

according to the Hazardous Products Regulations

Omarigliptin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/04/2023
4.1	09/30/2023	402445-00018	Date of first issue: 01/07/2016

SECTION 1. IDENTIFICATION

Product name	:	Omarigliptin Formulation
Other means of identification	:	No data available

Manufacturer or supplier's details

:	Merck & Co., Inc
:	126 E. Lincoln Avenue
	Rahway, New Jersey U.S.A. 07065
:	908-740-4000
:	1-908-423-6000
:	EHSDATASTEWARD@merck.com
	:

Recommended use of the chemical and restrictions on use

Recommended use	: Pl	harmaceutical
Restrictions on use	: N	ot applicable

SECTION 2. HAZARDS IDENTIFICATION

GHS classification in accordance with the Hazardous Products Regulations

Specific target organ toxicity	:	Category 2 (Stomach, Blood, Kidney)
 repeated exposure (Oral) 		

GHS label elements

Hazard pictograms



Signal Word	:	Warning

Hazard Statements : H373 May cause damage to organs (Stomach, Blood, Kidney) through prolonged or repeated exposure if swallowed.

:

Precautionary Statements

Prevention: P260 Do not breathe dust.

Response:

P314 Get medical attention if you feel unwell.

Disposal:

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

Other hazards

Dust contact with the eyes can lead to mechanical irritation. Contact with dust can cause mechanical irritation or drying of the skin. May form explosive dust-air mixture during processing, handling or other means.



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/04/2023
4.1	09/30/2023	402445-00018	Date of first issue: 01/07/2016

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

ComponentsChemical nameCommon
Name/SynonymCAS-No.Concentration (% w/w)CelluloseNo data availa-
ble9004-34-6>= 10 - < 30 *</td>OmarigliptinNo data availa-
ble1226781-44-7>= 10 - < 30 *</td>

Actual concentration or concentration range is withheld as a trade secret

SECTION 4. FIRST AID MEASURES

General advice	:	In the case of accident or if you feel unwell, seek medical advice immediately. When symptoms persist or in all cases of doubt seek medical advice.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention if symptoms occur.
In case of skin contact	:	Wash with water and soap.
		Get medical attention if symptoms occur.
In case of eye contact	:	If in eyes, rinse well with water.
		Get medical attention if irritation develops and persists.
If swallowed	:	If swallowed, DO NOT induce vomiting.
		Get medical attention if symptoms occur.
		Rinse mouth thoroughly with water.
Most important symptoms and effects, both acute and	:	May cause damage to organs through prolonged or repeated exposure if swallowed.
delayed		Contact with dust can cause mechanical irritation or drying of the skin.
		Dust contact with the eyes can lead to mechanical irritation.
Protection of first-aiders	:	First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment
Notos to physician		when the potential for exposure exists (see section 8).
Notes to physician		Treat symptomatically and supportively.

SECTION 5. FIRE-FIGHTING MEASURES

Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.
Specific hazards during fire fighting	:	Avoid generating dust; fine dust dispersed in air in sufficient concentrations, and in the presence of an ignition source is a potential dust explosion hazard. Exposure to combustion products may be a hazard to health.
Hazardous combustion prod-	:	Carbon oxides



Omarigliptin Formulation

Versio 4.1	on	Revision Date: 09/30/2023		9S Number: 2445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
U	ucts			Metal oxides	
	Specific extinguishing meth- ods		:	Use extinguishing measures that are appropriate to local cir- cumstances and the surrounding environment. Use water spray to cool unopened containers. Remove undamaged containers from fire area if it is safe to d so.	
	Special protective equipment for fire-fighters		:	Evacuate area. In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.	
SECT	ION 6.	ACCIDENTAL RELE	ASE	EMEASURES	
ti	Personal precautions, protec- tive equipment and emer- gency procedures		:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal protective equipment recommendations (see section 8).	
E	Inviron	mental precautions	:	Retain and dispos	akage or spillage if safe to do so. se of contaminated wash water. should be advised if significant spillages
		s and materials for nent and cleaning up	:	container for disper Avoid dispersal of with compressed Dust deposits sho surfaces, as these released into the a Local or national r disposal of this ma employed in the c determine which r Sections 13 and 1	dust in the air (i.e., clearing dust surfaces

SECTION 7. HANDLING AND STORAGE

Technical measures	:	Static electricity may accumulate and ignite suspended dust causing an explosion. Provide adequate precautions, such as electrical grounding and bonding, or inert atmospheres.
Local/Total ventilation	:	Use only with adequate ventilation.
Advice on safe handling	:	Do not breathe dust.
5		Do not swallow.
		Avoid contact with eyes.
		Avoid prolonged or repeated contact with skin.
		Handle in accordance with good industrial hygiene and safety
		practice, based on the results of the workplace exposure assessment
		Minimize dust generation and accumulation.
		Keep container closed when not in use.



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/04/2023
4.1	09/30/2023	402445-00018	Date of first issue: 01/07/2016
Cond	litions for safe storage rials to avoid	Keep away from Take precautio Take care to pr environment. Keep in proper Store in accord	m heat and sources of ignition. nary measures against static discharges. revent spills, waste and minimize release to the ly labeled containers. lance with the particular national regulations. th the following product types:

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis
Cellulose	9004-34-6	TWA	10 mg/m ³	CA AB OEL
		TWA (Total dust)	10 mg/m ³	CA BC OEL
		TWA (respir- able dust fraction)	3 mg/m³	CA BC OEL
		TWAEV (to- tal dust)	10 mg/m ³	CA QC OEL
		TWA	10 mg/m ³	ACGIH
Omarigliptin	1226781-44- 7	TWA	10 µg/m³	Internal
		Wipe limit	100 µg/100 cm ²	Internal

Ingredients with workplace control parameters

Engineering measures	:	Ensure adequate ventilation, especially in confined areas. Minimize workplace exposure concentrations. Apply measures to prevent dust explosions. Ensure that dust-handling systems (such as exhaust ducts, dust collectors, vessels, and processing equipment) are designed in a manner to prevent the escape of dust into the work area (i.e., there is no leakage from the equipment).
Personal protective equipment	nt	
	:	If adequate local exhaust ventilation is not available or exposure assessment demonstrates exposures outside the recommended guidelines, use respiratory protection. Particulates type
Material	:	Chemical-resistant gloves
Remarks	:	Choose gloves to protect hands against chemicals depending on the concentration specific to place of work. Breakthrough time is not determined for the product. Change gloves often! For special applications, we recommend clarifying the



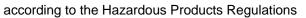
according to the Hazardous Products Regulations

Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023	SDS Number: 402445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016				
-		gloves with the breaks and at	hemicals of the aforementioned protective glove manufacturer. Wash hands before the end of workday.				
Еуе р	protection	: Wear the following personal protective equipment: Safety goggles					
Skin a	and body protection	: Skin should be washed after contact.					
Hygiene measures		 If exposure to chemical is likely during typical use, provide eye flushing systems and safety showers close to the working place. When using do not eat, drink or smoke. Wash contaminated clothing before re-use. 					

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	:	tablet
Color	:	yellow
Odor	:	No data available
Odor Threshold	:	No data available
рН	:	No data available
Melting point/freezing point	:	No data available
Initial boiling point and boiling range	:	No data available
Flash point	:	No data available
Evaporation rate	:	No data available
Flammability (solid, gas)	:	May form explosive dust-air mixture during processing, handling or other means.
Flammability (liquids)	:	No data available
Upper explosion limit / Upper flammability limit	:	No data available
Lower explosion limit / Lower flammability limit	:	No data available
Vapor pressure	:	No data available
Relative vapor density	:	No data available
Density	:	No data available
Solubility(ies) Water solubility	:	No data available
Partition coefficient: n-	:	No data available





Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023	SDS Nui 402445-		Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
	ol/water nition temperature	: No c	data available	e
Decon	nposition temperature	: No d	data available	e
	sity cosity, dynamic cosity, kinematic		data available data available	
Explos	Explosive properties		explosive	
	ing properties ular weight e size	: No c	substance o data available data available	

SECTION 10. STABILITY AND REACTIVITY

Reactivity Chemical stability Possibility of hazardous reac- tions	Not classified as a reactivity hazard. Stable under normal conditions. May form explosive dust-air mixture during proces handling or other means. Can react with strong oxidizing agents.	sing,
Conditions to avoid	Heat, flames and sparks. Avoid dust formation.	
Incompatible materials Hazardous decomposition products	Oxidizing agents No hazardous decomposition products are known	

SECTION 11. TOXICOLOGICAL INFORMATION

Information on likely routes Inhalation Skin contact Ingestion Eye contact	s of	exposure
Acute toxicity Not classified based on availa Components:	able	information.
Cellulose:		
Acute oral toxicity	:	LD50 (Rat): > 5,000 mg/kg
Acute inhalation toxicity	:	LC50 (Rat): > 5.8 mg/l Exposure time: 4 h Test atmosphere: dust/mist
		6/1/



according to the Hazardous Products Regulations

Omarigliptin Formulation

/ersion I.1	Revision Date: 09/30/2023	SDS Num 402445-0		Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
Acute	dermal toxicity	: LD50	(Rabbit): >	2,000 mg/kg
	i gliptin: oral toxicity	: LD50	(Rat): 750 ı	mg/kg
	corrosion/irritation	vailable informa	ation.	
<u>Comp</u>	onents:			
Omar i Result	igliptin:	: No ski	in irritation	
	us eye damage/eye assified based on av		ation.	
<u>Comp</u>	onents:			
	igliptin:			
Specie Result		-	e cornea e irritation	
Respi	ratory or skin sens	sitization		
	ensitization	vailable informa	ation.	
•	ratory sensitization		ation.	
<u>Comp</u>	onents:			
Omari Test T Specie Asses Result	es sment	: Mouse	e not cause s	e assay (LLNA) skin sensitization.
	cell mutagenicity assified based on av	vailable informa	ation.	
<u>Comp</u>	onents:			
Cellul	ose:			
Genot	oxicity in vitro		ype: Bacte t: negative	rial reverse mutation assay (AMES)
			ype: In vitro t: negative	o mammalian cell gene mutation test
Genot	oxicity in vivo	: Test T	ype: Mamr	nalian erythrocyte micronucleus test (in vi
			7 / 14	



according to the Hazardous Products Regulations

Omarigliptin Formulation

Component Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	ry in vitro ry in vivo nicity ed based on ava <u>nts:</u> Route	Result: negativ Test Type: Chr Test system: C Result: negativ : Test Type: Mut cytogenetic tes Species: Rat	e ute: Ingestion e cterial reverse mutation assay (AMES) e comosome aberration test in vitro chinese hamster ovary cells e cagenicity (in vivo mammalian bone-marrov ct, chromosomal analysis) ute: Intraperitoneal injection
Genotoxicit Genotoxicit Carcinoge Not classifie Componer Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result	ry in vitro ry in vivo nicity ed based on ava <u>nts:</u> Route	Result: negativ Test Type: Chr Test system: C Result: negativ : Test Type: Mut cytogenetic tes Species: Rat Application Ro Result: negativ ailable information. : Rat : Ingestion : 72 weeks	re comosome aberration test in vitro chinese hamster ovary cells re tagenicity (in vivo mammalian bone-marrov it, chromosomal analysis) ute: Intraperitoneal injection
Genotoxicit Carcinoge Not classifie Componer Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	y in vivo nicity ed based on ava <u>its:</u> Route	Result: negativ Test Type: Chr Test system: C Result: negativ : Test Type: Mut cytogenetic tes Species: Rat Application Ro Result: negativ ailable information. : Rat : Ingestion : 72 weeks	re comosome aberration test in vitro chinese hamster ovary cells re tagenicity (in vivo mammalian bone-marrov it, chromosomal analysis) ute: Intraperitoneal injection
Carcinoge Not classifie Componer Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	nicity ed based on ava nts: Route	Test system: C Result: negativ : Test Type: Mut cytogenetic tes Species: Rat Application Ro Result: negativ ailable information. : Rat : Ingestion : 72 weeks	chinese hamster ovary cells e cagenicity (in vivo mammalian bone-marrov t, chromosomal analysis) ute: Intraperitoneal injection
Carcinoge Not classifie Componer Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	nicity ed based on ava nts: Route	cytogenetic tes Species: Rat Application Ro Result: negativ ailable information. : Rat : Ingestion : 72 weeks	it, chromosomal analysis) ute: Intraperitoneal injection
Not classifie <u>Componer</u> <u>Cellulose:</u> Species Application Exposure ti Result <u>Omariglipt</u> Species Application Exposure ti Result Species Application	ed based on ava nts: Route	: Rat : Ingestion : 72 weeks	
Component Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	nts: Route	: Rat : Ingestion : 72 weeks	
Cellulose: Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application	Route	: Ingestion : 72 weeks	
Species Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application		: Ingestion : 72 weeks	
Application Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application		: Ingestion : 72 weeks	
Exposure ti Result Omariglipt Species Application Exposure ti Result Species Application		: 72 weeks	
Omariglipt Species Application Exposure ti Result Species Application		· negative	
Species Application Exposure ti Result Species Application		. nogative	
Application Exposure ti Result Species Application	in:		
Exposure ti Result Species Application		: Rat	
Result Species Application		: Oral : 2 Years	
Species Application	me	: 20 mg/kg body	weight
Application		: negative	
Application		: Mouse	
F		: Oral	
Exposure ti	me	: 2 Years	weight
Result		: 20 mg/kg body : negative	weight
Reproduct	ive toxicity		
-	-	ailable information.	
Componer	nts:		
Cellulose:			
Effects on f	ertility	: Test Type: One Species: Rat Application Ro	e-generation reproduction toxicity study



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023		DS Number: 2445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
Effec	ts on fetal development	:	Result: negative Test Type: Fertilit Species: Rat Application Route Result: negative	y/early embryonic development : Ingestion
	rigliptin: ts on fertility	:	Species: Rat Application Route	y/early embryonic development : Oral 100 mg/kg body weight
Effec	ts on fetal development	:	Species: Rabbit Developmental To Result: No effects Test Type: Embry Species: Rat Application Route Developmental To Result: Reduced food consumption	o-fetal development exicity: NOAEL: > 50 mg/kg body weight on fetal development. o-fetal development : Oral exicity: LOAEL: 100 mg/kg body weight offspring weight gain., Reduced maternal ., Skeletal malformations. ects were seen only at maternally toxic dos-

STOT-single exposure

Not classified based on available information.

STOT-repeated exposure

May cause damage to organs (Stomach, Blood, Kidney) through prolonged or repeated exposure if swallowed.

Components:

Omarigliptin:

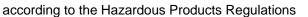
Routes of exposure		Ingestion Stomach Blood Kidney
Target Organs	•	Stomach, Blood, Kidney
Assessment	:	May cause damage to organs through prolonged or repeated exposure.

Repeated dose toxicity

Components:

Cellulose:

Species	:	Rat
NOAEL	:	>= 9,000 mg/kg
Application Route	:	Ingestion
Exposure time	:	90 Days





Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023	SDS Number: 402445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
Spec NOA Appli	EL cation Route sure time	: Rat : 100 mg/kg : Oral : 90 Days : No significant	adverse effects were reported
Expo	EL	: Rat : 10 mg/kg : 100 mg/kg : Oral : 180 Days : Blood, Kidney	/
Expo	EL	: Dog : 10 mg/kg : 75 mg/kg : Oral : 40 Days : Stomach	
Expo	EL	: Dog : 10 mg/kg : 75 mg/kg : Oral : 270 Days : Stomach	
	EL cation Route sure time	: Monkey : 9 mg/kg : Oral : 90 Days : No significant	adverse effects were reported

Aspiration toxicity

Not classified based on available information.

Experience with human exposure

Components:

Omarigliptin:

Ingestion

: Symptoms: Headache, stomach discomfort, Dizziness, Tiredness, Diarrhea, flu-like symptoms, Back pain, Vomiting, chills

SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity	
Components:	
Cellulose:	
Toxicity to fish	: LC50 (Oryzias latipes (Japanese medaka)): > 100 mg/l



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023	-	0S Number: 2445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
			Exposure time: 48 Remarks: Based o	h on data from similar materials
Тох	arigliptin: icity to daphnia and other atic invertebrates	:	EC50 (Daphnia m Exposure time: 48 Method: OECD Te	
			EC50 (Americamy Exposure time: 96 Method: US-EPA	5 h
Tox plar	icity to algae/aquatic its	:	EC50 (Pseudokiro mg/l Exposure time: 72 Method: OECD Te	
			NOEC (Pseudokir mg/l Exposure time: 72 Method: OECD Te	
Tox icity	icity to fish (Chronic tox-)	:	NOEC (Pimephale Exposure time: 32 Method: OECD Te	
aqu	icity to daphnia and other atic invertebrates (Chron- xicity)	:	NOEC (Daphnia n Exposure time: 21 Method: OECD Te	
Тох	icity to microorganisms	:	EC50: > 1,000 mg Exposure time: 3 l Test Type: Respire Method: OECD Te	n ation inhibition
			NOEC: 0.1 mg/l Exposure time: 3 l Test Type: Respir Method: OECD Te	ation inhibition
Per	sistence and degradabili	ity		
<u>Cor</u>	nponents:			
	lulose:		Desult De III - 1	
BIO	degradability	:	Result: Readily bio	Daegradable.
	arigliptin: degradability	:	Result: Not readily Biodegradation: 5 Exposure time: 11 Method: OECD Te	50 % d



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version 4.1	Revision Date: 09/30/2023		DS Number: 2445-00018	Date of last issue: 04/04/2023 Date of first issue: 01/07/2016
Bioaco	cumulative potential			
Comp	onents:			
Partitic	gliptin: on coefficient: n- I/water	:	log Pow: 0.525	
Mobili	ty in soil			
Comp	onents:			
Distrib	gliptin: ution among environ- compartments	:	log Koc: 4.01 Method: OECD T	est Guideline 106
•	adverse effects a available			
SECTION 1	3. DISPOSAL CONSI	DEF	RATIONS	
Dispos	sal methods			

Waste from residues	: Do not dispose of waste into sewer. Dispose of in accordance with local regulations.
Contaminated packaging	 Empty containers should be taken to an approved waste handling site for recycling or disposal.
	If not otherwise specified. Dispose of as unused product.

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.

Domestic regulation

TDG Not regulated as a dangerous good

Special precautions for user

Not applicable



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/04/2023
4.1	09/30/2023	402445-00018	Date of first issue: 01/07/2016

SECTION 15. REGULATORY INFORMATION

The ingredients of this product are reported in the following inventories:					
AICS	:	not determined			
DSL	:	not determined			
IECSC	:	not determined			

SECTION 16. OTHER INFORMATION

Full text of other abbreviations

ACGIH		USA. ACGIH Threshold Limit Values (TLV)
CA AB OEL	:	Canada. Alberta, Occupational Health and Safety Code (table 2: OEL)
CA BC OEL	:	Canada. British Columbia OEL
CA QC OEL	:	Québec. Regulation respecting occupational health and safe- ty, Schedule 1, Part 1: Permissible exposure values for air- borne contaminants
ACGIH / TWA	:	8-hour, time-weighted average
CA AB OEL / TWA	:	8-hour Occupational exposure limit
CA BC OEL / TWA	:	8-hour time weighted average
CA QC OEL / TWAEV	:	Time-weighted average exposure value

AIIC - Australian Inventory of Industrial Chemicals; ANTT - National Agency for Transport by Land of Brazil; ASTM - American Society for the Testing of Materials; bw - Body weight; CMR -Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; 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: 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; n.o.s. - Not Otherwise Specified; Nch - Chilean Norm; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NOM - Official Mexican Norm; 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; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TDG - Transporta-



according to the Hazardous Products Regulations

Omarigliptin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/04/2023
4.1	09/30/2023	402445-00018	Date of first issue: 01/07/2016

tion of Dangerous Goods; 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; WHMIS - Workplace Hazardous Materials Information System

Sources of key data used to compile the Material Safety Data Sheet	:	Internal technical data, data from raw material SDSs, OECD eChem Portal search results and European Chemicals Agen- cy, http://echa.europa.eu/
Revision Date Date format	:	09/30/2023 mm/dd/yyyy

The information provided in this Safety Data Sheet is correct to the best of our knowledge, information and belief at the date of its publication. The information is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and shall not be considered a warranty or quality specification of any type. The information provided relates only to the specific material identified at the top of this SDS and may not be valid when the SDS material is used in combination with any other materials or in any process, unless specified in the text. Material users should review the information and recommendations in the specific context of their intended manner of handling, use, processing and storage, including an assessment of the appropriateness of the SDS material in the user's end product, if applicable.

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