Version	Revision Date:	SDS Number:	Date of last issue: 2019/03/06
1.13	2019/03/06	100000014542	Date of first issue: 2018/08/20

SECTION 1. IDENTIFICATION

Substance name : SPRAVATO™
Nasal spray device Delivering 0.2 mL Solution, containing 32.3 mg of esketamine hydrochloride aqueous solution (28 mg of esketamine)

Manufacturer or supplier's details

Company name of supplier : Janssen Pharmaceuticals, Inc.

Address : 1125 Trenton-Harbourton Rd
Titusville NJ 08560
US

Telephone : (609) 730-2000

E-mail address Responsible/issuing person : SDSJanssen@its.jnj.com

Emergency telephone number : CHEMTREC US: 1-800-424-9300
CHEMTREC International: +1 703-527-3887

Recommended use of the chemical and restrictions on use

Recommended use : Finished Pharmaceutical Product
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 not 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 29 CFR 1910.1200

Long-term (chronic) aquatic hazard : Category 3

GHS label elements

Hazard statements : H412 Harmful to aquatic life with long lasting effects.

Precautionary statements :

Prevention:

P273 Avoid release to the environment.

Disposal:

P501 Dispose of contents/ container to an approved waste disposal plant.

Other hazards

None known.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Chemical nature : Liquid

Substance name : SPRAVATO™

Hazardous components

Chemical name	CAS-No.	Concentration (% w/w)
(S)-(+)-ketamine hydrochloride	33643-47-9	>= 10 - < 20

SECTION 4. FIRST AID MEASURES

If inhaled : If breathed in, move person into fresh air.
Consult a physician.

In case of skin contact : Take off contaminated clothing and shoes 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 : anxiety
Dissociation
Dizziness
Increased blood pressure
nausea
numbness
sedation
taste disorders
Vertigo
Vomiting
lethargy
anxiety

Notes to physician : Treat symptomatically.

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.
Further information	:	No information available.
Special protective equipment for firefighters	:	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. Evacuate personnel to safe areas.
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: Dam up. Soak up with inert absorbent material. Keep in properly labelled containers. Small spills: Gently cover the spill with an absorbent towel or pad. Large spills + Small spills: Keep in suitable, closed containers for disposal. Treat recovered material as described in the section "Disposal considerations".

SECTION 7. HANDLING AND STORAGE

Advice on protection against fire and explosion	:	No data available
Advice on safe handling	:	Do not break, crush or spill this Finished Pharmaceutical Product. To avoid thermal decomposition, do not overheat. Use personal protective equipment as required. Avoid inhalation, ingestion and contact with skin and eyes.
Conditions for safe storage	:	To maintain product quality, do not store in heat or direct sunlight. Store in original container. Keep containers tightly closed in a cool, well-ventilated place. Keep away from heat and sources of ignition. Store at room temperature.
Recommended storage temperature	:	15 - 25 °C

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components with workplace control parameters

Components	CAS-No.	Value type (Form of exposure)	Control parameters / Permissible concentration	Basis
(S)-(+)-ketamine hydrochloride	33643-47-9	TWA	0.024 mg/m3	J&J OEL/PBOEL HHC
		STEL	0.19 mg/m3	J&J OEL/PBOEL HHC
		PBOEL-HHC	2	J&J 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 2.			

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

Appearance : solution

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
Self-ignition	:	No data available
Upper explosion limit	:	No data available
Lower explosion limit	:	No data available
Vapour pressure	:	No data available
Relative vapour density	:	No data available
Relative density	:	No data available
Density	:	No data available
Solubility(ies)		
Water solubility	:	No data available
Solubility in other solvents	:	No data available
Partition coefficient: n-octanol/water	:	No data available
Decomposition temperature	:	No data available
Viscosity		
Viscosity, dynamic	:	Not applicable
Viscosity, kinematic	:	No data available

SECTION 10. STABILITY AND REACTIVITY

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.
Incompatible materials	:	None known.

Hazardous decomposition products : None known.

SECTION 11. TOXICOLOGICAL INFORMATION

Acute toxicity

Product:

Acute oral toxicity : Acute toxicity estimate: 3,817 mg/kg
Method: Calculation method

Components:

(S)-(+)-ketamine hydrochloride:

Acute oral toxicity : LD50 (Mouse): 616 mg/kg
Method: Acute oral toxicity

Acute inhalation toxicity : Remarks: No data available

Acute dermal toxicity : Remarks: No data available

Acute toxicity (other routes of administration) : LD50 (Rat): 200 mg/kg
Application Route: intraperitoneal; injection made in the abdominal area
Method: Acute toxicity study

LD50 (Rat): 35 mg/kg
Application Route: intravenous injection
Method: Acute toxicity study

Skin corrosion/irritation

Components:

(S)-(+)-ketamine hydrochloride:

Remarks: No data available

Serious eye damage/eye irritation

Components:

(S)-(+)-ketamine hydrochloride:

Remarks: No data available

Respiratory or skin sensitisation

Components:

(S)-(+)-ketamine hydrochloride:

Result: Not expected to cause skin sensitization

Germ cell mutagenicity

Components:

(S)-(+)-ketamine hydrochloride:

Genotoxicity in vitro : Method: Bacterial Reverse Mutation Test OECD 471
Result: negative
GLP: yes

: Method: In Vitro Mammalian Cell Gene Mutation Test (MLA TK) OECD 476
Result: positive
GLP: yes

: Method: In Vitro Mammalian Cell Micronucleus Test OECD 487
Result: positive
GLP: no

Genotoxicity in vivo : Method: In vivo Mammalian Erythrocyte Micronucleus Test OECD 474
Result: negative
GLP: yes

Method: In vivo Single Cell Gel Electrophoresis Assay (Comet Assay)
Result: negative
GLP: yes

Germ cell mutagenicity - Assessment : No evidence of mutagenicity based on weight of evidence.

Carcinogenicity

Components:

(S)-(+)-ketamine hydrochloride:

Species: Mouse, (male and female)
Application Route: Subcutaneous; injection made in the back or neck of animal
Exposure time: 26 weeks
Dose: 10, 25, 75 mpk
Method: carcinogenicity study
Result: No evidence of carcinogenicity in animal studies.
GLP: yes

Species: Rat, (male and female)
Application Route: intranasal (IN) administration
Exposure time: 104 weeks
Dose: 2,7-9-27 mg/kg
Method: carcinogenicity study
Result: No evidence of carcinogenicity in animal studies.
GLP: yes

Carcinogenicity - Assess- : No evidence of carcinogenicity.

ment

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.
OSHA	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.

Reproductive toxicity

Components:

(S)-(+)-ketamine hydrochloride:

Effects on fertility	:	Species: Rat, male and female Application Route: intranasal (IN) administration Dose: 2,7 - 9 - 27 mg/kg General Toxicity - Parent: NOAEL: 2.7 mg/kg General Toxicity F1: NOAEL: 27 mg/kg GLP: yes
Effects on foetal development	:	Species: Rat, female Application Route: intranasal (IN) administration Dose: 15, 50, 150 mg/kg General Toxicity Maternal: NOAEL: 15 mg/kg Teratogenicity: NOAEL: 150 mg/kg Method: Developmental Toxicity GLP: yes Species: Rabbit, female Application Route: intranasal (IN) administration Dose: 10, 30, 100 mg/kg General Toxicity Maternal: NOAEL: 10 mg/kg Teratogenicity: NOAEL: 10 mg/kg Method: Developmental Toxicity GLP: yes Species: Rat, female Application Route: intranasal (IN) administration Dose: 2,7 - 9 - 27 mg/kg General Toxicity Maternal: NOAEL: 27 mg/kg Teratogenicity: NOAEL: 27 mg/kg Method: Developmental Toxicity GLP: yes
Reproductive toxicity - Assessment	:	No evidence of reprotoxicity., No effects on or via lactation
Teratogenicity - Assessment	:	No evidence of adverse effects on development.

STOT - single exposure

Components:

(S)-(+)-ketamine hydrochloride:

Assessment: The substance or mixture is classified as specific target organ toxicant, single exposure, category 3 with narcotic effects.

STOT - repeated exposure

Components:

(S)-(+)-ketamine hydrochloride:

Remarks: No data available

Repeated dose toxicity

Components:

(S)-(+)-ketamine hydrochloride:

Species: Rat
NOAEL: 27 mg/kg
Application Route: intranasal (IN) administration
Exposure time: 3 month
Number of exposures: daily
Dose: 2,7 - 9 - 27 mg/kg
Subsequent observation period: 1 month
GLP: yes

Species: Dog
NOAEL: 10 mg/kg
Application Route: intranasal (IN) administration
Exposure time: 3 month
Number of exposures: daily
Dose: 3, 6, 10 mg/kg
Subsequent observation period: 1 month
GLP: yes

Species: Rat, male and female
NOAEL: 27 mg/kg
Application Route: intranasal (IN) administration
Exposure time: 6 month
Number of exposures: daily
Dose: 2,7 - 9 - 27 mg/kg
Subsequent observation period: 6 month
GLP: yes

Species: Dog, male and female
NOAEL: 10 mg/kg
Application Route: intranasal (IN) administration
Exposure time: 9 month
Number of exposures: daily
Dose: 3 - 6 - 10 mg/kg
Subsequent observation period: 9 month

GLP: yes

Species: Rat, female
Application Route: Oral
Exposure time: 2 weeks
Number of exposures: daily
Dose: 10 - 20 - 40 mg/kg
Target Organs: Liver, Kidney

Species: Rat, male
Application Route: Oral
Exposure time: 2 weeks
Number of exposures: daily
Dose: 40 - 80 - 160 mg/kg
Target Organs: Liver, Kidney

Aspiration toxicity

No data available

SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity

Components:

(S)-(+)-ketamine hydrochloride:

Toxicity to fish	:	LC50 (Oncorhynchus mykiss (rainbow trout)): 77.5 mg/l Exposure time: 96 h Test Type: static test Method: OECD Test Guideline 203 GLP: yes
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): 106.7 mg/l Exposure time: 48 h Test Type: Immobilization Method: OECD Test Guideline 202 GLP: yes
Toxicity to algae	:	ErC50 (Pseudokirchneriella subcapitata (green algae)): 90.9 mg/l End point: Growth rate Exposure time: 72 h Test Type: Growth inhibition Method: OECD Test Guideline 201 GLP: yes
Toxicity to fish (Chronic toxicity)	:	NOEC (Brachydanio rerio (zebrafish)): 0.341 mg/l Exposure time: 30 d Test Type: Fish early-life stage (FELS) toxicity test (OECD 210) Method: OECD Test Guideline 210 GLP: yes
Toxicity to daphnia and other	:	NOEC (Daphnia magna (Water flea)): 3.31 mg/l

aquatic invertebrates (Chronic toxicity)		Exposure time: 21 d Test Type: Daphnia reproduction test Method: OECD Test Guideline 211 GLP: yes
Toxicity to microorganisms	:	NOEC (activated sludge): 100 mg/l Exposure time: 3 h Test Type: Respiration inhibition Method: OECD Test Guideline 209 GLP: yes

Persistence and degradability

Components:

(S)-(+)-ketamine hydrochloride:

Biodegradability	:	aerobic Inoculum: activated sludge Result: Not readily biodegradable. Exposure time: 28 d Method: OECD Test Guideline 301B GLP: yes
Stability in water	:	Test Type: aerobic Degradation half life (DT50): 11.4 d Method: OECD Test Guideline 308 GLP: yes Remarks: Fresh water 1 Test Type: aerobic Degradation half life (DT50): 138 d Method: OECD Test Guideline 308 GLP: yes Remarks: total system 1 Test Type: aerobic Degradation half life (DT50): 20.4 d Method: OECD Test Guideline 308 GLP: yes Remarks: Fresh water 2 Test Type: aerobic Degradation half life (DT50): 230 d Method: OECD Test Guideline 308 GLP: yes Remarks: total system 2

Bioaccumulative potential

Components:

(S)-(+)-ketamine hydrochloride:

Bioaccumulation	:	Remarks: No data available

Partition coefficient: n-octanol/water	:	log Pow: 2.08 pH: 9 Method: OECD Test Guideline 107 GLP: yes

Mobility in soil

Components:

(S)-(+)-ketamine hydrochloride:

Distribution among environmental compartments	:	Adsorption/Soil Koc: 8.79 - 466.13Method: OECD Test Guideline 106

Other adverse effects

Components:

(S)-(+)-ketamine hydrochloride:

Results of PBT and vPvB assessment	:	Non-classified PBT substance Non-classified vPvB substance

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

California Prop 65

This product does not contain any chemicals known to State of California to cause cancer, birth defects, or any other reproductive harm.

Other regulations : Restricted to professional users.

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

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

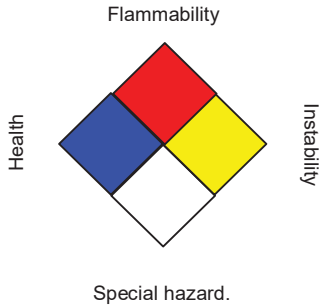
SECTION 16. OTHER INFORMATION

Full text of other abbreviations

AICS - Australian Inventory of Chemical Substances; 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; 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; 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

Further information

NFPA:



HMIS® IV:

HMIS® ratings are based on a 0-4 rating scale, with 0 representing minimal hazards or risks, and 4 representing significant hazards or risks. The "*" represents a chronic hazard, while the "/" represents the absence of a chronic hazard.

Revision Date : 2019/03/06

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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