

**Safety Data Sheet** 

# **CELLCEPT(R) Oral Suspension**

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

#### 1.1. Product identifier

Product name CELLCEPT(R) Oral Suspension

Product code CSE-3173

Synonyms - CELLCEPT Reconstituted Suspension (200mg/ml)

- CELLCEPT Oral Suspension (200mg/ml)

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

Use - pharmaceutical active substance (immunosuppressant) \*1

1.3. Details of the supplier of the safety data sheet

Company information Enquiries: Local representation:

Genentech, Inc. 1 DNA Way

South San Francisco USA-CA 94080

United States of America

Phone 001-(650) 225-1000 E-Mail info.sds@roche.com

US Chemtrec phone:

(800)-424-9300

1.4. Emergency telephone number

Emergency telephone number US Chemtrec phone: (800)-424-9300

\*1 referring to: Mycophenolate mofetil

## **SECTION 2: Hazards identification**

#### Classification of the substance or mixture / Label elements

**GHS** Classification

Health Hazards:

3.5Germ cell mutagenicity (Category 2)

H341 Suspected of causing genetic defects.

3.7 Reproductive toxicity (Category 1B) H360D May damage the unborn child.

3.9 Specific target organ toxicity - Repeated exposure (Category

H372 Causes damage to organs through prolonged or

repeated exposure.

Signalword: Danger

Label:



Precautionary statements:

- P201 Obtain special instructions before use.
- P273 Avoid release to the environment.
- P281 Use personal protective equipment as required.
- P309 + P311 IF exposed or you feel unwell: Call a POISON CENTER or doctor/physician.

#### Other hazards

Note - no information available

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

Characterization Mycophenolate mofetil and other inactive ingredients

Ingredients Concentration **GHS-Classification** 

(pure ingredient)

Mycophenolate mofetil 17.2 % - Combustible dust (No category), USH003 128794-94-5 - Acute toxicity (Category 4), H302

- Germ cell mutagenicity (Category 2), H341 - Reproductive toxicity (Category 1B), H360D - Specific target organ toxicity - Repeated

exposure (Category 1), H372

For the full text of the H-phrases mentioned in this Section, see Section 16.

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## **SECTION 4: First aid measures**

## 4.1. Description of first aid measures

Eye contact - rinse immediately with tap water for 10 minutes - open eyelids

forcibly

- consult a physician

Skin contact - remove immediately contaminated clothes, wash affected skin

with water and soap - do not use any solvents

Inhalation - remove the casualty to fresh air and keep him/her calm

- get medical treatment

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

Note - no information available

#### 4.3. Indication of any immediate medical attention and special treatment needed

Note to physician - treat symptomatically

- after accidental exposure women should get medical advice from

a physician

## **SECTION 5: Firefighting measures**

#### 5.1. Extinguishing media

Suitable extinguishing media - water spray jet, dry powder, foam, carbon dioxide

Flash point (liquid) not applicable

#### 5.2. Special hazards arising from the substance or mixture

Specific hazards - formation of toxic and corrosive combustion gases (nitrogen

oxides (NOx)) possible

- consider danger for the environment: dike spilled liquid

#### 5.3. Advice for firefighters

Protection of fire-fighters - precipitate gases/vapours/mists with water spray

Special method of fire-fighting - for reasons of environmental protection hold the extinguishing

agent back

#### **SECTION 6: Accidental release measures**

#### 6.1. Personal precautions, protective equipment and emergency procedures

Personal precautions - ensure adequate ventilation

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#### 6.2. Environmental precautions

Environmental protection - if the substance reaches waters or the sewer system, inform the

competent authority

 dilute the leaked substance by a water spray jet as far as necessary in order to minimize the hazard; hold the drainig water/mixture of substances back by all means available

#### 6.3. Methods and material for containment and cleaning up

Methods for cleaning up - collect spilled solutions with inert adsorbent and hand over to

waste removal

## **SECTION 7: Handling and storage**

## 7.1. Precautions for safe handling

Suitable materials - high density polyethylene (HDPE), glass

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

Storage conditions - room temperature

- do not freeze

Validity - ≤60 days, 25 °C

Packaging materials - high density polyethylene (HDPE) bottles with a child-resistant

polypropylene screw cap

## **SECTION 8: Exposure controls/personal protection**

#### 8.1. Control parameters

Threshold value (Roche) air - IOEL (Internal Occupational Exposure Limit): 0.01 mg/m3 \*1

8.2. Exposure controls

General protective and hygiene measures

- instruction of employees mandatory

Respiratory protection - Respiratory protection is recommended as a precaution to

minimize exposure. Effective engineering controls are considered to be the primary means to control worker exposure. Respiratory protection should not substitute for feasible engineering controls.

- in case of open handling or accidental release: particle mask or respirator with independent air supply

Hand protection - protective gloves (neoprene, nitrile or butyl rubber)

Eye protection - safety glasses

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Body protection - protective clothing

\*1 referring to: Mycophenolate mofetil

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

## 9.1. Information on basic physical and chemical properties

Color white to off-white

Form aqueous suspension

Solubility  $\leq$  22 mg/l, water (~ 22 °C, pH 6.3, HPLC, 24 h) \*1

≤36 mg/l, aquatic ecotoxicity media (~ 22 °C, HPLC, 24 h)

Partition coefficient log P<sub>ow</sub> 1.45 (n-octanol/water) pH 7.4

9.2. Other information

Note - no information available

\*1 referring to: Mycophenolate mofetil

## **SECTION 10: Stability and reactivity**

## 10.1. Reactivity

Note - no information available

10.2. Chemical stability

Stability - stable under the conditions mentioned in chapter 7

## 10.3. Possibility of hazardous reactions

Note - no information available

10.4. Conditions to avoid

Note - no information available

10.5. Incompatible materials

Materials to avoid - strong oxidizing agents \*1

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## 10.6. Hazardous decomposition products

Note - no information available

\*1 referring to: Mycophenolate mofetil

# **SECTION 11: Toxicological information**

## 11.1. Information on toxicological effects

Acute toxicity	<ul> <li>LD<sub>50</sub> 353 mg/kg (oral, rat); females, "trimmed Spearman-Karber method"</li> <li>LD<sub>50</sub> 500 mg/kg (oral, rat); males, "trimmed Spearman-Karber method"</li> <li>LD<sub>0</sub> 250 mg/kg (oral, rat)</li> <li>LD<sub>50</sub> &gt; 4'000 mg/kg (oral, mouse)</li> </ul>	*1 *1 *1 *1			
Local effects	<ul><li>skin: non-irritant</li><li>eye: non-irritant</li></ul>	*1 *1			
Sensitization	- non-sensitizing	*1			
Mutagenicity	<ul> <li>mutagenic (OECD No. 474 (Micronucleus Test)); threshold dose response</li> <li>mutagenic (mouse lymphoma cell mutation test; OECD No. 476 (Mammalian Cell Gene Mutation Test))</li> <li>does not induce chromosomal aberrations in vitro (OECD No. 473 (Mammalian Cytogenic Test))</li> <li>not mutagenic (Ames test)</li> </ul>				
Carcinogenicity	- not carcinogenic (several species)	*1			
Reproductive toxicity	- teratogenic (several species)	*1			
Note	<ul> <li>immunosuppressive agent</li> <li>average therapeutic dose 1 g twice daily</li> <li>elimination half-life: 16-18 h</li> <li>adverse effects at therapeutic dose: diarrhea, drop in white blookedle count, susceptibility to infections, vomiting</li> <li>high doses may irritate the digestive tract</li> <li>excretion predominantly renal</li> </ul>	*1 *1 *1 d *1 *1 *1			
Potential Health Effects	- Exposure: Inhalation, Ingestion, Skin contact, Eye contact				
	- Carcinogenicity: not listed by NTP, IARC or OSHA				
*1 referring to: *2 referring to:	Mycophenolate mofetil Mycophenolic acid				

## **SECTION 12: Ecological information**

#### 12.1. Toxicity

**Ecotoxicity** 

 barely toxic for planktonic crustaceans (nominal concentration = 100 mg/l), test performed with water accommodated fractions (Daphnia magna)

 $EC_{50}$  (48 h) > 100 mg/l (nominal concentration)

NOEC (48 h) 27.7 mg/l (average measured concentration) (OECD No. 202)

\*1

\*1

\*1

\*1

\*1

\*1

\*1

\*1

highly toxic for algae (Scenedesmus (=Desmodesmus) subspicatus)

 $\rm ErC_{50}$  (72 h)  $\,$  0.6 mg/l (average measured concentration)  $\rm EbC_{50}$  (72 h)  $\,$  0.2 mg/l (average measured concentration)

NOEC (72 h) 0.1 mg/l (nominal concentration) (OECD No. 201)

 adaptation/recovery of organisms upon prolongation of test duration (Scenedesmus (=Desmodesmus) subspicatus)
 LOEC (14 d) 1.6 mg/l (nominal concentration)

(OECD No. 201, prolonged)

acute fish toxicity in a limit test is lower than daphnid or algal toxicity, hence not relevant for classification (guppy)

NOEC (96 h) 1.7 mg/l (highest tested concentration)

(OECD No. 203)

no adverse influence on substrate biodegradation (activated sludge)

concentration (14 d) 100 mg/l (nominal concentration) (Manometric Respirometry Test, OECD No. 301 F)

#### 12.2. Persistence and degradability

Ready biodegradability

- not readily biodegradable

~ 14 %, 64 d

(FDA Technical Assistance Document No. 3.11)

- not readily biodegradable

primary degradation evidenced by HPLC

< 6 %, 28 d

(Manometric Respirometry Test, OECD No. 301 F)

Inherent biodegradability

- evidence for medium-term biodegradation in surface waters

(analogous to OECD 308, Transformation in natural

Abiotic degradation - rapid degradation, photodegradation, hydrolysis 22.3 mg/l, water;

water/sediment systems)

HPLC

~ 37 %, 120 h, ~ 22 °C, dark

~ 67 %, 120 h, ~ 22 °C, under illumination

#### 12.3. Bioaccumulative potential

Note - no information available

#### 12.4. Mobility in soil

Note - no information available

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#### 12.5. Results of PBT and vPvB assessment

Note - no information available

#### 12.6. Other adverse effects

Note - no information available

\*1 referring to: Mycophenolate mofetil

## **SECTION 13: Disposal considerations**

#### 13.1. Waste treatment methods

Waste from residues

- return to supplier or hand over to authorized disposal company
- observe local/national regulations regarding waste disposal
- incinerate in qualified installation with flue gas scrubbing
- DO NOT FLUSH unused medications or POUR them down a sink or drain. If available in your area, use takeback programs run by household hazardous waste collection programs or community pharmacies to dispose of unused and expired medicines. If you don't have access to a takeback program, dispose of these medicines in the household trash by removing them from their original containers and mixing them with an undesirable substance, such as used coffee grounds or kitty litter.

## **SECTION 14: Transport information**

IATA	Class	UN/ID	PG		PI	Label	Mark	
	9	3082	III		964/964	9	EHS	
IMDG	Class	UN	PG	EmS	PI	Label	Mark	
	9	3082	III	F-A S-F	P001/IBC03	9	marine po	ollutant
RID/ADR	Class	UN	PG	Haz.no	PI	Label	Mark	Classif. code
	9	3082	Ш	90	P001/IBC03	9	EHS	M6

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DOT	Class	UN/ID	PG	PI	RQ	Label	Haz.no
	9	3082	III			9	

DOT Remark: - NON-REGULATED IN NON-BULK PACKAGINGS

TRANSPORTED BY MOTOR VEHICLES, RAIL CARS OR

AIRCRAFT (49CFR 171.4(c)).

Proper shipping name ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID,

N.O.S.

Technical name Mycophenolate mofetil

## **SECTION 15: Regulatory information**

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

TSCA Status - FDA Exemption - not on inventory

Reporting Requirements - The United States Environmental Protection Agency (USEPA) has

not established a Reportable Quantity (RQ) for releases of this

material.

- In New Jersey, report all releases which are likely to endanger the public health, harm the environment or cause a complaint to the

NJDEPE Hotline (1-609-292-5560) and to local officials.

- State and local regulations vary and may impose additional

reporting requirements.

## **SECTION 16: Other information**

Full text of H-Statements referred to under section 3

H302 Harmful if swallowed.

H341 Suspected of causing genetic defects.

H360D May damage the unborn child.

H372 Causes damage to organs through prolonged or repeated

exposure.

USH003 May form combustible dust concentrations in the air

Note

Edition documentation - first edition

The information in this safety data sheet is based on current scientific knowledge. It should not be taken as expressing or implying any warranty concerning product characteristics.

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