganic, alkaline compounds and toxic particles (eg. EN 14387). When

workers are facing concentrations above the exposure limit they must use appropriate certified

respirators.

Thermal hazards Wear appropriate thermal protective clothing, when necessary.

Always observe good personal hygiene measures, such as washing after handling the material Hygiene measures

and before eating, drinking, and/or smoking. Routinely wash work clothing and protective

equipment to remove contaminants.

Environmental exposure controls

Hazard guidance and control recommendations Contain spills and prevent releases and observe national regulations on emissions. Environmental

manager must be informed of all major releases.

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Appearance

Physical state Solid. **Form** Tablet.

Not available. Colour Odour Not available. **Odour threshold** Not available. Not available. pН Not available. Melting point/freezing point Not available. Initial boiling point and boiling

range

Flash point

Evaporation rate

Not available. Not available.

Flammability (solid, gas) Not available.

Upper/lower flammability or explosive limits

Flammability limit - lower

Not available.

Flammability limit - upper

Not available.

(%)

Not available. Vapour pressure Vapour density Not available. Relative density Not available.

Solubility(ies)

Not available. Solubility (water) Not available Solubility (other) Partition coefficient Not available.

(n-octanol/water)

Auto-ignition temperature Not available.

Material name: ZOFRAN TABLETS

Decomposition temperatureNot available.ViscosityNot available.Explosive propertiesNot available.Oxidizing propertiesNot available.

9.2. Other information No relevant additional information available.

SECTION 10: Stability and reactivity

10.1. ReactivityThe 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. Fluorine.

10.6. Hazardous Irritating and/or toxic fumes and gases may be emitted upon the products decomposition.

decomposition products

SECTION 11: Toxicological information

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

Information on likely routes of exposure

Ingestion Health injuries are not known or expected under normal use. Expected to be a low ingestion

hazard. However, ingestion is not likely to be a primary route of occupational exposure.

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.

Symptoms Exposure may cause temporary irritation, redness, or discomfort.

11.1. Information on toxicological effects

Acute toxicity Health injuries are not known or expected under normal use.

Components Species Test results

HYDROXYPROPYL METHYL 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

MICROCRYSTALLINE CELLULOSE (CAS 9004-34-6)

Acute

Dermal

LD50 Rabbit > 2000 mg/kg

Oral

LD50 Rat > 2000 mg/kg

ONDANSETRON HYDROCHLORIDE DIHYDRATE (CAS 103639-04-9)

Acute

Oral

LD50 Rat 100 - 150 mg/kg

Chronic

Oral

LD Rat > 36 mg/kg/day

 LOEL
 Dog
 1 mg/kg/day, 52 weeks

 NOAEL
 Rat
 1 mg/kg/day, 18 months

Skin corrosion/irritation Based on available data, the classification criteria are not met.

Material name: ZOFRAN TABLETS SDS MALTA

^{*} Estimates for product may be based on additional component data not shown.

Corrosivity

ONDANSETRON HYDROCHLORIDE DIHYDRATE 50 %, formulated in soft paraffin.

Result: Non-irritant Species: Guinea pig

Irritation Corrosion - Skin: P.I.I. value

MAGNESIUM STEARATE 0

Serious eye damage/eye Based on available data, the classification criteria are not met.

irritation

Eye

ONDANSETRON HYDROCHLORIDE DIHYDRATE **OFCD 405**

Result: Severe Irritant

Species: Rabbit

Eye / Kay and Calandra class - Intact

MAGNESIUM STEARATE

Recovery Period: 2 days

Respiratory sensitisation Not available.

Skin sensitisation Based on available data, the classification criteria are not met.

Maximisation assay (Magnusson and Kligman)

HYDROXYPROPYL METHYL CELLULOSE Result: negative

Species: Guinea pig

Sensitisation

ONDANSETRON HYDROCHLORIDE DIHYDRATE Split adjuvant assay

> Result: negative Species: Guinea pig

Germ cell mutagenicity Based on available data, the classification criteria are not met.

Mutagenicity

ONDANSETRON HYDROCHLORIDE DIHYDRATE Ames

Result: negative

Chromosomal Aberration Assay In Vitro

Result: positive

HPRT gene mutation in human lymphocytes

Result: negative Micronucleus test Result: negative Species: Mouse

V79 Cell Mutagenicity Assay

Result: negative

Due to partial or complete lack of data the classification is not possible. Carcinogenicity

ONDANSETRON HYDROCHLORIDE DIHYDRATE ICH S1B

Result: negative Species: Mouse ICH S1B Result: negative Species: Rat

Reproductive toxicity Based on available data, the classification criteria are not met.

Reproductivity

ONDANSETRON HYDROCHLORIDE DIHYDRATE **Embryofetal Development**

> Result: No effect Species: Rabbit

Embryofetal Development

Result: No effect Species: Rat Fertility Result: No effect

Species: Rat

Pre- and Post-natal development

Result: negative Species: Rat

Specific target organ toxicity -

single exposure

None known.

Specific target organ toxicity -

repeated exposure

None known.

Not likely, due to the form of the product. **Aspiration hazard**

Mixture versus substance

information

No information available.

Other information

Not available.

Material name: ZOFRAN TABLETS 110606 Version No.: 11 Revision date: 11-August-2014 Issue date: 11-August-2014

SECTION 12: Ecological information

12.1. Toxicity Contains a substance which causes risk of hazardous effects to the environment.

Components Species Test results

HYDROXYPROPYL METHYL CELLULOSE (CAS 9004-65-3)

Aquatic

Acute

Fish EC50 Fish > 100 mg/l, 96 hours

MAGNESIUM STEARATE (CAS 557-04-0)

Aquatic

Acute

Fish EC50 Orange-red killfish (Adult Oryzias 130 mg/l, 96 hours

latipes)

Microtox EC50 Microtox 12,5 mg/l, 15 minutes

ONDANSETRON HYDROCHLORIDE DIHYDRATE (CAS 103639-04-9)

Aquatic

Acute

Activated Sludge IC50 Residential sludge > 1000 mg/l, 3 hours OECD 209 Respiration

Algae EC50 Green algae (Selenastrum 0,87 mg/l, 72 hours Measured, OECD

capricornutum)

NOEC Green algae (Selenastrum 0,31 mg/l, 72 hours Static test

capricornutum)

Crustacea EC50 Water flea (Daphnia pulex) 28 mg/l, 48 hours Static test, TAD 4.08

NOEC Water flea (Daphnia pulex) 16 mg/l, 48 hours Static test

Fish EC50 Rainbow trout (Adult Oncorhyncus 6,5 mg/l, 96 hours Static test, OECD 203

mykiss)

NOEC Rainbow trout (Adult Oncorhyncus 2,6 mg/l, 96 hours Measured

mykiss)

Chronic

Crustacea EC50 Water flea (Ceriodaphnia dubia) 1,4 mg/l, 8 days Static renewal test, EPA

1002

LOEC Water flea (Ceriodaphnia dubia) 1 mg/l, 8 days

NOEC Water flea (Ceriodaphnia dubia) 0,32 mg/l, 8 days

12.2. Persistence and degradability

No data is available on the degradability of this product.

Photolysis

Half-life (Photolysis-atmospheric)

MAGNESIUM STEARATE 17 Hours Estimated

UV/visible spectrum wavelength

MAGNESIUM STEARATE 210 nm

ONDANSETRON HYDROCHLORIDE DIHYDRATE 305 nm, pH 5-9

Hydrolysis

Half-life (Hydrolysis-neutral)

ONDANSETRON HYDROCHLORIDE DIHYDRATE > 1 years

Biodegradability

Percent degradation (Aerobic biodegradation-inherent)

MAGNESIUM STEARATE 77 %, 28 days BOD

ONDANSETRON HYDROCHLORIDE DIHYDRATE 18,9 %, 28 days Semi-continuous activated sludge (SCAS),

Activated sludge

Percent degradation (Aerobic biodegradation-ready)

MAGNESIUM STEARATE 95 %, 22 days Sturm test

Percent degradation (Aerobic biodegradation-soil)

MAGNESIUM STEARATE 50 %, 13 days

ONDANSETRON HYDROCHLORIDE DIHYDRATE 20,3 - 99,9 %, 64 days, Soil

12.3. Bioaccumulative potential No data available for this product.

Partition coefficient n-octanol/water (log Kow)

HYDROXYPROPYL METHYL CELLULOSE -5
ONDANSETRON HYDROCHLORIDE DIHYDRATE 0.995

Material name: ZOFRAN TABLETS

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Bioconcentration factor (BCF)

HYDROXYPROPYL METHYL CELLULOSE 3,2 Estimated
MAGNESIUM STEARATE > 9999 Estimated

12.4. Mobility in soil No data available.

Adsorption

Sludge/biomass distribution coefficient - log Kd

ONDANSETRON HYDROCHLORIDE DIHYDRATE 3,95 - 4,23 Calculated

Soil/sediment sorption - log Koc

MAGNESIUM STEARATE 5,86 Estimated

ONDANSETRON HYDROCHLORIDE DIHYDRATE 4,22 - 4,51 Measured

Mobility in general Not available.

Volatility

Henry's law

HYDROXYPROPYL METHYL CELLULOSE 0 atm m3/mol Estimated

Distribution

Octanol/water distribution coefficient log DOW

ONDANSETRON HYDROCHLORIDE DIHYDRATE 0,23, pH 5

0,99, pH 7 1,26, pH 9

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).

Contaminated packaging Empty containers should be taken to an approved waste handling site for recycling or disposal.

Since emptied containers may retain product residue, follow label warnings even after container is

emptied.

EU waste codeThe Waste code should be assigned in discussion between the user, the producer and the waste

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 precautionsDispose in accordance with all applicable regulations.

SECTION 14: Transport information

General Classifications are for the material when offered for transport as fully regulated. Depending on

the specific transport details (Ship-From/Ship To locations, quantities being shipped, type of packaging and mode of transport) it may be possible to ship this material in a manner other than fully regulated. (One example is IATA Limited or Excepted Quantity. There are others.) Be sure to review all regulatory agency packaging instructions and special provisions, referenced in this

section, to identify options applicable to the specifics of your shipment.

ADR

14.1. UN number UN3077

14.2. UN proper shipping Environmentally hazardous substances, solid, n.o.s. (ONDANSETRON HYDROCHLORIDE

name TABLETS)

14.3. Transport hazard class(es)

Class 9 Subsidiary risk -Label(s) 9

Hazard No. (ADR) Not available.

Tunnel code Not available.

14.4. Packing group III **14.5. Environmental hazards** No.

14.6. Special precautions Not available.

for user

IATA

14.1. UN number UN3077

14.2. UN proper shipping Environmentally hazardous substance, solid, n.o.s. (ONDANSETRON HYDROCHLORIDE

name TABLETS)

Material name: ZOFRAN TABLETS

SDS MALTA

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14.3. Transport hazard 9

class(es)

Subsidiary class(es) Ш 14.4. Packing group Labels required 9 14.5. Environmental hazards No. 9L **ERG Code**

14.6. Special precautions

for user

Not available.

Other information

Allowed. Cargo aircraft only

Additional Information:

Passenger & cargo Allowed.

IMDG

14.1. UN number UN3077

14.2. UN proper shipping ENVIRONMENTALLY HAZARDOUS SUBSTANCE, SOLID, N.O.S. (ONDANSETRON

HYDROCHLORIDE TABLETS)

14.3. Transport hazard class(es)

Class Subsidiary risk 9 Label(s) 14.4. Packing group Ш 14.5. Environmental hazards

Marine pollutant F-A, S-F Not available. 14.6. Special precautions

for user

14.7. Transport in bulk according to Annex II of

MARPOL73/78 and the IBC Code

MARPOL 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.

ADR; IATA; IMDG



Marine pollutant



General information

Classifications are for the material when offered for transport as fully regulated. Depending on the specific transport details (Ship-From/Ship To locations, quantities being shipped, type of packaging and mode of transport) it may be possible to ship this material in a manner other than fully regulated. (One example is IATA Limited or Excepted Quantity. There are others.) Be sure to review all regulatory agency packaging instructions and special provisions, referenced in this section, to identify options applicable to the specifics of your shipment.

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.

Material name: ZOFRAN TABLETS SDS MALTA Regulation (EC) No. 1005/2009 on substances that deplete the ozone layer, Annex II

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

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 3 as amended

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(1) 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

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

Follow national regulation for work with chemical agents. No Chemical Safety Assessment has been carried out.

15.2. Chemical safety

assessment

List of abbreviations

SECTION 16: Other information

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

R10 Flammable.

R22 Harmful if swallowed. R25 Toxic if swallowed.

R41 Risk of serious damage to eyes.

R50/53 Very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic

environment.

R51/53 Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic

environment.

R68/20/21/22 Harmful: possible risk of irreversible effects through inhalation, in contact with skin

and if swallowed.

H225 Highly flammable liquid and vapour.

H301 Toxic if swallowed.

H318 Causes serious eye damage.

Material name: ZOFRAN TABLETS SDS MALTA H371 May cause damage to organs. H400 Very toxic to aquatic life.

H411 Toxic to aquatic life with long lasting effects. May form combustible dust concentrations in air.

Revision information Product and Company Identification: Product and Company Identification

Composition / Information on Ingredients: Ingredients

Physical & Chemical Properties:

Toxicological Information: Toxicological Property Data

Ecological Information: Ecotoxicity

Transport Information: Material Transportation Information

Regulatory Information: United States

Material Attributes & Uses; Experimental Data: Material Uses

GHS: Classification

Consumer Products: CPSC Hazard Categories

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.

Material name: ZOFRAN TABLETS