

TOPAMAX

 Version
 Revision Date:
 SDS Number:
 Date of last issue: 2022/05/19

 2.14
 2022/09/25
 100000014488
 Date of first issue: 2018/04/12

SECTION 1. IDENTIFICATION

Product name : TOPAMAX

Substance name : TOPAMAX 50 mg tablets

Topiramate Tablets 50 mg

Manufacturer or supplier's details

Company name of supplier : Janssen Pharmaceuticals, Inc.

Address : 1125 Trenton-Harbourton Rd

Titusville NJ 08560

USA

Telephone : +16097302000

E-mail address of person responsible for the SDS

SDSJanssen@its.jnj.com

Emergency telephone : CHEMTREC US: 1-800-424-9300

number CHEMTREC International: +1 703-741-5970

Recommended use of the chemical and restrictions on use

Recommended use : Finished Pharmaceutical Product

Pharmacotherapeutic group: Antiepileptics

This SDS is only intended for occupational use and not for consumer use (see patient packaging insert for consumer use). This SDS is written to provide environmental, health and safety information for personnel that will be handling this finished pharmaceutical product. For health and safety information during manufacturing of this product we refer to

the appropriate SDS for each component.

This dosage form is exempt from the requirements of the OSHA Hazard Communication Standard (US OSHA Standard

29 CFR Part 1910.1200).

SECTION 2. HAZARDS IDENTIFICATION

GHS classification in accordance with the OSHA Hazard Communication Standard (29 CFR 1910.1200)

Reproductive toxicity : Category 2

GHS label elements



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Hazard pictograms

Signal word : Warning

Hazard statements : H361 Suspected of damaging fertility or the unborn child.

Precautionary statements : Prevention:

P201 Obtain special instructions before use.

P202 Do not handle until all safety precautions have been read

and understood.

P280 Wear protective gloves/ protective clothing.

Response:

P308 + P313 IF exposed or concerned: Get medical advice/

attention.

Storage:

P405 Store locked up.

Disposal:

P501 Dispose of contents/ container to an approved waste

disposal plant.

Additional Labelling

The following percentage of the mixture consists of ingredient(s) with unknown acute toxicity: 15.43%

Other hazards

Refer to the pharmacotherapeutic group (section 1.2) and the patient packaging insert to evaluate the possible workplace hazards when this Finished Pharmaceutical Product is accidently leaking, broken or crushed.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Chemical nature : Solid

Components

Chemical name	CAS-No.	Concentration (% w/w)
TOPIRAMATE	97240-79-4	>= 50 - < 70
Cellulose	9004-34-6	>= 10 - < 20
Octadecanoic acid, magnesium salt	557-04-0	>= 0.1 - < 1
TOPIRAMATE	97240-79-4	>= 50 - < 70
Cellulose	9004-34-6	>= 10 - < 20
Octadecanoic acid, magnesium salt	557-04-0	>= 0.1 - < 1



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Actual concentration is withheld as a trade secret

SECTION 4. FIRST AID MEASURES

If inhaled : Health injuries are not known or expected under normal use.

If breathed in, move person into fresh air.

Consult a physician.

In case of skin contact : Take off all contaminated clothing immediately.

Wash off with soap and water.

If symptoms persist, call a physician.

In case of eye contact : Rinse immediately with plenty of water, also under the eyelids,

for at least 5 minutes. Remove contact lenses.

If eye irritation persists, consult a specialist.

If swallowed : If swallowed, rinse mouth with water (only if the person is

conscious).

Call a physician immediately.

Most important symptoms and effects, both acute and

delayed

Consult the patient packaging insert for more information

about this Finished Pharmaceutical Product.

Abdominal pain anorexia

insomnia
Disorientation
Dizziness
Tremors
Blurred vision
nausea
Diarrhoea
constipation

stomach discomfort

dry mouth Spasm Fatigue

nasopharyngitis

Cough

weight decrease

Notes to physician : Treat symptomatically.

Consult the patient packaging insert for more information

about this Finished Pharmaceutical Product.

SECTION 5. FIREFIGHTING MEASURES

Suitable extinguishing media : Use extinguishing measures that are appropriate to local

circumstances and the surrounding environment.

Hazardous combustion

products

No information available.



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Further information No information available.

for firefighters

Special protective equipment : In the event of fire, wear self-contained breathing apparatus.

SECTION 6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures

In the event of an accidental release the emergency response team must respond based on a risk assessment and use

personal protective equipment as appropriate.

Environmental precautions Should not be released into the environment.

Do not flush into surface water or sanitary sewer system.

Methods and materials for containment and cleaning up Large spills + Small spills: Keep in suitable, closed containers for disposal. Treat recovered material as described in the

section "Disposal considerations".

Large spills: Sweep up (intact) or vacuum with HEPA filter (broken or crushed) or via wet cleaning into suitable

containers for disposal. Pick up and arrange without creating

dust. Keep in properly labelled containers.

Small spills: Moisten a towel, cover the spill, pick up the spill

or use HEPA vacuum.

SECTION 7. HANDLING AND STORAGE

Advice on protection against

fire and explosion

No data available

Advice on safe handling To avoid thermal decomposition, do not overheat.

> Avoid inhalation, ingestion and contact with skin and eyes. Do not break, crush or spill this Finished Pharmaceutical

Product.

Use personal protective equipment as required.

To maintain product quality, do not store in heat or direct Conditions for safe storage

sunlight.

Store in original container.

Keep containers tightly closed in a dry, cool and well-

ventilated place.

Keep away from heat and sources of ignition.

Keep locked up.

Recommended storage

temperature

< 77 °F / < 25 °C

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components with workplace control parameters



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Components	CAS-No.	Value type	Control	Basis	
Components	CAS-NO.	(Form of	parameters /	Dasis	
		exposure)	Permissible		
		exposure)	concentration		
TOPIRAMATE	97240-79-4	TWA	0.120 mg/m3	J&J	
TOPINAMATE	31240-13-4	1 1 1 1 1 1	0.120 mg/m3	OEL/PBOEL	
				HHC	
		PBOEL-HHC	1 B	J&J	
		1 BOLL 11110		OEL/PBOEL	
				HHC	
	Further inform	ation: J&J has a	hazard banding nota		
			ied by J&J as being l		
			potential to have adv		
		n and fetal deve			
Cellulose	9004-34-6	TWA	10 mg/m3	ACGIH	
		TWA	5 mg/m3	NIOSH REL	
		(Respirable)	-		
		TWA (total)	10 mg/m3	NIOSH REL	
		TWA (total	15 mg/m3	OSHA Z-1	
		dust)			
		TWA	5 mg/m3	OSHA Z-1	
		(respirable			
		fraction)			
		TWA (Total	15 mg/m3	OSHA P0	
		dust)			
		TWA	5 mg/m3	OSHA P0	
		(respirable			
		dust fraction)			
Octadecanoic acid,	557-04-0	TWA	10 mg/m3	ACGIH	
magnesium salt		(Inhalable			
		particulate			
		matter) TWA	2	ACCILI	
			3 mg/m3	ACGIH	
		(Respirable particulate			
		matter)			
TOPIRAMATE	97240-79-4	TWA	0.120 mg/m3	J&J	
I OT II ON III C	31270-13-4	1 4 4 7	0.120 mg/m3	OEL/PBOEL	
				HHC	
		PBOEL-HHC	1 B	J&J	
		. 2022 11110		OEL/PBOEL	
				HHC	
	Further information: J&J has a hazard banding notation: PBOEL HHC. This substance is classified by J&J as being PBOEL HHC 1B., Notation REPRO: has the potential to have adverse effects on reproduction and fetal development				
Cellulose	9004-34-6	TWA	10 mg/m3	ACGIH	
		TWA	5 mg/m3	NIOSH REL	
		(Respirable)			
		TWA (total)	10 mg/m3	NIOSH REL	
		TWA (total	15 mg/m3	OSHA Z-1	
		dust)			



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		TWA (respirable fraction)	5 mg/m3	OSHA Z-1
		TWA (Total dust)	15 mg/m3	OSHA P0
		TWA (respirable dust fraction)	5 mg/m3	OSHA P0
Octadecanoic acid, magnesium salt	557-04-0	TWA (Inhalable particulate matter)	10 mg/m3	ACGIH
		TWA (Respirable particulate matter)	3 mg/m3	ACGIH

Engineering measures : All personal protective equipment should be based on a risk

assessment. Consult a Environment Health Safety expert if

necessary.

If this product is processed not in accordance with the prescribed use, contact the Industrial Hygiene / Environment

Health Safety Expert to assess the situation.

Validated Industrial Hygiene Analytical methods are developed to monitor and quantify inhalable exposure to the

Active Pharmaceutical Ingredient. For more information contact Maxxam Analytics (www.maxxamlabs.com) or the Laboratory of Occupational and Environmental Hygiene

(www.lamh.be).

Personal protective equipment

Respiratory protection : No personal respiratory protective equipment normally

required.

Hand protection

Remarks : Disposable gloves

Eye protection : No special precautions required.

Skin and body protection : closed work clothing

Protective measures : The type of protective equipment must be selected based on

the Environmental Health and Safety risk assessment. Consult a Environmental Health and Safety expert if

necessary.

Hygiene measures : Handle in accordance with good industrial hygiene and safety

practice.

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES



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Appearance : tablet, Coated

Colour : No data available

Odour : No data available

Odour Threshold : No data available

pH : No data available

Melting point/range : No data available

Boiling point/boiling range : No data available

Flash point : No data available

Evaporation rate : No data available

Flammability (solid, gas) : No information available.

Self-ignition : No data available

Upper explosion limit / Upper

flammability limit

No data available

Lower explosion limit / Lower

flammability limit

No data available

Vapour pressure : No data available

Relative density : No data available

Density : No data available

Solubility(ies)

Water solubility : No data available

Partition coefficient: n-

octanol/water

No data available

Decomposition temperature : No data available

Viscosity

Viscosity, dynamic : Not applicable

Viscosity, kinematic : Not applicable

Explosive properties : No data available

Oxidizing properties : No data available

SECTION 10. STABILITY AND REACTIVITY



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Reactivity : None reasonably foreseeable.

Chemical stability : Stable under normal conditions.

Possibility of hazardous

reactions

: No dangerous reaction known under conditions of normal use.

Conditions to avoid : To avoid thermal decomposition, do not overheat.

Heat, flames and sparks. Exposure to moisture

Incompatible materials : None known.

Hazardous decomposition

products

None known.

SECTION 11. TOXICOLOGICAL INFORMATION

Acute toxicity

Product:

Acute oral toxicity : Acute toxicity estimate: 4,418 mg/kg

Method: Calculation method

Components:

TOPIRAMATE:

Acute oral toxicity : LD50 (Rat): 2,436 mg/kg

Acute inhalation toxicity : Remarks: May be harmful if inhaled.

Acute dermal toxicity : Remarks: No data available

Cellulose:

Acute oral toxicity : LD50 (Rat): > 5,000 mg/kg

TOPIRAMATE:

Acute oral toxicity : LD50 (Rat): 2,436 mg/kg

Acute inhalation toxicity : Remarks: May be harmful if inhaled.

Acute dermal toxicity : Remarks: No data available

Cellulose:

Acute oral toxicity : LD50 (Rat): > 5,000 mg/kg



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Skin corrosion/irritation

Components:

TOPIRAMATE:

Species : Rabbit

Result : No skin irritation

Cellulose:

Species : Rabbit

Remarks : No skin irritation

TOPIRAMATE:

Species : Rabbit

Result : No skin irritation

Cellulose:

Species : Rabbit

Remarks : No skin irritation

Serious eye damage/eye irritation

Components:

TOPIRAMATE:

Species : Rabbit

Result : No eye irritation

Cellulose:

Species : Rabbit

Remarks : No eye irritation

TOPIRAMATE:

Species : Rabbit

Result : No eye irritation

Cellulose:

Species : Rabbit

Remarks : No eye irritation

Respiratory or skin sensitisation

Components:

TOPIRAMATE:

Result : Not a sensitizer



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Cellulose:

Species : Guinea pig

Remarks : Did not cause sensitisation on laboratory animals.

TOPIRAMATE:

Result : Not a sensitizer

Cellulose:

Species : Guinea pig

Remarks : Did not cause sensitisation on laboratory animals.

Germ cell mutagenicity

Components:

TOPIRAMATE:

Genotoxicity in vitro : Test Type: Ames test

Result: negative

Test Type: Chromosome aberration test in vitro

Test system: Human lymphocytes

Result: negative

Test Type: A mouse lymphoma test

Result: negative

Genotoxicity in vivo : Test Type: In vivo micronucleus test

Species: Rat Result: negative

TOPIRAMATE:

Genotoxicity in vitro : Test Type: Ames test

Result: negative

Test Type: Chromosome aberration test in vitro

Test system: Human lymphocytes

Result: negative

Test Type: A mouse lymphoma test

Result: negative

Genotoxicity in vivo : Test Type: In vivo micronucleus test

Species: Rat Result: negative



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Carcinogenicity

Components:

TOPIRAMATE:

Species : Rat
Application Route : Oral
Exposure time : 2 years

Dose : 120 mg/kg/day

Remarks : no

carcinogenic effects

TOPIRAMATE:

Species : Rat
Application Route : Oral
Exposure time : 2 years

Dose : 120 mg/kg/day

Remarks : no

carcinogenic effects

IARC No component of this product present at levels greater than or equal to 0.1% is

identified as probable, possible or confirmed human carcinogen by IARC.

No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.

OSHANo component of this product present at levels greater than or equal to 0.1% is

on OSHA's list of regulated carcinogens.

No component of this product present at levels greater than or equal to 0.1% is

on OSHA's list of regulated carcinogens.

NTP No component of this product present at levels greater than or equal to 0.1% is

identified as a known or anticipated carcinogen by NTP.

No component of this product present at levels greater than or equal to 0.1% is

identified as a known or anticipated carcinogen by NTP.

Reproductive toxicity

Components:

TOPIRAMATE:

Effects on foetal : Species: Rabbit development : Remarks: positive

Species: Rat Remarks: positive

Species: Mouse Remarks: positive



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Reproductive toxicity -

Assessment

Some evidence of adverse effects on development, based on

animal experiments.

Teratogenicity - Assessment : Limited evidence of adverse effects on development in animal

studies and/ or human studies.

TOPIRAMATE:

Effects on foetal development

Species: Rabbit Remarks: positive

Species: Rat Remarks: positive

Species: Mouse Remarks: positive

Reproductive toxicity -

Assessment

Some evidence of adverse effects on development, based on

animal experiments.

Teratogenicity - Assessment : Limited evidence of adverse effects on development in animal

studies and/ or human studies.

STOT - single exposure

Components:

TOPIRAMATE:

Remarks : No data available

TOPIRAMATE:

Remarks : No data available

STOT - repeated exposure

No data available

Repeated dose toxicity

Components:

TOPIRAMATE:

Species : Rat
NOAEL : 10 mg/kg
Exposure time : 3 months

Species : Dog NOAEL : 10 mg/kg Exposure time : 3 months

Species : Dog NOAEL : 10 mg/kg Exposure time : 12 months



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TOPIRAMATE:

Species : Rat

NOAEL : 10 mg/kg

Exposure time : 3 months

Species : Dog NOAEL : 10 mg/kg Exposure time : 3 months

Species : Dog NOAEL : 10 mg/kg Exposure time : 12 months

Aspiration toxicity

No data available

Experience with human exposure

No data available

Toxicology, Metabolism, Distribution

No data available

Neurological effects

No data available

Further information

No data available

Other health hazards

No data available

SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity

Components:

TOPIRAMATE:

Toxicity to fish : NOEC (Lepomis macrochirus (Bluegill sunfish)): < 75 mg/l

Method: FDA 4.11

LC50 (Lepomis macrochirus (Bluegill sunfish)): > 2,400 mg/l

Exposure time: 96 h Method: FDA 4.11

Toxicity to daphnia and other :

aquatic invertebrates

NOEC (Daphnia (water flea)): 1,000 mg/l

Method: FDA 4.08

EC50 (Daphnia (water flea)): 1,000 mg/l

Method: FDA 4.08



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Toxicity to algae/aquatic

plants

: EbC50 (Pseudokirchneriella subcapitata (green algae)): > 93

mg/l

Test Type: Cell multiplication inhibition test

Method: OECD Test Guideline 201

NOECb (Pseudokirchneriella subcapitata (green algae)): 93

mg/l

Test Type: Cell multiplication inhibition test

Method: OECD Test Guideline 201

ErC50 (Pseudokirchneriella subcapitata (green algae)): > 93

mg/l

Test Type: Growth inhibition

Method: OECD Test Guideline 201

NOECr (Pseudokirchneriella subcapitata (green algae)): 93

mg/l

Test Type: Growth inhibition

Method: OECD Test Guideline 201

Toxicity to microorganisms : NOEC (activated sludge): >= 1,000 mg/l

Exposure time: 3 h

Method: OECD Test Guideline 209

EC50 (activated sludge): > 1,000 mg/l

Exposure time: 3 h

Method: OECD Test Guideline 209

EC50: > 100 mg/l

Test Type: Growth inhibition

Method: similar to OECD Test Guideline 209

MIC: > 10 mg/l

Test Type: Growth inhibition

Method: similar to OECD Test Guideline 209

Cellulose:

Toxicity to fish : Remarks: No data available

TOPIRAMATE:

Toxicity to fish : NOEC (Lepomis macrochirus (Bluegill sunfish)): < 75 mg/l

Method: FDA 4.11

LC50 (Lepomis macrochirus (Bluegill sunfish)): > 2,400 mg/l

Exposure time: 96 h Method: FDA 4.11

Toxicity to daphnia and other :

aquatic invertebrates

NOEC (Daphnia (water flea)): 1,000 mg/l

Method: FDA 4.08

EC50 (Daphnia (water flea)): 1,000 mg/l

Method: FDA 4.08



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Toxicity to algae/aquatic

plants

EbC50 (Pseudokirchneriella subcapitata (green algae)): > 93

mg/l

Test Type: Cell multiplication inhibition test

Method: OECD Test Guideline 201

NOECb (Pseudokirchneriella subcapitata (green algae)): 93

mg/l

Test Type: Cell multiplication inhibition test

Method: OECD Test Guideline 201

ErC50 (Pseudokirchneriella subcapitata (green algae)): > 93

mg/l

Test Type: Growth inhibition

Method: OECD Test Guideline 201

NOECr (Pseudokirchneriella subcapitata (green algae)): 93

mg/l

Test Type: Growth inhibition

Method: OECD Test Guideline 201

Toxicity to microorganisms : NOEC (activated sludge): >= 1,000 mg/l

Exposure time: 3 h

Method: OECD Test Guideline 209

EC50 (activated sludge): > 1,000 mg/l

Exposure time: 3 h

Method: OECD Test Guideline 209

EC50: > 100 mg/l

Test Type: Growth inhibition

Method: similar to OECD Test Guideline 209

MIC: > 10 mg/l

Test Type: Growth inhibition

Method: similar to OECD Test Guideline 209

Cellulose:

Toxicity to fish : Remarks: No data available

Persistence and degradability

Components:

TOPIRAMATE:

Biodegradability : Remarks: No data available

TOPIRAMATE:

Biodegradability : Remarks: No data available



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Bioaccumulative potential

Components:

TOPIRAMATE:

Bioaccumulation : Remarks: No data available

Partition coefficient: n-

octanol/water

Remarks: No data available

Octadecanoic acid, magnesium salt:

Partition coefficient: n- : Remarks: No data available

octanol/water

TOPIRAMATE:

Bioaccumulation : Remarks: No data available

Partition coefficient: n-

octanol/water

Remarks: No data available

Octadecanoic acid, magnesium salt:

Partition coefficient: n- : Remarks: No data available

octanol/water

Mobility in soil

Components:

TOPIRAMATE:

Distribution among : log Koc: 2.1

environmental compartments Method: OECD Test Guideline 121

Remarks: high mobility

TOPIRAMATE:

Distribution among : log Koc: 2.1

environmental compartments Method: OECD Test Guideline 121

Remarks: high mobility

Other adverse effects

Components:

TOPIRAMATE:

Results of PBT and vPvB

assessment

No information available.

Additional ecological

information

: No data available No data available



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TOPIRAMATE:

Results of PBT and vPvB

assessment

No information available.

Additional ecological

information

No data available No data available

SECTION 13. DISPOSAL CONSIDERATIONS

Disposal methods

Waste from residues : In accordance with National, Federal, State and Local

regulations.

Contaminated packaging : Empty containers should be taken to an approved waste

handling site for recycling or disposal.

SECTION 14. TRANSPORT INFORMATION

International Regulations

UNRTDG

Not regulated as a dangerous good

IATA-DGR

Not regulated as a dangerous good

IMDG-Code

Not regulated as a dangerous good

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied.

National Regulations

49 CFR

Not regulated as a dangerous good

SECTION 15. REGULATORY INFORMATION

SARA 302 Extremely Hazardous Substances Threshold Planning Quantity

This material does not contain any components with a section 302 EHS TPQ.

SARA 313 : This material does not contain any chemical components with

known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

US State Regulations

Massachusetts Right To Know



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Cellulose 9004-34-6

Massachusetts Right To Know

Cellulose 9004-34-6

Pennsylvania Right To Know

TOPIRAMATE 97240-79-4
Cellulose 9004-34-6
alpha-D-Glucopyranose, 4-O-beta-D-galactopyranosyl-, 5989-81-1

hydrate (1:1)

NATRIUM ZETMEELGLYCOLAAT 9063-38-1

Pennsylvania Right To Know

TOPIRAMATE 97240-79-4
Cellulose 9004-34-6
alpha-D-Glucopyranose, 4-O-beta-D-galactopyranosyl-,
hydrate (1:1)
NATRIUM ZETMEELGLYCOLAAT 9063-38-1

Maine Chemicals of High Concern

Product does not contain any listed chemicals

Vermont Chemicals of High Concern

Product does not contain any listed chemicals

Washington Chemicals of High Concern

Product does not contain any listed chemicals

New Jersey Right To Know

TOPIRAMATE 97240-79-4
Cellulose 9004-34-6
alpha-D-Glucopyranose, 4-O-beta-D-galactopyranosyl-,
hydrate (1:1)
NATRIUM ZETMEELGLYCOLAAT 9063-38-1

New York City Hazardous Substances

No components listed on the New York City Hazardous Substances List

New York City Hazardous Substances

Carnauba wax 8015-86-9

California Prop. 65

WARNING: This product contains a chemical known to the State of California to cause birth defects or other reproductive harm.

WARNING: This product can expose you to chemicals including TOPIRAMATE, which is/are known to the State of California to cause birth defects or other reproductive harm. For more information go to www.P65Warnings.ca.gov.

California Permissible Exposure Limits for Chemical Contaminants

Cellulose 9004-34-6

California Permissible Exposure Limits for Chemical Contaminants

Cellulose 9004-34-6

The components of this product are reported in the following inventories:



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> This product is not subject to TSCA and TSCA 12(b) Export notification because Food. Drugs and cosmetic products are exempt.

Other regulations

Medicinal products in the finished state, intended for the final user, are not subject to GHS labeling.

Restricted to professional users.

SECTION 16. OTHER INFORMATION

Full text of other abbreviations

USA. ACGIH Threshold Limit Values (TLV)

J&J OEL/PBOEL HHC J&J OEL/PBOEL HHC

NIOSH REL USA, NIOSH Recommended Exposure Limits

OSHA PO USA. Table Z-1-A Limits for Air Contaminants (1989 vacated

USA. Occupational Exposure Limits (OSHA) - Table Z-1 OSHA Z-1

Limits for Air Contaminants

8-hour, time-weighted average ACGIH / TWA

J&J OEL/PBOEL HHC / TWA : Time weighted average **PBOEL-HHC**

J&J OEL/PBOEL HHC /

PBOEL-HHC

NIOSH REL / TWA Time-weighted average concentration for up to a 10-hour

workday during a 40-hour workweek

OSHA P0 / TWA 8-hour time weighted average OSHA Z-1 / TWA 8-hour time weighted average

AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight; CERCLA - Comprehensive Environmental Response, Compensation, and Liability Act; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN -Standard of the German Institute for Standardisation; DOT - Department of Transportation; DSL -Domestic Substances List (Canada); ECx - Concentration associated with x% response; EHS -Extremely Hazardous Substance; ELx - Loading rate associated with x% response; EmS -Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx -Concentration associated with x% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; HMIS - Hazardous Materials Identification System; IARC - International Agency for Research on Cancer; IATA -International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO -International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO -International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; MSHA - Mine Safety and Health Administration; n.o.s. - Not Otherwise Specified; NFPA - National Fire Protection Association; NO(A)EC - No Observed (Adverse) Effect Concentration: NO(A)EL - No Observed (Adverse) Effect Level: NOELR - No Observable Effect Loading Rate; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals;



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OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; RCRA - Resource Conservation and Recovery Act; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RQ - Reportable Quantity; SADT - Self-Accelerating Decomposition Temperature; SARA - Superfund Amendments and Reauthorization Act; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TECI - Thailand Existing Chemicals Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative

Revision Date : 2022/09/25

Date and Number Formats

This document uses the following notation for printing dates and numbers:

Date: Dec 31th, 2012 as 2012/12/31

Numbers: 123456,78 as 123,456.78

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