

Material Safety Data Sheet AVASTIN(R) Vials (100 mg)

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

1.1. Product identifier

Product name AVASTIN(R) Vials (100 mg)

Product code SAP-10086726

Synonyms - AVASTIN(R) Vials (100 mg/4 ml)

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

Use - formulated pharmaceutical active substance (antineoplastic)

1.3. Details of the supplier of the safety data sheet

Company information Enquiries: Local representation:

Hoffmann-La Roche Inc. 340 Kingsland Street

USA-Nutley, N.J. 07110-1199 United States of America

Phone 001-973/235 50 00 E-Mail info.sds@roche.com

US Emergency phone: (800)-827-6243 US Chemtrec phone: (800)-424-9300

1.4. Emergency telephone number

Emergency telephone number US emergency phone: (800)-827-6243

SECTION 2: Hazards identification

Emergency Overview

Form aqueous solution

sterile liquid

Color colorless

Hazard Overview - May cause allergic reactions.

- May cause birth defects based on animal data.

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Potential Health Effects - Exposure: Inhalation, Ingestion, Skin contact, Eye contact

- Target Organs: Hematopoietic/blood system, Immune System

- Acute Effects: May cause allergic reactions., This material is not likely to be significantly absorbed via occupational routes of entry due to its chemical structure and large molecular weight.

- Chronic Effects: May cause blood system effects.

- Carcinogenicity: not listed by NTP, IARC or OSHA

Classification of the substance or mixture / Label elements

GHS Classification no classification and labelling according to GHS

Other hazards

Additional Health Information

- Conditions aggravated: Hypersensitivity to this material and other
 - materials in its chemical class.

Reproductive Toxicity: May cause birth defects. Since this
material may effect the developing fetus, females planning to have
a child and pregnant women should exercise caution regarding
exposure.

It is also advisable for nursing mothers to exercise caution regarding exposure.

SECTION 3: Composition/information on ingredients

Characterization bevacizumab and other inactive ingredients

Ingredients Concentration

Bevacizumab ~ 2 %

CAS: 216974-75-3

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

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

- in the event of symptoms get medical treatment

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

Note - no information available

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

Note to physician - treat symptomatically

SECTION 5: Firefighting measures

5.1. Extinguishing media

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

- adapt extinguishing media to surrounding fire conditions

Flash point (liquid) not applicable

5.2. Special hazards arising from the substance or mixture

Specific hazards - no particular hazards known

5.3. Advice for firefighters

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

SECTION 6: Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

Personal precautions - no special precautions required

6.2. Environmental precautions

Environmental protection - no special environmental precautions required

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 - aluminium, glass, enamel, stainless steel

Note - do not shake solution

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7.2. Conditions for safe storage, including any incompatibilities

Storage conditions - 2 - 8 °C

do not freezeprotected from light

Validity - 2 to 8 °C, in the unopened original container, see "best use before"

date stated on the label

Packaging materials - vials

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

Threshold value (Roche) air - IOEL (Internal Occupational Exposure Limit): 0.05 mg/m3 *1

8.2. Exposure controls

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.

- respiratory protection not necessary during normal operations

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

Eye protection - safety glasses

*1 referring to: Bevacizumab

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Color colorless

Form aqueous solution

sterile liquid

Density 1.031 g/ml

pH value 5.9 to 6.3

Boiling temperature ~ 100 °C

9.2. Other information

Note - no information available

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SECTION 10: Stability and reactivity

10.1. Reactivity

Note - no information available

10.2. Chemical stability

Stability - does not contain any antimicrobial preservative; therefore, care

must be taken to ensure the sterility of the prepared solution

10.3. Possibility of hazardous reactions

Note - no information available

10.4. Conditions to avoid

Note - no information available

10.5. Incompatible materials

Note - no information available

10.6. Hazardous decomposition products

Note - no information available

SECTION 11: Toxicological information

11.1. Information on toxicological effects

Acute toxicity - not bioavailable by oral administration *1

- NOEL 50 mg/kg (i.v., cynomolgus monkey)

*1

*1

Chronic toxicity - LOAEL 2 mg/kg/w (i.v., cynomolgus monkey; 26 weeks) *1

Reproductive toxicity - teratogenic and embryotoxic (i.v., rabbit) *1

- should be administered during pregnancy only if the potential benefit justifies the potential risk to the fetus

benefit justifies the potential fish to the letus

Note - humanized monoclonal antibody which binds to and inactivates

the vascular endothelial growth factor (VEGF)

- bevacizumab is effective in the treatment of advanced stages of colon and rectum carcinoma

colon and rectum carcinoma *1
- therapeutic dose: 5 mg/kg/2w *1
- elimination half-life: 20 d *1

- side effect(s) during therapy: tendency to bleeding,

thrombophlebitis, proteinuria *1

*1 referring to: Bevacizumab

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SECTION 12: Ecological information

12.1. Toxicity

Ecotoxicity - no adverse influence on substrate biodegradation (activated

sludge)

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

 barely toxic for algae (nominal concentration = 100 mg/l), growth inhibition possibly due to turbidity caused by test substance

(Scenedesmus (=Desmodesmus) subspicatus) ErC $_{50}$ (72 h) > 100 mg active substance/l EbC $_{50}$ (72 h) ~ 100 mg active substance/l NOEC (72 h) < 100 mg active substance/l

(OECD No. 201)

- barely toxic for planktonic crustaceans (nominal concentration

= 100 mg/l) (Daphnia magna)

 EC_{50} (48 h) > 100 mg active substance/l NOEC (48 h) 100 mg active substance/l

(OECD No. 202)

12.2. Persistence and degradability

Ready biodegradability - readily biodegradable

78 % BOD/ThOD, 28 d 96 % DOC. 28 d

(Manometric Respirometry Test, OECD No. 301 F)

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

SECTION 13: Disposal considerations

13.1. Waste treatment methods

Waste from residues - observe local/national regulations regarding waste disposal

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SECTION 14: Transport information

Note - not classified by transport regulations, proper shipping name

non-regulated

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

Edition documentation - changes from previous version in sections 1

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Material Safety Data Sheet AVASTIN(R) Vials (400 mg)

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Product code SAP-10062575

Synonyms - AVASTIN(R) Vials (400 mg/16 ml)

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Use - formulated pharmaceutical active substance (antineoplastic)

1.3. Details of the supplier of the safety data sheet

Company information Enquiries: Local representation:

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