

Version 3.8	Revision Date: 10.10.2020		S Number: 612-00012	Date of last issue: 23.03.2020 Date of first issue: 07.01.2016
Section 1:	Identification			
Produ	ct name	:	Olmesartan / Hyd	drochlorothiazide Formulation
Manu	facturer or supplier's d	etai	ls	
Comp	any	:	Organon & Co.	
Addre	SS	:	30 Hudson Stree Jersey City, New	t, 33nd floor Jersey, U.S.A 07302
Telepl	hone	:	551-430-6000	
Emerç	gency telephone number	:	215-631-6999	
E-mai	l address	:	EHSSTEWARD@	⊉organon.com
	mmended use of the ch nmended use	nemi :	i cal and restrictic Pharmaceutical	ons on use
Section 2:	Hazard identification			
GHS (Classification			
Repro	ductive toxicity	:	Category 1A	
	fic target organ toxicity - ted exposure	:	Category 2 (Kidn	ey, Parathyroid gland)
GHS I	abel elements			
Hazar	d pictograms	:		
Signal	l word	:	Danger	
Hazar	d statements	:	H373 May cause	age the unborn child. damage to organs (Kidney, Parathyroid olonged or repeated exposure.
Preca	utionary statements	:	P202 Do not han and understood. P260 Do not brea P281 Use person Response:	cial instructions before use. dle until all safety precautions have been rea athe dust. hal protective equipment as required. exposed or concerned: Get medical advice/



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P405 Store locked up.

Disposal:

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

Other hazards which do not result in classification

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.

Section 3: Composition/information on ingredients

Substance / Mixture : Mixture

Components

Chemical name	CAS-No.	Concentration (% w/w)
Olmesartan	144689-63-4	>= 0.3 -< 10
Cellulose	9004-34-6	< 10
Hydrochlorothiazide	58-93-5	>= 1 -< 10

Section 4: First-aid measures

General advice	:	In the case of accident or if you feel unwell, seek medical ad- vice immediately. When symptoms persist or in all cases of doubt seek medical advice.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention.
In case of skin contact	:	In case of contact, immediately flush skin with soap and plenty of water. Remove contaminated clothing and shoes. Get medical attention. Wash clothing before reuse. Thoroughly clean shoes before reuse.
In case of eye contact	:	
If swallowed	:	
Most important symptoms and effects, both acute and delayed	:	May damage the unborn child. May cause damage to organs through prolonged or repeated exposure. Contact with dust can cause mechanical irritation or drying of the skin.
Protection of first-aiders	:	Dust contact with the eyes can lead to mechanical irritation. First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment when the potential for exposure exists (see section 8).
Notes to physician	:	Treat symptomatically and supportively.

Section 5: Fire-fighting measures

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Suitable extinguishing media		:	Water spray Alcohol-resistant Carbon dioxide (C Dry chemical		
	Unsuita media	able extinguishing	:	None known.	
	Specific fighting	c hazards during fire-	:	concentrations, and potential dust exp	dust; fine dust dispersed in air in sufficient nd in the presence of an ignition source is a losion hazard. pustion products may be a hazard to health.
	Hazard ucts	ous combustion prod-	:	Carbon oxides Nitrogen oxides (I Chlorine compour Sulphur oxides	
	Specific ods	c extinguishing meth-	:	cumstances and t Use water spray t	measures that are appropriate to local cir- he surrounding environment. o cool unopened containers. ged containers from fire area if it is safe to do
	Special for firef	protective equipment ighters	:	In the event of fire	e, wear self-contained breathing apparatus. ective equipment.

Section 6: Accidental release measures

Personal precautions, protec- tive equipment and emer- gency procedures	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal pro- tective equipment recommendations (see section 8).
Environmental precautions	:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Retain and dispose of contaminated wash water. Local authorities should be advised if significant spillages cannot be contained.
Methods and materials for containment and cleaning up	:	Sweep up or vacuum up spillage and collect in suitable con- tainer for disposal. Avoid dispersal of dust in the air (i.e., clearing dust surfaces with compressed air). Dust deposits should not be allowed to accumulate on surfac- es, as these may form an explosive mixture if they are re- leased into the atmosphere in sufficient concentration. Local or national regulations may apply to releases and dis- posal of this material, as well as those materials and items employed in the cleanup of releases. You will need to deter- mine which regulations are applicable. Sections 13 and 15 of this SDS provide information regarding certain local or national requirements.

Section 7: Handling and storage

- Technical measures
- : Static electricity may accumulate and ignite suspended dust



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Local/Total ventilation		and bonding, o	blosion. ate precautions, such as electrical grounding or inert atmospheres. htilation is unavailable, use with local exhaust
Advice on safe handling		Do not breathe Do not swallow Avoid contact Wash skin tho Handle in acco practice, base sessment Keep containe Keep containe Keep away fro Take precautio Do not eat, dri	v.
Hygi	ene measures	flushing syster place. When using do Wash contami The effective of engineering co appropriate de industrial hygie	chemical is likely during typical use, provide eye ns and safety showers close to the working o not eat, drink or smoke. nated clothing before re-use. operation of a facility should include review of ontrols, proper personal protective equipment, gowning and decontamination procedures, ene monitoring, medical surveillance and the trative controls.
	ditions for safe storage erials to avoid	Store locked u Keep tightly cl Store in accord	osed. dance with the particular national regulations. ith the following product types:
		Strong oxidizir	ig agents

Section 8: Exposure controls/personal protection

Components with workplace control parameters

:

Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis
Olmesartan	144689-63-4	TWA	30 µg/m3 (OEB 3)	Internal
		Wipe limit	300 µg/100 cm ²	Internal
Cellulose	9004-34-6	WES-TWA	10 mg/m3	NZ OEL
		TWA	10 mg/m3	ACGIH
Hydrochlorothiazide	58-93-5	TWA	100 µg/m3 (OEB 2)	Internal

Engineering measures

All engineering controls should be implemented by facility design and operated in accordance with GMP principles to



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		Containment are required t				
Pers	onal protective equipr	nent				
Resp	iratory protection	sure assessm	: If adequate local exhaust ventilation is not available or exposure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection.			
	lter type I protection	: Particulates type				
M	aterial	: Chemical-res	istant gloves			
	emarks protection	If the work en mists or aeros Wear a faces	ble gloving. glasses with side shields or goggles. vironment or activity involves dusty conditions, sols, wear the appropriate goggles. hield or other full face protection if there is a lirect contact to the face with dusts, mists, or			
Skin	and body protection	: Work uniform Additional boot task being pe posable suits	or laboratory coat. dy garments should be used based upon the rformed (e.g., sleevelets, apron, gauntlets, dis-) to avoid exposed skin surfaces. ate degowning techniques to remove potentially clothing.			

Section 9: Physical and chemical properties

Appearance	:	powder
Colour	:	white to off-white
Odour	:	No data available
Odour 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	:	Not applicable
Evaporation rate	:	Not applicable
Flammability (solid, gas)	:	May form explosive dust-air mixture during processing, han- dling or other means.
Flammability (liquids)	:	No data available

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		explosion limit / Upper bility limit	:	No data available)
		explosion limit / Lower bility limit	:	No data available	
	Vapour	pressure	:	Not applicable	
	Relative	e vapour density	:	Not applicable	
	Relative	e density	:	No data available)
	Density		:	No data available)
	Solubili Wat	ty(ies) er solubility	:	No data available	9
	Partition octanol	n coefficient: n-	:	Not applicable	
		nition temperature	:	No data available)
	Decom	position temperature	:	No data available)
	Viscosi Visc	ty osity, kinematic	:	Not applicable	
	Explosi	ve properties	:	Not explosive	
	Oxidizir	ng properties	:	The substance of	r mixture is not classified as oxidizing.
	Molecu	lar weight	:	Not applicable	
	Particle	size	:	No 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 processing, han- dling or other means. Can react with strong oxidizing agents.
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

Exposure routes	:	Inhalation
		Skin contact



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			Ingestion Eye contact	
	e toxicity	- 1 -		
Not ci Produ	assified based on availa	ble	information.	
	oral toxicity	:	Acute toxicity e Method: Calcu	estimate: > 2,000 mg/kg lation method
<u>Comp</u>	oonents:			
Olme	sartan:			
Acute	oral toxicity	:	LD50 (Rat): > 2	2,000 mg/kg
			LD50 (Mouse)	: > 2,000 mg/kg
			LD50 (Dog): >	1,500 mg/kg
Acute	inhalation toxicity	:	Remarks: No o	lata available
Acute	dermal toxicity	:	Remarks: No c	data available
Cellu	lose:			
Acute	oral toxicity	:	LD50 (Rat): >	5,000 mg/kg
Acute	inhalation toxicity	:	LC50 (Rat): > Exposure time Test atmosphe	:4h
Acute	e dermal toxicity	:	LD50 (Rabbit):	> 2,000 mg/kg
Hydro	ochlorothiazide:			
Acute	oral toxicity	:	LD50 (Rat): > 2	2,750 mg/kg
			LD50 (Mouse)	: > 2,830 mg/kg
	toxicity (other routes of histration)	:	LD50 (Rat): 99 Application Ro	0 mg/kg ute: Intravenous
			LD50 (Mouse) Application Ro	: 590 mg/kg ute: Intravenous
	corrosion/irritation lassified based on availa	bla	information	
	oonents:	UIG		
	sartan:			
Rema		:	No data availa	ble

Hydrochlorothiazide:



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Spec Resu		: Rabbit : No skin irritatio	n
	ous eye damage/eye lassified based on av		
Com	ponents:		
Olme	esartan:		
Spec Resu Meth	lt	: Rabbit : Moderate eye i : Draize Test	ritation
Hydr	ochlorothiazide:		
Spec Resu		: Rabbit : Mild eye irritatio	n
Resp	iratory or skin sens	tisation	
	sensitisation lassified based on av	ailable information.	
-	iratory sensitisatior lassified based on av		
Com	ponents:		
	e sartan: sure routes arks	: Skin contact : No data availat	le
Chro	nic toxicity		
	n cell mutagenicity lassified based on av	ailable information.	
Com	ponents:		
	esartan: otoxicity in vitro	: Test Type: Bac Result: negative	terial reverse mutation assay (AMES) e
		Test Type: Mut Result: negative	agenicity (in vitro mammalian cytogenetic test) e
			omosome aberration test in vitro hinese hamster lung cells
		Test Type: Mou Result: negative	
Geno	otoxicity in vivo	: Test Type: Mice Species: Mouse	



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		Cell type: Bone n Application Route Result: negative	
	cell mutagenicity -	: Weight of eviden cell mutagen.	ce does not support classification as a germ
Cellu	lose:		
Geno	toxicity in vitro	: Test Type: Bacte Result: negative	erial reverse mutation assay (AMES)
		Test Type: In vitr Result: negative	o mammalian cell gene mutation test
Geno	toxicity in vivo	: Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative	
Hydro	ochlorothiazide:		
Geno	toxicity in vitro	: Test Type: Bacte Result: negative	erial reverse mutation assay (AMES)
			nosomal aberration inese hamster ovary cells
			chromatid exchange assay inese hamster ovary cells
		Test Type: in vitre Test system: more Result: positive	o assay use lymphoma cells
Geno	toxicity in vivo	: Test Type: Chror Species: Chinese Cell type: Bone n Result: negative	
		Test Type: in vivo Species: Mouse Cell type: Bone n Result: negative	
	cell mutagenicity -	: Weight of eviden cell mutagen.	ce does not support classification as a germ

Carcinogenicity

Not classified based on available information.



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<u>Comp</u>	oonents:			
Olme	sartan:			
Speci	es	:	Rat	
	ation Route	:	Oral	
Expos Resul	sure time t	:	2 Years negative	
Speci		:	Mouse	
	ation Route	:	Oral	
Expos Resul	sure time t	:	6 Months negative	
Cellul	lose:			
Speci		:	Rat	
	ation Route	:	Ingestion	
Expos Resul	sure time t	:	72 weeks negative	
Resul	L .	•	negative	
-	ochlorothiazide:			
Speci	es ation Route	:	Mouse, female Oral	
	sure time	:	2 Years	
Resul		:	negative	
Speci	es	:	Mouse, male	
	ation Route	:	Oral	
Expos Resul	sure time	:	2 Years equivocal	
		•		
Speci	es ation Route	:	Rat, male and fe Oral	male
	sure time	:	2 Years	
Resul		:	negative	
Repro	oductive toxicity			
May d	lamage the unborn child	d.		
<u>Comp</u>	oonents:			
	sartan:			
Effect	s on fertility	:	Test Type: Fertil Species: Rat	ity
			Application Rout	e: Oral
			Fertility: NOAEL:	: 1,000 mg/kg body weight
			Result: No effect	s on fertility
Effect	s on foetal develop-	:	Test Type: Deve	lopment
ment	·		Species: Rat	
			Application Rout	e: Oral gram per kilogram
			Result: No terato	



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		Test Type: Development Species: Rabbit Application Route: Oral Dose: 1 milligram per kilogram Result: No teratogenic effects	
		Test Type: Development Species: Rat Application Route: Oral Developmental Toxicity: LOAEL: >= 1.6 mg/kg body weigh Symptoms: Malformations were observed., Reduced body weight Result: Effects on postnatal development	
	productive toxicity - As- sment	: Positive evidence of adverse effects on development from human epidemiological studies.	ı
Cel	lulose:		
Effe	ects on fertility	: Test Type: One-generation reproduction toxicity study Species: Rat Application Route: Ingestion Result: negative	
Effe me	ects on foetal develop- nt	: Test Type: Fertility/early embryonic development Species: Rat Application Route: Ingestion Result: negative	
Цv	drochlorothiazide:		
-	ects on fertility	: Test Type: Fertility Species: Rat, male and female Application Route: oral (feed) Fertility: NOAEL: 4 mg/kg body weight Result: Effects on fertility	
		Test Type: Fertility Species: Mouse, male and female Application Route: oral (feed) Fertility: NOAEL: 100 mg/kg body weight Result: Effects on fertility	
Effe me	ects on foetal develop- nt	 Test Type: Development Species: Mouse Application Route: Oral Developmental Toxicity: NOAEL: 3,000 mg/kg body weigh Result: No teratogenic effects 	ht
		Test Type: Development Species: Rat Application Route: Oral Developmental Toxicity: NOAEL: 1,000 mg/kg body weigh Result: No teratogenic effects	ht



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STOT	- single exposure			
Not c	lassified based on ava	ailable i	nformation.	
	repeated exposur cause damage to orga		Iney, Parathyro	oid gland) through prolonged or repeated expo
	oonents:			
Hydro	ochlorothiazide:			
Targe	et Organs ssment		Kidney, Paratł Causes dama exposure.	nyroid gland ge to organs through prolonged or repeated
Repe	ated dose toxicity			
<u>Com</u>	oonents:			
Olme	sartan:			
Speci			Rat	
NOAE	L cation Route		2,000 mg/kg Oral	
	sure time		24 Months	
Rema		:	No significant	adverse effects were reported
Cellu	lose:			
Speci		:	Rat	
NOAE	EL cation Route		>= 9,000 mg/k Ingestion	g
	sure time		90 Days	
Hydro	ochlorothiazide:			
Speci		:	Rat, male and	female
LOAE		:	10 mg/kg	
	cation Route sure time	:	Oral 2 yr	
	et Organs	:	Kidney, Parath	nyroid gland
Speci	es	:	Mouse, male a	and female
NOAE	ΞL	:	300 - 550 mg/	
	cation Route	:	Oral	
Expos	sure time arks	:	2 yr No significant	adverse effects were reported
Speci	es	:	Dog 50 - 200 mg/kg	n
Applia	cation Route	:	Oral	3
	sure time		9 Months	
Targe	et Organs	:	Parathyroid gl	and
Aspir	ation toxicity			

Not classified based on available information.



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<u>Com</u>	oonents:					
-	ochlorothiazide: piration toxicity classific	atio	n			
Expe	rience with human exp	osi	ire			
Com	oonents:					
Olme	sartan:					
Eye c Inges	ontact tion	:	 Symptoms: Eye irritation Symptoms: hypotension Remarks: May cause harm to the unborn child. Based on Human Evidence 			
Hydro	ochlorothiazide:					
Eye c Inges	ontact tion	:	Symptoms: Eye irritation Symptoms: Dizziness, Headache, Fatigue, Nausea, Ab- dominal pain, hypotension, dry mouth, electrolyte imbalanc eye pain			
ection 12	2: Ecological informati	on				
Ecoto	oxicity					
<u>Com</u>	oonents:					
Cellu	lose:					
Toxic	ity to fish	:	Exposure time: 4	tipes (Japanese medaka)): > 100 mg/l 8 h on data from similar materials		
Hydro	ochlorothiazide:					
Toxic	ity to fish	:	LC50 (Pimephale Exposure time: 9	es promelas (fathead minnow)): > 500 mg/l 6 h		
	ity to daphnia and other ic invertebrates	:	EC50 (Daphnia r Exposure time: 4	nagna (Water flea)): > 500 mg/l 8 h		
Persi	stence and degradabil	ity				
Com	oonents:					
Cellu	lose:					
Biode	gradability	:	Result: Readily b	iodegradable.		
-	ochlorothiazide: ity in water	:	Hydrolysis: 46.2	%(96 h)		
	ccumulative potential ata available					



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Mobil	ity in soil					
No da	ta available					
	adverse effects ata available					
Section 13	3: Disposal consider	ations				
Dispo	osal methods					
	e from residues minated packaging	: Empty contain dling site for r	accordance with local regulations. ners should be taken to an approved waste han- recycling or disposal. se specified: Dispose of as unused product.			
Section 14	4: Transport informa	tion				
Interr	national Regulations					
UNRT Not re	DG egulated as a dangero	us good				
IATA- Not re	DGR egulated as a dangero	us good				
-	-Code egulated as a dangero	us good				
	port in bulk accordi	-	ARPOL 73/78 and the IBC Code			
Natio	nal Regulations					
NZS : Not re	5433 egulated as a dangero	us good				
Section 1	5: Regulatory inform	ation				
Safet ture	y, health and enviro	nmental regulations	/legislation specific for the substance or mix-			
) Approval Number 00425 Pharmaceutic	al Active Ingredients	Group Standard 2017			
	Controls	-				
Certified handler certificate not required. Tracking hazardous substance not required. Refer to the Health and Safety at Work (Hazardous Substances) Regulations 2017, for further in- formation.						

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

not determined
r

DSL	:	not determined
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IEC	CSC	:	not determined	
Section	16: Other information			
Fu	rther information			
CO	ources of key data used to mpile the Safety Data leet	:		data, data from raw material SDSs, OECD arch results and European Chemicals Agen- ropa.eu/
Da	ate format : dd.mm.yyyy			
Fu	Il text of other abbreviati	ons		
	CGIH Z OEL	:		eshold Limit Values (TLV) orkplace Exposure Standards for Atmospher-
	CGIH / TWA 2 OEL / WES-TWA	:	 8-hour, time-weighted average Workplace Exposure Standard - Time Weighted average 	

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 - Transportation of Dangerous Goods; 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

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



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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.

NZ / EN