

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

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SECTION 2: Hazards identification

Classification of the substance or mixture / Label elements

GHS Classification

Health Hazards:

- 3.5 Germ 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 1)
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 + P310 IF exposed or if you feel unwell: Immediately 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
128794-94-5

17.2 %

- Combustible dust (No category), USH003
- 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 'Hazard statements' 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 | <ul style="list-style-type: none">- rinse immediately with tap water for at least 20 minutes - open eyelids forcibly- consult a physician |
| Skin contact | <ul style="list-style-type: none">- remove immediately contaminated clothes, wash affected skin with water and soap - do not use any solvents |
| Inhalation | <ul style="list-style-type: none">- 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 | <ul style="list-style-type: none">- no information available |
|------|--|

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

- | | |
|-------------------|--|
| Note to physician | <ul style="list-style-type: none">- treat symptomatically- after accidental exposure women should get medical advice from a physician |
|-------------------|--|

SECTION 5: Firefighting measures

5.1. Extinguishing media

- | | |
|------------------------------|---|
| Suitable extinguishing media | <ul style="list-style-type: none">- 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 | <ul style="list-style-type: none">- formation of toxic and corrosive combustion gases (nitrogen oxides (NO_x)) possible- consider danger for the environment: dike spilled liquid |
|------------------|--|

5.3. Advice for firefighters

- | | |
|---------------------------------|---|
| Protection of fire-fighters | <ul style="list-style-type: none">- precipitate gases/vapours/mists with water spray |
| Special method of fire-fighting | <ul style="list-style-type: none">- 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 | <ul style="list-style-type: none">- 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 draining 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
- PNEC
- 0.068 µg/l, based on acute data, surface waters *2
 - 312.5 µg/l, based on chronic data, provisional antibiotic resistance PNEC *2

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)

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Eye protection - safety glasses
Body protection - protective clothing

*1 referring to: Mycophenolate mofetil
*2 referring to: Mycophenolic acid

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Color white to off-white
Form aqueous suspension
Solubility ≤ 22 mg/l, water (~ 22 °C, pH 6.3, HPLC, 24 h) *1
 ≤ 36 mg/l, aquatic ecotoxicity media (~ 22 °C, HPLC, 24 h) *1
Partition coefficient $\log P_{ow}$ 2.38 (n-octanol/water) pH 7.4 *1

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	- LD ₅₀ 353 mg/kg (oral, rat); females, "trimmed Spearman-Karber method"	*1
	- LD ₅₀ 500 mg/kg (oral, rat); males, "trimmed Spearman-Karber method"	*1
	- LD ₀ 250 mg/kg (oral, rat)	*1
	- LD ₅₀ > 4'000 mg/kg (oral, mouse)	*1
Local effects	- skin: non-irritant	*1
	- eye: non-irritant	*1
Sensitization	- non-sensitizing	*1
Mutagenicity	- mutagenic (OECD No. 474 (Micronucleus Test)); threshold dose response	*2
	- mutagenic (mouse lymphoma cell mutation test; OECD No. 476 (Mammalian Cell Gene Mutation Test))	*2
	- does not induce chromosomal aberrations in vitro (OECD No. 473 (Mammalian Cytogenic Test))	*2
	- not mutagenic (Ames test)	*2
Carcinogenicity	- not carcinogenic (several species)	*1
Reproductive toxicity	- teratogenic (several species)	*1
STOT-single exposure	- no information available	
STOT-repeated exposure	- no information available	
Aspiration hazard	- no information available	
Note	- immunosuppressive agent	*1
	- average therapeutic dose 1 g twice daily	*1
	- elimination half-life: 16-18 h	*1
	- adverse effects at therapeutic dose: diarrhea, drop in white blood cell count, susceptibility to infections, vomiting	*1
	- high doses may irritate the digestive tract	*1
	- excretion predominantly renal	*1

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- Potential Health Effects
- Exposure: Inhalation, Ingestion, Skin contact, Eye contact
 - Carcinogenicity: not listed by NTP, IARC or OSHA

*1 referring to: Mycophenolate mofetil
*2 referring to: 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)
EC₅₀ (48 h) > 100 mg/l (nominal concentration)
NOEC (48 h) 27.7 mg/l (average measured concentration) (OECD No. 202) *1
 - highly toxic for algae (Scenedesmus (=Desmodesmus) subspicatus)
ErC₅₀ (72 h) 0.6 mg/l (average measured concentration)
EbC₅₀ (72 h) 0.2 mg/l (average measured concentration)
NOEC (72 h) 0.1 mg/l (nominal concentration) (OECD No. 201) *1
 - 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) *1
 - 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) *1
 - no adverse influence on substrate biodegradation (activated sludge)
concentration (14 d) 100 mg/l (nominal concentration) (Manometric Respirometry Test, OECD No. 301 F) *1

12.2. Persistence and degradability

- Ready biodegradability
- not readily biodegradable
~ 14 %, 64 d
(FDA Technical Assistance Document No. 3.11) *1
 - not readily biodegradable
primary degradation evidenced by HPLC
< 6 %, 28 d
(Manometric Respirometry Test, OECD No. 301 F) *1
- Inherent biodegradability
- evidence for medium-term biodegradation in surface waters (analogous to OECD 308, Transformation in natural water/sediment systems) *1
- Abiotic degradation
- rapid degradation, photodegradation, hydrolysis 22.3 mg/l, water; HPLC
~ 37 %, 120 h, ~ 22 °C, dark
~ 67 %, 120 h, ~ 22 °C, under illumination *1

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12.3. Bioaccumulative potential

Note - no information available

12.4. Mobility in soil

Note - no information available

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 pollutant

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RID/ADR	Class	UN	PG	Haz.no	PI	Label	Mark	Classif. code
	9	3082	III	90	P001/IBC03	9	EHS	M6
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		<ul style="list-style-type: none"> - 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 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		- changes from previous version in sections 8, 15						
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.								