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

1.1 Product identifier Trade name	: Doravirine / Lamivudine / Tenofovir Disoproxil Fumarate Bi- layer Formulation
1.2 Relevant identified uses of t Use of the Sub- stance/Mixture	he substance or mixture and uses advised against : Pharmaceutical
Recommended restrictions on use	: Not applicable
1.3 Details of the supplier of the Company	e safety data sheet : MSD 120 Moorgate EC2M 6UR London, United Kingdom
Telephone	: +44 (0) 2081548000

E-mail address of person		EHSDATASTEWARD@msd.com
responsible for the SDS		

1.4 Emergency telephone number

1-908-423-6000

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008) as amended by GB-CLP Regulation, UK SI 2019/720, and UK SI 2020/1567)

Eye irritation, Category 2 Reproductive toxicity, Category 2 Specific target organ toxicity - repeated exposure, Category 2 H319: Causes serious eye irritation. H361d: Suspected of damaging the unborn child. H373: May cause damage to organs through prolonged or repeated exposure.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008) as amended by GB-CLP Regulation, UK SI 2019/720, and UK SI 2020/1567)



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Haz	ard pictograms		
Sigr	al word	: Warning	•
Haz	ard statements	: H319 H361d H373	Causes serious eye irritation. Suspected of damaging the unborn child. May cause damage to organs through prolonged or repeated exposure.
Prec	autionary statements	: Preventio P201 P260 P264 P280	Don: Obtain special instructions before use. Do not breathe dust. Wash skin thoroughly after handling. Wear protective gloves/ protective clothing/ eye protection/ face protection.
		Respons P308 + P P337 + P	313 IF exposed or concerned: Get medical advice/ attention.

Hazardous components which must be listed on the label: Lamivudine Tenofovir

2.3 Other hazards

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

May form explosive dust-air mixture during processing, handling or other means.

SECTION 3: Composition/information on ingredients

3.2 Mixtures

Components

Chemical name	CAS-No. EC-No. Index-No. Registration number	Classification	Concentration (% w/w)
Lamivudine	134678-17-4	Repr. 2; H361d STOT RE 2; H373 (Blood)	>= 10 - < 20
Tenofovir	202138-50-9	Acute Tox. 4; H302 Eye Irrit. 2; H319 STOT RE 2; H373 (Bone, Kidney)	>= 10 - < 20



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Dorav	virine	1338225-9	7-0 >= 1 - < 10
Subst	tances with a workpla	ce exposure limit :	
Cellul	ose	9004-34-6	>= 20 - < 30
		232-674-9	

For explanation of abbreviations see section 16.

SECTION 4	4: First aid	measures
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4.1 Description of 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.			
Protection of first-aiders :	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).			
If inhaled :	If inhaled, remove to fresh air. Get medical attention.			
In case of skin contact :	In case of contact, immediately flush skin with 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 :	In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. If easy to do, remove contact lens, if worn. Get medical attention.			
If swallowed :	If swallowed, DO NOT induce vomiting. Get medical attention. Rinse mouth thoroughly with water.			
4.2 Most important symptoms and effects, both acute and delayed				
Risks :	Causes serious eye irritation. Suspected of damaging the unborn child. May cause damage to organs through prolonged or repeated exposure.			
4.3 Indication of any immediate medical attention and special treatment needed				
Treatment :	Treat symptomatically and supportively.			



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SECTION 5: Firefighting measures

5.1 Extinguishing media

Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.

5.2 Special hazards arising from the substance or mixture

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- ucts	:	Carbon oxides Nitrogen oxides (NOx) Halogenated compounds Metal oxides			
5.3 Advice for firefighters					
Special protective equipment for firefighters	: :	In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.			
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 do so. Evacuate area.			

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

• •		
Personal precautions	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal pro-
		tective equipment recommendations (see section 8).

6.2 Environmental precautions

:	Avoid release to the environment.
	Prevent further leakage or spillage if safe to do so.
	Retain and dispose of contaminated wash water.
	If spillage enters rivers or watercourses, inform the Environ-
	ment Agency (emergency telephone number 0800 807060).
	:



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6.3 Methods and material for containment and cleaning up

Methods for cleaning up	ta A C e le L P e S	weep up or vacuum up spillage and collect in suitable con- ainer for disposal. woid dispersal of dust in the air (i.e., clearing dust surfaces with compressed air). bust deposits should not be allowed to accumulate on surfac- s, as these may form an explosive mixture if they are re- eased into the atmosphere in sufficient concentration. ocal or national regulations may apply to releases and dis- osal of this material, as well as those materials and items mployed in the cleanup of releases. You will need to deter- nine which regulations are applicable. eections 13 and 15 of this SDS provide information regarding ertain local or national requirements.

6.4 Reference to other sections

See sections: 7, 8, 11, 12 and 13.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

T	
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 get on skin or clothing.
	Do not breathe dust.
	Do not swallow.
	Do not get in eyes.
	Wash skin thoroughly after handling.
	Handle in accordance with good industrial hygiene and safety
	practice, based on the results of the workplace exposure as-
	sessment
	Minimize dust generation and accumulation.
	Keep container closed when not in use.
	Keep away from heat and sources of ignition.
	Take precautionary measures against static discharges.
	Take care to prevent spills, waste and minimize release to the
	environment.
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 contami-
	nated clothing before re-use.
	The effective operation of a facility should include review of
	engineering controls, proper personal protective equipment,
	appropriate degowning and decontamination procedures,
	industrial hygiene monitoring, medical surveillance and the
	use of administrative controls.



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7.2 Cond	itions for safe storage,	inc	luding any incom	patibilities
	irements for storage s and containers	:		labelled containers. Store locked up. Store in the particular national regulations.
Advi	ce on common storage	:	Do not store with Strong oxidizing	the following product types: agents
•	f ic end use(s) ific use(s)	:	No data available	9

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

dust of any kind	10 mg/m3 Value type (Form of exposure): TWA (Inhalable) Basis: GB EH40
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4 mg/m3 Value type (Form of exposure): TWA (Respirable fraction) Basis: GB EH40

Components	CAS-No.	Value type (Form of exposure)	Control parameters	Basis
Cellulose	9004-34-6	TWA (inhalable dust)	10 mg/m3	GB EH40
		TWA (Respirable dust)	4 mg/m3	GB EH40
		STEL (inhalable dust)	20 mg/m3	GB EH40
Lamivudine	134678-17- 4	TWA	100 µg/m3 (OEB 2)	Internal
Tenofovir	202138-50- 9	TWA	150 ug/m3 (OEB 2)	Internal
Doravirine	1338225- 97-0	TWA	500 ug/m3 (OEB2)	Internal

8.2 Exposure controls

Engineering measures

Use feasible engineering controls to minimize exposure to compound. All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment.

Personal protective equipment

Eye/face protection

Wear safety glasses with side shields or goggles.

:



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Hanc	I protection	Wear a face	osols, wear the appropriate goggles. shield or other full face protection if there is a direct contact to the face with dusts, mists, or
Ma	aterial	: Chemical-res	sistant gloves
	and body protection iratory protection	: If adequate I sure assessi ommended g	n or laboratory coat. ocal exhaust ventilation is not available or expo- ment demonstrates exposures outside the rec- guidelines, use respiratory protection. hould conform to BS EN 143
Fil	ter type	: Particulates	

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Appearance Colour Odour Odour Threshold	:	powder No data available No data available No data available
рН	:	No data available
Melting point/freezing point	:	No data available
Initial boiling point and boiling	:	No data available
range Flash point	:	Not applicable
Evaporation rate	:	Not applicable
Flammability (solid, gas)	:	May form explosive dust-air mixture during processing, han- dling or other means.
Upper explosion limit / Upper flammability limit	:	No data available
Lower explosion limit / Lower flammability limit	:	No data available
Vapour pressure	:	Not applicable
Relative vapour density	:	Not applicable
Relative density	:	No data available
Density	:	No data available
Solubility(ies) Water solubility	:	No data available



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octan	ion coefficient: n- ol/water ignition temperature	: Not applicable : No data availa			
Deco	mposition temperature	: No data availa	able		
Explo	sity scosity, kinematic sive properties zing properties	: Not explosive	: Not explosive		
Flam	information mability (liquids) cular weight cle size	 No data availa No data availa No data availa 	able		

SECTION 10: Stability and reactivity

10.1 Reactivity

Not classified as a reactivity hazard.

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

Hazardous reactions	:	May form explosive dust-air mixture during processing, han- dling or other means. Can react with strong oxidizing agents.
10.4 Conditions to avoid		
Conditions to avoid	:	Heat, flames and sparks. Avoid dust formation.
10.5 Incompatible materials		
Materials to avoid	:	Oxidizing agents

10.6 Hazardous decomposition products

No hazardous decomposition products are known.

SECTION 11: Toxicological information

11.1 Information on toxicological effects

Information on likely routes of : Inhalation



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	exposu	re		Skin contact Ingestion Eye contact	
	Acute t	oxicity			
	Not clas	ssified based on availa	ble	information.	
	Produc				
	Acute o	oral toxicity	:	Acute toxicity estin Method: Calculation	mate: > 2,000 mg/kg on method
	<u>Compo</u>	onents:			
	Lamivu				
	Acute o	oral toxicity	:	LD50 (Rat): > 2,00	00 mg/kg
				LD50 (Mouse): 4,0 Remarks: No mor	000 mg/kg tality observed at this dose.
	Acute to adminis	oxicity (other routes of stration)	:	LD50 (Rat): > 2,00 Application Route	
	Tenofovir:				
	Acute o	oral toxicity	:	LD50 (Rat): > 1,50	00 mg/kg
				LD50 (Dog): 30 m	g/kg
	Doravii	rine:			
	Acute o	oral toxicity	:	LD50 (Rat): > 750 Remarks: No mor	mg/kg tality observed at this dose.
				(Rat): Method: Pr Remarks: No evid	nototoxicity ence of phototoxicity was observed
				LD50 (Dog): > 1,0 Remarks: No mor	00 mg/kg tality observed at this dose.
				LD50 (Mouse): > 4 Remarks: No mor	450 mg/kg tality observed at this dose.
	Cellulo	se:			
	Acute o	oral toxicity	:	LD50 (Rat): > 5,00	00 mg/kg
	Acute ir	nhalation toxicity	:	LC50 (Rat): > 5.8 Exposure time: 4 Test atmosphere:	h
	Acute d	lermal toxicity	:	LD50 (Rabbit): > 2	2,000 mg/kg



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Skin d	corrosion/irritation			
Not cl	assified based on ava	ilable	information.	
Comp	oonents:			
Lamiv	/udine:			
Specie Resul		:	Rabbit Mild skin irritation	
Tenof	iovir:			
Specie Resul		:	Rabbit Mild skin irritation	
Dorav	virine:			
Rema	rks	:	No data available	
	us eye damage/eye i es serious eye irritation		ion	
<u>Comp</u>	oonents:			
Lamiv	/udine:			
Specie Resul		:	Rabbit No eye irritation	
Tenof	fovir:			
Specie Resul		:	Rabbit Severe irritation	
Dorav	virine:			
Rema	rks	:	No data available	
Respi	iratory or skin sensit	isatio	on	
Skin s	sensitisation			
Not cl	assified based on ava	ilable	information.	
•	iratory sensitisation assified based on ava	ilable	information.	
Comp	oonents:			
Lamiv	/udine:			
Expos Specio Resul		:	Dermal Guinea pig Not a skin sensitiz	zer.
Tenof		-		



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		: Skin cor : Guinea	
	avirine: narks	: No data	available
	m cell mutagenicity classified based on availa	able informati	on.
<u>Con</u>	nponents:		
Lam	ivudine:		
Gen	otoxicity in vitro		be: Bacterial reverse mutation assay (AMES) negative
			be: Mouse Lymphoma equivocal
Gen	otoxicity in vivo	Species Applicat	be: Micronucleus test : Rat ion Route: Oral negative
		mamma Species	be: Unscheduled DNA synthesis (UDS) test with lian liver cells in vivo : Rat negative
Ten	ofovir:		
Gen	otoxicity in vitro		pe: Bacterial reverse mutation assay (AMES) equivocal
		Test Tyj Result:	pe: In vitro mammalian cell gene mutation test positive
Gen	otoxicity in vivo	cytogen Species Applicat	be: Mutagenicity (in vivo mammalian bone-marrow etic test, chromosomal analysis) : Mouse ion Route: Intraperitoneal injection negative
	m cell mutagenicity- As- sment		of evidence does not support classification as a germ
Dor	avirine:		
	otoxicity in vitro	: Test Ty	be: Bacterial reverse mutation assay (AMES)



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				Result: negative		
					nosomal aberration nese hamster ovary cells	
Genotoxicity in vivo		:	Test Type: Micronucleus test Species: Rat Cell type: Bone marrow Application Route: Oral Result: negative			
С	Cellulo	se:				
G	Genoto	xicity in vitro	:	Test Type: Bacter Result: negative	ial reverse mutation assay (AMES)	
				Test Type: In vitro Result: negative	o mammalian cell gene mutation test	
G	Genotoxicity in vivo		:	Test Type: Mammalian erythrocyte micronucleus test (in vivo cytogenetic assay) Species: Mouse Application Route: Ingestion Result: negative		
		ogenicity ssified based on availa	able	information.		
<u>C</u>	Compo	nents:				
L	_amivu	dine:				
E	Species Exposu Result		: : :	Rat 2 Years negative		
E	Species Exposu Result		:	Mouse 2 Years negative		
т	Tenofo	vir:				
A E	Species Applicat Exposu Result	tion Route	::	Mouse Oral 104 weeks negative		
A E	Species Applicat Exposu Result	tion Route	: :	Rat Oral 104 weeks negative		

SAFETY DATA SHEET

According to REACH Regulation (EC) No 1907/2006, as amended by UK REACH Regulations SI 2019/758



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Spec Appli	cation Route sure time It	:	Mouse Oral 6 Months negative No significant adv	verse effects were reported	
Spec Appli Expo	Cellulose: Species Application Route Exposure time Result		Rat Ingestion 72 weeks negative		
-	oductive toxicity	unho	rn child		
•	ected of damaging the ponents:	unbo	m child.		
	vudine:				
	ts on fertility	:	Species: Rat Application Route Fertility: NOAEL:	900 mg/kg body weight on fertility and early embryonic develop-	
	Effects on foetal develop-		Species: Rabbit Application Route Symptoms: Preim Result: Embryoto spring were detec Test Type: Embry Species: Rat Application Route Developmental To	plantation loss, Skeletal malformations xic effects and adverse effects on the off- ted. ro-foetal development :: Oral oxicity: LOAEL: 45 mg/kg body weight	
			Symptoms: Effect Result: positive	s on foetal development	
Repr	oductive toxicity - As- nent	:	Some evidence o animal experimer	f adverse effects on development, based on its.	
Teno	fovir:				
	ts on fertility	:	Test Type: Fertilit Species: Rat Application Route Result: No effects		
Effec	ts on foetal develop-	:	Test Type: Embry	ro-foetal development	



rsion B	Revision Date: 28.09.2024	SDS Number: 9371580-00009	Date of last issue: 06.04.2024 Date of first issue: 27.08.2021
ment			at Route: Oral adverse effects
		Species: Ra	Embryo-foetal development abbit adverse effects
Dora	virine:		
Effect	s on fertility	Fertility: NC	Fertility at, male and female AEL: 450 mg/kg body weight effects on fertility
Effect ment	s on foetal develop-	Species: Ra Application Developme	Embryo-foetal development at Route: Oral ntal Toxicity: NOAEL: 450 mg/kg body weight adverse effects
		Species: Ra Application Developme	Embryo-foetal development abbit Route: Oral ntal Toxicity: NOAEL: 300 mg/kg body weight adverse effects
Cellu	lose:		
Effect	s on fertility	Species: Ra	Route: Ingestion
Effect ment	s on foetal develop-	Species: Ra	Route: Ingestion

Not classified based on available information.

STOT - repeated exposure

May cause damage to organs through prolonged or repeated exposure.

Components:

Lamivudine:

Exposure routes	: Ingestion
Target Organs	: Blood
Assessment	: May cause damage to organs through prolonged or repeated
	exposure.

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Tar	ofovir: get Organs essment	: Bone, Kidney : May cause dan exposure.	nage to organs through prolonged or repeated
Rep	peated dose toxicity		
<u>Co</u>	nponents:		
Spe NO App Exp Tar Syr	nivudine: ecies AEL olication Route oosure time get Organs nptoms narks		l discomfort, Breathing difficulties, Fatality city observed in testing
LÖ/ App Exp Tar	ecies AEL blication Route bosure time get Organs nptoms	: Dog : 90 mg/kg : Oral : 12 Months : Blood, spleen, : Salivation, Diar	
NO App Exp	ecies AEL dication Route posure time get Organs	: Mouse : 500 mg/kg : Oral : 1 Months : Blood	
Spe NO LO/ App Exp	ofovir: ecies AEL AEL blication Route bosure time get Organs	: Rat : 30 mg/kg : 300 mg/kg : Oral : 13 Weeks : Bone	
NO LO/ App Exp	ecies AEL AEL olication Route oosure time get Organs	: Dog : 3 mg/kg : >= 10 mg/kg : Oral : 42 Weeks : Kidney	
	ecies AEL	: Monkey : 10 mg/kg	



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Expos	cation Route sure time et Organs	: Subcutan : 10 Month : Bone	
Dora	virine:		
	EL cation Route sure time	: Rat : 450 mg/k : Oral : 6 Months : No signifi	g cant adverse effects were reported
	EL cation Route sure time	: Mouse : > 450 mg : Oral : 3 Months : No signifi	/kg cant adverse effects were reported
	EL cation Route sure time	: Dog : > 1,000 n : Oral : 9 Months : No signifi	ng/kg cant adverse effects were reported
Cellu	lose:		
		: Rat : >= 9,000 : Ingestion : 90 Days	mg/kg
-	ration toxicity lassified based on ava	ilable informatio	٦.
Expe	rience with human e	xposure	
Com	oonents:		
	vudine:		
Inges	tion	: Symptom rhoea, Co	s: Headache, Fatigue, Respiratory disorders, Diar- bugh
Teno	-		
Inges	tion	: Symptom ache, Ras	s: Nausea, Diarrhoea, Vomiting, flatulence, Head- sh
Dora	virine:		
Inges	tion		s: confusion, Headache, Dizziness, Nausea, Rash, dreams, flushing, Neurological disorders, mental n



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SECTION 12: Ecological information

12.1 Toxicity

Components:		
Lamivudine: Toxicity to fish	:	LC50 (Pimephales promelas (fathead minnow)): > 97.7 mg/l Exposure time: 96 h Method: OECD Test Guideline 203
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): > 100 mg/l Exposure time: 48 h Method: OECD Test Guideline 202
Toxicity to algae/aquatic plants	:	EC50 (Pseudokirchneriella subcapitata (green algae)): > 96.9 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
		NOEC (Pseudokirchneriella subcapitata (green algae)): 96.9 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
Tenofovir:		
Toxicity to algae/aquatic plants	:	EC50 (Raphidocelis subcapitata (freshwater green alga)): 69 mg/l End point: Growth Exposure time: 72 h Method: OECD Test Guideline 201
		NOEC (Raphidocelis subcapitata (freshwater green alga)): 18 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
Toxicity to microorganisms	:	EC50 : > 1,000 mg/l Exposure time: 3 h Test Type: Respiration inhibition Method: OECD Test Guideline 209
		NOEC : > 1,000 mg/l Exposure time: 3 h Test Type: Respiration inhibition Method: OECD Test Guideline 209
Toxicity to fish (Chronic tox- icity)	:	NOEC: 9 mg/l Exposure time: 32 d Species: Pimephales promelas (fathead minnow)



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				Method: OECD Te	est Guideline 210
		/ to daphnia and other invertebrates (Chron- ity)	:	NOEC: 12 mg/l Exposure time: 21 Species: Daphnia Method: OECD Te	magna (Water flea)
	Doravi	rine:			
		<i>t</i> to daphnia and other invertebrates	:	Exposure time: 48 Method: OECD Te	
				EC50 (Americamy Exposure time: 96	
	Toxicity plants	/ to algae/aquatic	:	mg/l Exposure time: 72 Method: OECD Te	
				mg/l Exposure time: 72 Method: OECD Te	
	Toxicity	<i>i</i> to microorganisms	:	EC50 : > 1,000 m Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
				NOEC : 1,000 mg Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
	Toxicity icity)	/ to fish (Chronic tox-	:	Method: OECD Te	ales promelas (fathead minnow)
	•	/ to daphnia and other invertebrates (Chron- ity)	:	Method: OECD Te	magna (Water flea)



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	Cellulc Toxicity	o se: y to fish	:	Exposure time: 4	tipes (Japanese medaka)): > 100 mg/l 8 h on data from similar materials			
12.2	12.2 Persistence and degradability							
	Compo	onents:						
	Lamivu	udine:						
	Biodeg	radability	:	Result: Not readil Biodegradation: Exposure time: 2	4 %			
	Tenofo	ovir:						
	Biodeg	radability	:	Result: Not readil Biodegradation: Exposure time: 2 Method: OECD T	3.66 %			
	Doravi	rine:						
	Biodeg	radability	:	Result: Not readil Biodegradation: Exposure time: 2	2 %			
	Cellulo	ose:						
	Biodeg	radability	:	Result: Readily b	iodegradable.			
12.3	Bioaco	cumulative potential						
	Compo	onents:						
	Lamivu	udine:						
	Partitio octanol	n coefficient: n- I/water	:	log Pow: -1.44				
	Tenofo							
	Partitio octanol	n coefficient: n- I/water	:	log Pow: 1.06 pH: 7				
	Doravi	rine:						
		n coefficient: n-	:	log Pow: 2.08				
12.4	Mobili	ty in soil						
	Compo	onents:						
	Lamivu	udine:						



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	bution among environ- al compartments	:	log Koc: 2.03	
Tenofovir: Distribution among environ- mental compartments		:	0	est Guideline 106
Distri	virine: bution among environ- al compartments	:	log Koc: 2.86	
12.5 Resu	llts of PBT and vPvB a	sse	ssment	
Prod	uct:			
Asse	ssment	:	to be either persis	nixture contains no components considered stent, bioaccumulative and toxic (PBT), or nd very bioaccumulative (vPvB) at levels of
12.6 Othe	r adverse effects			
Prod	uct:			
Endo tial	crine disrupting poten-	:	ered to have end	nixture does not contain components consid- ocrine disrupting properties for environment REACH Article 57(f).
	l 13: Disposal consi	dera	ations	
13.1 Wast	e treatment methods			

Product	 Dispose of in accordance with local regulations. According to the European Waste Catalogue, Waste Codes are not product specific, but application specific. Waste codes should be assigned by the user, preferably in
	discussion with the waste disposal authorities. Do not dispose of waste into sewer.
Contaminated packaging	: Empty containers should be taken to an approved waste han- dling site for recycling or disposal. If not otherwise specified: Dispose of as unused product.

SECTION 14: Transport information

14.1 UN number

ADN	:	Not regulated as a dangerous good
ADR	:	Not regulated as a dangerous good
RID	:	Not regulated as a dangerous good
IMDG	:	Not regulated as a dangerous good



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ΙΑΤΑ		: Not regulated as	a dangerous good			
14.2 UN p	roper shipping name	-				
ADN		: Not regulated as	a dangerous good			
ADR		-	a dangerous good			
RID		: Not regulated as	a dangerous good			
IMDG	ì	: Not regulated as	a dangerous good			
ΙΑΤΑ		: Not regulated as	a dangerous good			
14.3 Trans	sport hazard class(es)					
ADN		: Not regulated as	a dangerous good			
ADR		: Not regulated as	a dangerous good			
RID		: Not regulated as	a dangerous good			
IMDG	ì	: Not regulated as	a dangerous good			
ΙΑΤΑ		: Not regulated as	a dangerous good			
14.4 Pack	ing group					
ADN		: Not regulated as	a dangerous good			
ADR		: Not regulated as	a dangerous good			
RID		: Not regulated as	a dangerous good			
IMDG	ì	: Not regulated as	a dangerous good			
ΙΑΤΑ	(Cargo)	: Not regulated as	a dangerous good			
ΙΑΤΑ	(Passenger)	: Not regulated as	a dangerous good			
14.5 Environmental hazards						
Not regulated as a dangerous good						
14.6 Special precautions for user Not applicable						
	sport in bulk accordin					
Rema	arks	: Not applicable for	or product as supplied.			
SECTION 15: Regulatory information						

15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture

Relevant EU provisions transposed through retained EU law

UK REACH List of restrictions (Annex 17)	:	Not applicable
UK REACH Candidate list of substances of very high concern (SVHC) for Authorisation	:	Not applicable



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	5	lutants Regulations (ret as amended for Great		Not applicable
,	llation (EC) on substar	nces that deplete the oz	one :	Not applicable
UK R	EACH List of substan	ces subject to authorisa	ition :	Not applicable
	xport and import of hamed Consent (PIC) Re	zardous chemicals - Pr gulation	ior :	Not applicable
Contr	rol of Major Accident H	lazards Regulations 20 Not applicable	15 (COM	AH)

Other regulations:

Take note of The Management of Health and Safety at Work Regulations 1999 (requirements relating to new and expectant mothers at work contained in Regulation 16 to 18) and of the Pregnant Workers Directive 92/85/EEC.

Take note of The Management of Health and Safety at Work Regulations 1999 (requirements relating to protection of young people at work contained in Regulation 19) and of Directive 94/33/EC on the protection of young people at work.

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

AICS	:	not determined
DSL	:	not determined
IECSC	:	not determined

15.2 Chemical safety assessment

A Chemical Safety Assessment has not been carried out.

SECTION 16: Other information

Other information	:	Items where changes have been made to the previous version are highlighted in the body of this document by two vertical lines.
Full text of H-Statements		
H302	:	Harmful if swallowed.
H319	:	Causes serious eye irritation.
H361d	:	Suspected of damaging the unborn child.
H373	:	May cause damage to organs through prolonged or repeated exposure if swallowed.
Full text of other abbreviatior	าร	
Acute Tox.	:	Acute toxicity
Eye Irrit.	:	Eye irritation
Repr.	:	Reproductive toxicity
STOT RE	:	Specific target organ toxicity - repeated exposure
GB EH40	:	UK. EH40 WEL - Workplace Exposure Limits
GB EH40 / TWA	:	Long-term exposure limit (8-hour TWA reference period)



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GB EH40 / STEL : Short-term exposure limit (15-minute reference period)

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways; ADR - Agreement concerning the International Carriage of Dangerous Goods by Road; AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight; CLP - Classification Labelling Packaging Regulation; Regulation (EC) No 1272/2008; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECHA -European Chemicals Agency; EC-Number - European Community number; 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; 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; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; 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; RID - Regulations concerning the International Carriage of Dangerous Goods by Rail; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; SVHC - Substance of very high concern; 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

Further information

Sources of key data used to	:	Internal technical data, data from raw material SDSs, OECD
compile the Safety Data		eChem Portal search results and European Chemicals Agen-
Sheet		cy, http://echa.europa.eu/

Classification	of the n	nixture:

Eye Irrit. 2	H319
Repr. 2	H361d
STOT RE 2	H373

Classification procedure: Calculation method Calculation method Calculation method

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.

GB / EN