Commission Delegated Regulation (EU) 2020/1182 of 19 May 2020 amending, for the purposes of its adaptation to technical and scientific progress, Part 3 of Annex VI to Regulation (EC) No 1272/2008 of the European Parliament and of the Council on classification, labelling and packaging of substances and mixtures (Text with EEA relevance)

COMMISSION DELEGATED REGULATION (EU) 2020/1182

of 19 May 2020

amending, for the purposes of its adaptation to technical and scientific progress, Part 3 of Annex VI to Regulation (EC) No 1272/2008 of the European Parliament and of the Council on classification, labelling and packaging of substances and mixtures

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006⁽¹⁾, and in particular Article 37(5) thereof,

Whereas:

- (1) Table 3 of Part 3 of Annex VI to Regulation (EC) No 1272/2008 contains the list of harmonised classification and labelling of hazardous substances based on the criteria set out in Parts 2 to 5 of Annex I to that Regulation.
- (2) Proposals to introduce harmonised classification and labelling of certain substances and to update or delete the harmonised classification and labelling of certain other substances have been submitted to the European Chemicals Agency ('Agency') pursuant to Article 37 of Regulation (EC) No 1272/2008. Based on the opinions⁽²⁾ on those proposals issued by the Committee for Risk Assessment of the Agency (RAC), as well as on the comments received from the parties concerned, it is appropriate to introduce, update or delete the harmonised classification and labelling of certain substances. Those RAC opinions are:
 - Opinion of 8 June 2018 concerning nitric acid ... $\%[C \le 70 \%]$;
 - Opinion of 9 March 2018 concerning silicon carbide fibres (with diameter < $3 \mu m$, length > $5 \mu m$ and aspect ratio $\geq 3:1$);
 - Opinion of 8 June 2018 concerning trimethoxyvinylsilane; trimethoxy(vinyl)silane;
 - Opinion of 8 June 2018 concerning tris(2-methoxyethoxy)vinylsilane; 6-(2-methoxyethoxy)-6-vinyl-2,5,7,10-tetraoxa-6-silaundecane;
 - Opinion of 8 June 2018 concerning dimethyl disulphide;

- Opinion of 8 June 2018 concerning granulated copper;
- Opinion of 30 November 2018 concerning bis(N-hydroxy-N-nitrosocyclohexylaminato-O,O')copper; bis(N-cyclohexyl-diazenium-dioxy)-copper; [Cu-HDO];
- Opinion of 14 September 2018 concerning dioctyltin dilaurate; [1] stannane, dioctyl-, bis(coco acyloxy) derivs. [2];
- Opinion of 30 November 2018 concerning dibenzo[def,p]chrysene; dibenzo[a,l]pyrene;
- Opinion of 9 March 2018 concerning ipconazole (ISO); (1RS,2SR,5RS;1RS,2SR,5SR)-2-(4-chlorobenzyl)-5-isopropyl-1-(1H-1,2,4-triazol-1-ylmethyl)cyclopentanol;
- Opinion of 8 June 2018 concerning bis(2-(2-methoxyethoxy)ethyl)ether; tetraglyme;
- Opinion of 8 June 2018 concerning paclobutrazol (ISO); (2RS,3RS)-1-(4-chlorophenyl)-4,4-dimethyl-2-(1H-1,2,4-triazol-1-yl)pentan-3-ol;
- Opinion of 8 June 2018 concerning 2,2-bis(bromomethyl)propane-1,3-diol;
- Opinion of 14 September 2018 concerning geraniol; (2*E*)-3,7-dimethylocta-2,6-dien-1-ol;
- Opinion of 28 January 2019 concerning 2-(4-tert-butylbenzyl)propionaldehyde;
- Opinion of 9 March 2018 concerning MCPA-thioethyl (ISO);
 S-ethyl (4-chloro-2-methylphenoxy)ethanethioate;
 S-ethyl 4-chloro-o-tolyloxythioacetate;
- Opinion of 9 March 2018 concerning diisooctyl phthalate;
- Opinion of 14 September 2018 concerning 4-{[(6-chloropyridin-3-yl)methyl]
 (2,2-difluoroethyl) amino} furan-2(5H)-one; flupyradifurone;
- Opinion of 30 November 2018 concerning thiencarbazone-methyl (ISO); methyl 4- [(4,5-dihydro-3-methoxy-4-methyl-5-oxo-1*H*-1,2,4-triazol-1-yl)carbonylsulfamoyl]-5-methylthiophene-3-carboxylate;
- Opinion of 9 March 2018 concerning L-(+)-lactic acid; (2S)-2hydroxypropanoic acid;
- Opinion of 9 March 2018 concerning 2-methoxyethyl acrylate;
- Opinion of 8 June 2018 concerning glyoxylic acid ...%;
- Opinion of 14 September 2018 concerning sodium N-(hydroxymethyl)glycinate; [formaldehyde released from sodium N-(hydroxymethyl)glycinate];
- Opinion of 30 November 2018 concerning potassium (oxido-NNO-azoxy)cyclohexane; cyclohexylhydroxydiazene 1-oxide, potassium salt; [K-HDO];
- Opinion of 14 September 2018 concerning mecetronium etilsulfate; *N*-ethyl-*N*,*N*-dimethylhexadecan-1-aminium ethyl sulfate; mecetronium ethyl sulphate [MES];

- Opinion of 9 March 2018 concerning (2RS)-2-[4-(4-chlorophenoxy)-2-(trifluoromethyl)phenyl]-1-(1H-1,2,4-triazol-1-yl)propan-2-ol; mefentrifluconazole;
- Opinion of 30 November 2018 concerning oxathiapiprolin (ISO); 1-(4-{4-[5-(2,6-difluorophenyl)-4,5-dihydro-1,2-oxazol-3-yl]-1,3-thiazol-2-yl}piperidin-1-yl)-2-[5-methyl-3-(trifluoromethyl)-1*H*-pyrazol-1-yl]ethanone;
- Opinion of 14 September 2018 concerning pyrithione zinc; (*T*-4)-bis[1-(hydroxy-.kappa.*O*) pyridine-2(1*H*)-thionato-.kappa.*S*]zinc;
- Opinion of 30 November 2018 concerning 3-chloro-4-(chloromethyl)-1-[3-trifluoromethyl)phenyl]pyrrolidin-2-one; flurochloridone (ISO);
- Opinion of 30 November 2018 concerning 4,5-dichloro-2-octyl-2*H*-isothiazol-3-one; [DCOIT];
- Opinion of 8 June 2018 concerning 2-methyl-1,2-benzothiazol-3(2*H*)-one; [MBIT];
- Opinion of 30 November 2018 concerning 3-(difluoromethyl)-1-methyl-*N*-(3',4',5'-trifluorobiphenyl-2-yl)pyrazole-4-carboxamide; fluxapyroxad;
- Opinion of 8 June 2018 concerning *N*-(hydroxymethyl)acrylamide; methylolacrylamide; [NMA];
- Opinion of 15 October 2018 concerning 5-fluoro-1,3-dimethyl-*N*-[2-(4-methylpentan-2-yl)phenyl]-1*H*-pyrazole-4-carboxamide; 2'-[(*RS*)-1,3-dimethylbutyl]-5-fluoro-1,3-dimethylpyrazole-4-carboxanilide; penflufen;
- Opinion of 30 November 2018 concerning iprovalicarb(ISO); isopropyl [(2S)-3-methyl-1-{[1-(4-methylphenyl)ethyl]amino}-1-oxobutan-2-yl]carbamate;
- Opinion of 30 November 2018 concerning silthiofam (ISO); *N*-allyl-4,5-dimethyl-2-(trimethylsilyl)thiophene-3-carboxamide;
- Opinion of 9 March 2018 concerning Margosa, ext. [cold-pressed oil of Azadirachta indica seeds without shells extracted with super-critical carbon dioxide];
- Opinion of 8 June 2018 concerning nitric acid ...%[C> 70 %];
- Opinion of 9 March 2018 concerning octamethylcyclotetrasiloxane; [D4];
- Opinion of 30 November 2018 concerning pirimiphos-methyl (ISO); *O*-[2-(diethylamino)-6-methylpyrimidin-4-yl] *O,O*-dimethyl phosphorothioate;
- Opinion of 30 November 2018 concerning phosphine;
- Opinion of 14 September 2018 concerning dichlorodioctylstannane;
- Opinion of 30 November 2018 concerning 2-ethylhexyl 10-ethyl-4,4-dioctyl-7-oxo-8-oxa-3,5-dithia-4-stannatetradecanoate; [DOTE];
- Opinion of 30 November 2018 concerning lead;
- Opinion of 14 September 2018 concerning 2-butoxyethanol; ethylene glycol monobutyl ether;

- Opinion of 30 November 2018 concerning *m*-bis(2,3-epoxypropoxy)benzene; resorcinol diglycidyl ether;
- Opinion of 14 September 2018 concerning tribenuron-methyl (ISO); methyl 2- [N-(4-methoxy-6-methyl-1,3,5-triazin-2-yl)-N-methylcarbamoylsulfamoyl]benzoate;
- Opinion of 8 June 2018 concerning azoxystrobin (ISO); methyl (E)-2-{2-[6-(2-cyanophenoxy)pyrimidin-4-yloxy]phenyl}-3-methoxyacrylate;
- Opinion of 9 March 2018 concerning ethofumesate (ISO); (RS)-2-ethoxy-2,3-dihydro-3,3-dimethylbenzofuran-5-yl methanesulfonate;
- Opinion of 30 November 2018 concerning 2,4-dinitrophenol;
- Opinion of 14 September 2018 concerning mesotrione (ISO); 2-[4-(methylsulfonyl)-2-nitrobenzoyl]-1,3-cyclohexanedione;
- Opinion of 30 November 2018 concerning octhilinone (ISO); 2-octyl-2*H*-isothiazol-3-one; [OIT];
- Opinion of 14 September 2018 concerning hymexazol (ISO); 3-hydroxy-5methylisoxazole;
- Opinion of 30 November 2018 concerning hexythiazox (ISO); *trans*-5-(4-chlorophenyl)-*N*-cyclohexyl-4-methyl-2-oxo-3-thiazolidine-carboxamide;
- Opinion of 9 March 2018 concerning pymetrozine (ISO); (*E*)-4,5-dihydro-6-methyl-4-(3-pyridylmethylene amino)-1,2,4-triazin-3(2*H*)-one;
- Opinion of 9 March 2018 concerning imiprothrin (ISO); reaction mass of: [2,4-dioxo-(2-propyn-1-yl)imidazolidin-3-yl]methyl(1*R*)-*cis*-chrysanthemate; [2,4-dioxo-(2-propyn-1-yl)imidazolidin-3-yl]methyl(1*R*)-*trans*-chrysanthemate;
- Opinion of 14 September 2018 concerning butanone oxime; ethyl methyl ketoxime; ethyl methyl ketone oxime;
- Opinion of 8 June 2018 concerning bis(α , α -dimethylbenzyl) peroxide;
- Opinion of 9 March 2018 concerning branched hexatriacontane;
- Opinion of 30 November 2018 concerning hexyl 2-(1-(diethylaminohydroxyphenyl) methanoyl)benzoate; hexyl 2-[4-(diethylamino)-2-hydroxybenzoyl]benzoate.
- (3) With regard to the substance lead (CAS number 7439-92-1 and index numbers 082-013-00-1 (lead powder; [particle diameter < 1 mm];) and 082-014-00-7 (lead massive; [particle diameter ≥ 1 mm];)), RAC proposed in its opinion of 30 November 2018 to apply the same environmental classification to the massive and the powder form. However, in view of the lower dissolution rate of the massive form, the malleable structure of lead, the specific intentional production of the powder and the different environmental classification between massive and powder forms for existing entries in Annex VI for other metals, further assessment needs to be done by RAC on whether to apply the same environmental classification to the massive as to the powder form of lead. In addition, new scientific data has been made available suggesting that the environmental classification for the massive form as recommended in the RAC opinion might not be appropriate Therefore, the environmental classification for the massive

- form will not be included in Annex VI to Regulation (EC) No 1272/2008 until RAC has had the opportunity to deliver a revised opinion.
- (4) With regard to the substance 2-butoxyethanol; ethylene glycol monobutyl ether; (CAS number 111-76-2), new scientific data has been made available for the hazard class 'acute toxicity (inhalation)' which suggests that the classification for this hazard class as recommended in the RAC opinion, which is based on older data, might not be appropriate. Therefore, this hazard class should not be modified in Annex VI to Regulation (EC) No 1272/2008 until RAC has had the opportunity to deliver a revised opinion based on the new information, while all other hazard classes covered by the RAC opinion should be included.
- (5) Regulation (EC) No 1272/2008 should therefore be amended accordingly.
- (6) Compliance with the new or updated harmonised classifications should not be required immediately as a certain period of time is necessary to allow suppliers to adapt the labelling and packaging of substances and mixtures to the new or revised classifications and to sell existing stocks subject to the pre-existing regulatory requirements. That period of time is also necessary to allow suppliers sufficient time to take the actions required to ensure continuing compliance with other legal requirements following the changes made under this Regulation. Such requirements may include those set out in point (f) of Article 22(1) of Regulation (EC) No 1907/2006 of the European Parliament and of the Council⁽³⁾ or those set out in Article 50 of Regulation (EU) No 528/2012 of the European Parliament and of the Council⁽⁴⁾.
- (7) Suppliers should, however, have the possibility to apply the new classification, labelling and packaging provisions on a voluntary basis before the date of application of this Regulation. This is consistent with the approach taken under Article 61(2) of Regulation (EC) No 1272/2008,

HAS ADOPTED THIS REGULATION:

Article 1 U.K.

Amendments to Regulation (EC) No 1272/2008

Table 3 of Part 3 of Annex VI to Regulation (EC) No 1272/2008 is amended as set out in the Annex to this Regulation.

Article 2 U.K.

Entry into force and application

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

It shall apply from 1 March 2022.

By way of derogation from the second paragraph of this Article, substances and mixtures may, before 1 March 2022 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by this Regulation.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 19 May 2020.

For the Commission

The President

Ursula VON DER LEYEN

ANNEX U.K.

In Annex VI to Regulation (EC) No 1272/2008, Table 3 of Part 3 is amended as follows:

(1) the following entries are inserted:

Index	Chem	ChemicEC CAS			fication		Specif	idNotes		
No	name	No	No	Hazar	d Hazar	d Pictog	rahhazar	d Suppl.	Conc.	Limits,
				Class	statem	e Si gnal	statem	e h tazar	վ M-	
				and				s)statem		5
				Catego		Code(s)	Code(s)and	
				Code(ATE	
'007-0	3 60i+60 0c-3	231-71	47 -6 97-3		H272	GHS03		EUH07		B'
	acid			Liq. 3		GHS06	l		Liq.	
	% [C≤			Acute Tox.	H314	GHS05) Н314		3; H272:	
	[C ≤ 70 %]			3		Dgr			11272. C≥	
	70 70]			Skin					65 %	
				Corr.					inhalat	ion:
				1A					ATE	
									=	
									2,65	
									mg/L (vapou	re)
									Skin	13)
									Corr.	
									1A;	
									H314:	
									$C \ge$	
									20 % Skin	
									Corr.	
									1B;	
									H314:	
									5 %	
									$\leq C <$	
									20 %	
'014 - 0	48iH000+6				B H350i		H350i'			
	carbide	}	308076	5-74-6		Dgr				
	fibres									
	(with diamet	er								
	< 3	C1								
	μm,									
	length									
	> 5									
	μm									
	and									
	aspect ratio									
	≥ 3:1)									
	/									

014-0	4 9 i 60 e (0				H317		H317'			
	trimeth	oxy(vin	yl)silan	esens. 1B		Wng				
014-0	6-(2- methox	yethoxy yethoxy ,5,7,10- a-6-	y)vinyls y)-6-		H360F	IGHS08 Dgr	3H360F	D'		
016-0	9 8i-000+13 disulph		16:204-92	Liq. 2 Acute Tox. 3 Acute Tox. 3 STOT SE 3	c		H331 H301		inhalat ATE = 5 mg/L (vapou oral: ATE = 190 mg/ kg bw M = 1 M = 10'	
*029-0 X	copper [particlength: from 0,9 mm to 6,0 mm; particle width: from 0,494 to 0,949 mm]	e	97 -6 40-5	0A&quati Chroni 2	сH411 с	GHS09)H411'			

				Г		1	<u> </u>	1		
'029-02	25is00V-5 23	9-70	33-412600) 18 19a+18.	H228	GHS02	2H228		oral:	
	hydroxy- <i>I</i>	V-	15627-	039 o.5 1	H302	GHS07	'H302		ATE	
	nitrosocyo	clohe	xylamir	atoute	DH373p	eGHS08	3H373(li	ver)	= 360	
	bis(N-			Tox.		GHS05			mg/	
	cyclohexy	/1_		4	H318	GHS09			kg bw	
	diazenium			STOT		Dgr	11.10		M = 1	
	dioxy)-	1-		RE 2	H410	Dgi			M = 1	
	copper;			Eye	11710				1'	
	[Cu-			Dam.					1	
	HDO]			1	_					
				Aquati	C					
				Acute						
				1						
				Aquati						
				Chroni	c					
				1						
<u>'050_0'</u>	3 di-00t-9 1t <u>0</u> 2	2-88	33,6 4,Ω_1	SQ &nr	H360D	GHSUS	H360D			
050-0.	dilauratel		[1]	а че рг. 1В	H372	Dgr	H372			
			19 5 648-		(immu	_	(immun			
				RE 1						
	stannan [2]	J	[2]	KE I	system	,	system)			
	dioctyl-,									
	bis(coco									
	acyloxy)									
	derivs.									
	[2]									
601-0°	9 2 iH 0:e n#0 2 #	1 <i>5£</i> 88	6684886	enCarc 1	IB:1350	GHS08	H350		Carc. 1	B·
001 0	dibenzo[a			Muta.	H341	Dgr	H341		H350:	Σ,
		, 1PJ		2	115 11	251	113 11		C ≥	
				_					0,001	
									%,	
									70	
'603-2i	3 ip@ona zol	e	125225				3H360D		oral:	
	(ISO);		115850		H302	GHS07	'H302		ATE	
	(1RS,2SR,	,5 <i>RS</i> ;	11 R.S , 23 37	R ASSESCHRO -	2H373	GHS09	H373		= 500	
	(4-			Tox.	(eyes,	Dgr	(eyes,		mg/	
	chloroben	zyl)-	5-	4	skin,		skin,		kg bw	
	isopropyl-			STOT	liver)		liver)		$\widetilde{M} =$	
	(1H-1,2,4)			RE 2	H410		H410		100'	
	triazol-1-			Aquati						
	ylmethyl)	cyclo	pentand							
				1						
(605.5	310: 0/2 0 2 3		41 800 0 0		110 60-	DOLLC :	112 627			
603-2	3 8но(2- 9 20	15-59	41-473-24		H360F		H360FI)´		
	(2-		اید	1B		Dgr				
	methoxye	- 4	y)ethyl)	ether;						
	tetraglyme	e								
·602 2	3 9 a0loHutra	2701	76738-	ADeAr	Н3614	GHS08	Н3614		inhalat	ion.
003-2.	(ISO);	aZUI	10130-	2	H332	GHS07			ATE	1011.
		\ ₁				GHS09			AIE =	
	(2RS,3RS)	<i>)</i> -1-		Acute Tox.	H302 H319	Wng	H302 H319			
				1/17	11119	vv no l	11119 L		3,13	
	(4-	1\	4.4			"""5				
	chlorophe dimethyl-		4,4-	4	H400 H410	,,,,,,	H410		mg/L (dusts	

	(1 <i>H</i> -1,2, triazol-1 yl)pentar	-		Acute Tox. 4 Eye Irrit. 2 Aquati Acute 1 Aquati Chroni 1	c				or mists) oral: ATE = 490 mg/ kg bw M = 10 M = 10'	
'603-24 X	4 0,-2 00- 2 bis(brom diol		7 3-2 796-9 yl)prop			GHS08 Dgr	3H350 H340'			
·603-24	dimethyl dien-1-	-		- \$ kin Sens. 1	Н317	GHS07 Wng	'H317'			
['] 605-0	4 2- 00-3 2 (4- <i>tert</i> - butylben			1B	H360F	dGHS08 Dgr	3H360F	ď'		
`607-7	thioethyl (ISO); S- ethyl (4- chloro-2 methylpl S- ethyl 4- chloro-o tolyloxyt	- henoxy	v)ethane	Tox. 4 STOT RE. 2 Aquati Acute	H410 c	GHS08 GHS09 Wng	3H373		oral: ATE = 450 mg/ kg bw M = 10 M = 10'	
	1 0 ii3002t3 phthalate		23 554-	Жер т. 1В	H360F	IGHS08 Dgr	3H360F	D'		
'607-7 <i>i</i>	44-00-4 {[(6- chloropy yl)methy (2,2- difluoroe one; flupyrad	/l] ethyl)a	mino}f	Tox. 4 STOT RE 2	H400 H410 cH)-	GHS08 GHS08 GHS09 Wng	3H373	e)	oral: ATE = 500 mg/ kg bw M = 10 M = 10'	

	10.00	21701	. 02 14	11400	CHICO	11110		1.6	
	4 2 hi 20 hearbazone	-31/813			GHS09	/H410		M = 1000	
X	methyl		Acute	H410	Wng			1000	
	(ISO);		1					M =	
	methyl		Aquati					1000'	
	4-		Chroni	c					
	[(4,5-		1						
	dihydro-3-								
	methoxy-4-								
	methyl-5-								
	oxo-1 <i>H</i> -1,2,4-								
	triazol-1-		_						
	yl)carbonylsul		-5-						
	methylthiophe	ne-3-							
	carboxylate								
·607-7	4B-00-5 201-19	<i>6</i> 7 -2 -33-4	4Skin	H314	GHS05	H314	EUH07	71'	
,	(+)-		Corr.	H318	Dgr			-	
	lactic		1C	10	- 0-				
	acid;		Eye						
	(2S)-2-		Dam.						
	hydroxypropar	noic	1						
	acid								
(607.7		m 201	4D#	11226	GHGOG	11006	FILLO	77 1 1 .	•
607-7	42- 00-0 221-49	93-1321-6		H226	GHS02		EUH0	inhalat	ion:
	methoxyethyl		Liq. 3		GHS05		ь.	ATE	
	acrylate		Muta.			H360F	D	=2,7	
			2	H331	GHS08			mg/L	
			Repr.	H302	Dgr	H302		(vapou	rs)
			1B	H314		H314		oral:	
			Acute			H317		ATE	
			Tox.	H317				= 404	
			3					mg/	
			Acute					kg	
			Tox.					bw'	
			4						
			Skin						
			Corr.						
			1C						
			Eye						
			Dam.						
			1						
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			Sens.						
			1						
607-7	4 51-00 x61206-05	829 8-12	- H ye	H318	GHS05	H318			B'
	acid		Dam.	H317	GHS07				
	%		1		Dgr				
			Skin		-				
			Sens.						
			1B						

								1		1
'607 - 74	lsediluih	274-35	<i>77-</i> 8 161-			GHS08			inhalat	i&n:
	<i>N</i> -			Muta.	H341	GHS07			ATE	9'
			yl)glycii		H332	Dgr	H332		= 3	
		dehyde		Acute			H302		mg/L	
	release	d		Tox.	H335		H335		(dusts	
	from			4	H315		H315		or	
	sodium	-		Acute	H319		H319		mists)	
	<i>N</i> -			Tox.	H317		H317		oral:	
	(hydro)	cymethy	yl)glycii	n 4 te]					ATE	
				STOT					=	
				SE 3					1100	
				Skin					mg/	
				Irrit.					kg bw	
				2						
				Eye						
				Irrit.						
				2						
				Skin						
				Sens.						
				1						
·611 19	apedosti	um	66603-	III am	H228	GHS02	H228		oral:	
011-10	oxido-		00003-	Sol. 1	H301	GHS06			ATE	
	azoxy)		vana.	Acute	H373	GHS08			= 136	
			roxydia		(liver)		(liver)		mg/	
	1-	ZXYIIIYU	ioxydia	3	H315	GHS09			kg	
	oxide,			STOT		Dgr	H318		bw'	
	potassi	um		RE 2	H411	Dgi	H411		UW	
	salt;	GIII		Skin	11111		11111			
	[K-			Irrit.						
	HDO]			2						
				Eye						
				Dam.						
				1						
				Aquati	c					
				Chroni						
				2						
(612.5	14.00.5	221.15	0.000		11011	CITCO		DITTO	73.6	
612-29	I		6-6 06-1		H314	GHS05		EUH07		
	etilsulf	ate;		Corr.	H318	GHS09	H410		100	
	N-	7.3.7		1	H400	Dgr			M =	
	ethyl-N			Eye	H410				1000'	
		,	ecan-1-	Dam.						
	aminiu	m		1						
	ethyl			Aquati	c					
	sulfate			Acute						
	mecetro	onium		1						
	ethyl			Aquati						
	sulphat	e;		Chroni	c					
	[MES]			1						
⁶¹³⁻³²	3(12 RO)62	2-	141778	25H033 -6	H317	GHS07	H317		M = 1	
	[4-(4-			Sens.	H400	GHS09			M =	
	chlorop	henoxy)-2-	1	H410	Wng			1'	
1	~	-		•				. '		

	(1 <i>H</i> -1,2 triazol- yl)prop ol;	2,4- 1-	yl)pheny azole	/ I duati Acute 1 Aquati Chroni 1	c					
`613-3 .	dihydro oxazol- yl]-1,3- thiazol- yl}pipe yl)-2- [5- methyl	ophenylo-1,232- eridin-1- romethy)-4,5-	&G์นิสติ Chroni 1		GHS09 Wng)H410		M = 1'	
' 613-3:	3 βy0Ωh7 0 zinc;	o 238 6-67	11- 3 463-	4Repr. 1B	H360D H330	GHS08 GHS06	H360D)	inhalati ATE	ion:
(612.2)	(T-4)- bis[1- (hydror thionat	okapp	pa.O)py a.S]zinc	Acute Tox. ridine-2 Acute Tox. 3 STOT RE 1 Eye Dam. 1 Aquati Acute 1 Aquati Chroni	H301 H372 (HB)8 H400 H410	GHS09 Dgr	3H301 9H372 H318 H410	D	= 0,14 mg/L (dusts or mists) oral: ATE = 221 mg/ kg bw M = 1000 M = 10'	
·613-3:	[3-	4- methyl)		1B Acute Tox. 4 Skin	H302 H317 H400 H410	IIGHS08 GHS07 GHS09 Dgr		D	oral: ATE = 500 mg/ kg bw M = 100 M = 100'	

'613-33 4 , 5 00-8 264-84 dichloro-2-octyl-2 <i>H</i> -isothiazol-3-one; [DCOIT]	Aquati Acute 1 Aquati Chroni 1 36-\$359-84-e5te Tox. 2 Acute Tox. 4 Skin Corr. 1 Eye Dam. 1 Skin Sens. 1A Aquati Acute 1 Aquati Chroni 1	C C C H330 H302 H314 H318 H317 H400 H410	GHS06 GHS09 Dgr	H302	EUH07	7inhalat ATE = 0,16 mg/L (dusts or mists) oral: ATE = 567 mg/ kg bw Skin Irrit. 2; H315: 0,025 % ≤ C < 5 % Eye Irrit. 2; H319: 0,025 % ≤ C < 3 % Skin Sens. 1A; H317: C ≥ 0,0015 % M = 100 M =	ion:
						M = 100'	
'613-33 2 -00-3 methyl-1,2- benzothiazol-3 one; [MBIT]	2527-66A&ute Tox. (2 <i>H</i>)- 4 Acute Tox. 3	H312 H301 H314 H318 H317 H400	GHS06 GHS05 GHS09 Dgr	H301	EUH07	ATE = 1100 mg/ kg bw	:

			Skin Corr. 1C Eye Dam. 1 Skin Sens. 1A Aquati Acute 1 Aquati Chroni 2	c				oral: ATE = 175 mg/ kg bw Skin Sens. 1A; H317: C ≥ 0,0015 % M = 1'	
	28-00-4 (difluoromethy methyl- <i>N</i> - (3',4',5'- trifluorobipher yl)pyrazole-4- carboxamide; fluxapyroxad	iyl-2-	Aquati Acute 1 Aquati Chroni 1	H410 c c	GHS09 Wng)H362 H410		M = 1 M = 1'	
'616-2.	My-00-5 213-10 (hydroxymethymethylolacryla [NMA]	/l)acryla		BH350 H340 H372 (periph nervou system	S	H350 H340 H372 (periph nervous	S		
`616-2.	fluoro-1,3- dimethyl-N- [2-(4- methylpentan-yl)phenyl]-1H- pyrazole-4- carboxamide; 2'- [(RS)-1,3- dimethylbutyl] fluoro-1,3- dimethylpyraz- carboxanilide; penflufen	2- - -5-	Aquati Acute 1 Aquati Aquati Chroni 1	cH400 H410	GHS08 GHS09 Wng			M = 1 M = 1'	
·616-2.	(ISO); isopropyl [(2S)-3- methyl-1- {[1- (4-	140923	3 . (1∄F €. 2	H351	GHS08 Wng	3Н351'			

	methylphenyl) oxobutan-2- yl]carbamate	ethyl]ar	nino}-1	_				
·616-2.	38il00offam (ISO); N- allyl-4,5- dimethyl-2- (trimethylsilyl carboxamide		RE 2 Aquati Chroni 2 ne-3-	H411 c	GHS08 GHS09 Wng	3H373 9H411'		
`650-0 .	ext. [cold-pressed oil of Azadirachta indica seeds without shells extracted with super-critical carbon dioxide]	48-7 1696-	2 Ֆ գնյati Chroni 3			H412'		

(2) the entries corresponding to index numbers 007-004-00-1; 014-018-00-1; 015-134-00-5; 015-181-00-1; 050-021-00-4; 050-027-00-7; 082-013-00-1; 605-019-00-3; 603-014-00-0; 603-065-00-9; 607-177-00-9; 607-256-00-8; 607-314-00-2; 609-041-00-4; 609-064-00-X; 613-112-00-5; 613-115-00-1; 613-125-00-6; 613-202-00-4; 613-259-00-5; 616-014-00-0 and 617-006-00-X are replaced by the following entries respectively:

Index	Chem	ic E IC	CAS	Classi	fication	Labell	ling		Specif	idNotes
No	name	No	No	Hazar	d Hazar	d Pictog	ra Ha zar	d Suppl	Conc.	Limits,
				Class	statem	e S tignal	staten	ne h tazar	d M-	
				and	Code(s)Word	Code(s)statem	efactor	S
				Catego	ory	Code(s)	Code(
				Code(s)				ATE	
'007-0	Ora i 100 Oc- 1	231-71	476 97-3	79 <u>3</u> x.	H272	GHS03	H272	EUH0'	7Ox.	B'
	acid			Liq. 2	H330	GHS06	H330		Liq.	
	%			Acute	H314	GHS05	H314		2;	
	[C >			Tox.		Dgr			H272:	
	70 %]			1					$C \ge$	
				Skin					99 %	
				Corr.					Ox.	
				1A					Liq.	
									3;	
									H272:	
									70 %	

						≤ C < 99 %	
·014-0	1 8eUUnet209e y3 16t76 r66/ [D4]	Repne; 2 Aquati Chroni 1	*** cH410	GHS08 GHS09 Wng		M = 10'	
	methyl (ISO); O-[2- (diethylamino)-6- methyl pyrimidin-4- yl] O,O- dimethyl phosphorothioate	Tox. 4 STOT RE 1 Aquati Acute 1 Aquati Chroni 1	H372 (nervot system H400 cH410 c c c H220 H330 H314 H400		3H372 (nervousystem) H410 2H220 H330 5H314 5H400	oral: ATE = 1414 mg/ kg bw M = 1000 M = 1000' inhalat ATE = 10 ppmV (gases)	idn':
°050-02	2di-Clot-01120e5\$ St242e3	Repr. 1B Acute Tox. 2 STOT RE 1 Aquati Chroni 3	H330 H372 ** H412	GHS06 GHS06 Dgr	3H360D 5H330 H372 ** H412	Repr. 1B; H360 D: C≥ 0,03 % inhalat ATE = 0,098 mg/L (dusts or mists)'	ion:
'050-02	2 7- 00-7 239-62 21-3 571- ethylhexyl 10- ethyl-4,4-	5%eþ r. 1B STOT RE 1	H360D H372 (immur system	GHS09 nDgr	H360D H372 (immur system)		

·082-0	dioctyl-7- oxo-8- oxa-3,5- dithia-4- stannatetradeca [DOTE] Beato-1 231-10 powder; [particle diameter < 1 mm]		Aquati-Acute 1 Aquati-Chroni 1 Repr. 1A Lact. Aquati-Acute 1 Aquati-Chroni 1	H410 c c H360F H362 H400 cH410	IØHS08 GHS09 Dgr		D	Repr. 1A; H360D C ≥ 0,03 % M = 1 M = 10'):
°603-0	butoxyethanol; ethylene glycol monobutyl ether		Tox. 4* Acute Tox. 4 Skin Irrit. 2 Eye Irrit. 2	H332 H302 H315 H319	GHS07 Wng	7H332 H302 H315 H319		oral: ATE = 1200 mg/ kg bw'	
	bis(2,3- epoxypropoxy) resorcinol diglycidyl ether)benzen	Muta. Acute Tox. Acute Tox. 4 Skin Irrit. 2 Eye Irrit. 2 Skin Sens. 1 Aquati Chroni 3	H341 H311 H302 H315 H319 H317 H412	GHS08 GHS06 Dgr	6H341 H311 H302 H315 H319 H317 H412		dermal ATE = 300 mg/ kg bw oral: ATE = 500 mg/ kg bw'	
['] 607-1'	7 G-iD0n9 r 401 -19 methyl (ISO);	01-0 1200	STROOT RE 2	H373 H317 H400	GHS08 GHS07 GHS09	'H317		M = 100	

	methyl 2-[N-(4-methoxy-6-methyl-1,3,5-triazin-2-yl)-N-methyl carbam	oylsulfa	Skin Sens. 1 Aquati Acute 1 Aquati 100yljbi	c	Wng		M = 100'	
·607-2	faz00y8trobin (ISO); methyl (E)-2- {2- [6-(2- cyanophenoxy yloxy]phenyl) methoxyacryl	y)pyrimi -3-	Aduse Tox. 3 Aquati Acute 1 diaquati Chroni	H400 H410 c	GHS09 GHS09 Dgr		inhalat ATE = 0,7 mg/L (dusts or mists) M = 10 M = 10'	ion:
·607-3	ethoxy-2,3-dihydro-3,3-dimethylbenzyl	ofuran-5	Acute 1 Aquati Chroni	H410 c	GHS09 Wng)H410	M = 1 M = 1'	
°609-04	42,40-4 200-03 dinitrophenol	3 75-1 7-28-	5Acute Tox. 3 * Acute Tox. 3 Acute Tox. 2 STOT RE 1 Aquati Acute 1	H311 H300 H372 H400	GHS06 GHS09 GHS09 Dgr	H311	dermal ATE = 300 mg/ kg bw oral: ATE = 30 mg/ kg bw'	
'609-00 X	6th@trione (ISO); 2-[4- (methylsulfon nitrobenzoyl]- cyclohexaned	1,3-	2 STOT RE 2	H361d H373 (eyes, nervou csystem H400 H410	GHS09 Wng s	H361d H373 (eyes, nervous system) H410	M = 10 M = 10'	

·613-1	laedoHsi (ISO); 2- octyl-2 isothia one; [OIT]		12-6530-	Aquati-Chroni 1 20elite Tox. 2 Acute Tox. 3 Acute Tox. 3 Skin Corr. 1 Eye Dam. 1 Skin Sens. 1A Aquati-Acute 1 Aquati-Chroni 1	H330 H311 H301 H314 H318 H317 H400 H410	GHS06 GHS09 Dgr	H311	EUH07	7 Inhalat ATE = 0,27 mg/L (dusts or mists) dermal ATE = 311 mg/ kg bw oral: ATE = 125 mg/ kg bw Skin Sens. 1A; H317: C ≥ 0,0015 % M = 100 M =	
·613-1	I Kannoet	a 23 B-00	0L6004_	421eihr	H361d	GHS08	3H361d		oral:	
	(ISO); 3- hydrox methyl	y-5- isoxazo	le	Acute Tox. 4 Eye Dam. 1 Skin Sens. 1 Aquati Chroni 2	H302 H318 H317 H411	GHS05 GHS09 Dgr	7H302 5H318 9H317 H411		ATE = 1600 mg/ kg bw'	
'613-1 <u>1</u>	2beQQtb (ISO); trans-5 (4- chlorop			OSeQuati Acute 1	сН400 Н410	GHS09 Wng)H410		M = 1 M = 1'	

·613-20	cyclohexyl-4-methyl-2-oxo-3-thiazolidine-carboxamide Dayonet ozine (ISO); (E)-4,5-dihydro-6-methyl-4-(3-pyridylmethyletriazin-3(2H)-		Aquati Chroni 1 282r0 2 Repr. 2 Aquati Chroni 1 o)-1,2,4	H351 H361fq H410 c	GHS08 IGHS09 Wng	3H351 9H361fo H410	l	M = 1'	
·613-2	one 59x09x5t428-79 (ISO); reaction mass of: [2,4- dioxo- (2- propyn-1- yl)imidazolidin yl]methyl(1R)- chrysanthemat [2,4- dioxo- (2- propyn-1- yl)imidazolidin yl]methyl(1R)- chrysanthemat	n-3- .cis- e; n-3- .trans-		H332 H302 H371 (nervor system oral, inhalat H400 cH410	,	H332	,	inhalate ATE = 1,4 mg/L (dusts or mists) oral: ATE = 550 mg/ kg bw M = 10 M = 10'	ion:
·616-0	oxime; ethyl methyl ketoxime; ethyl methyl ketone oxime		7Carc. 1 Acute Tox. 4 Acute Tox. 3 STOT SE 3 STOT SE 1 STOT RE 2 Skin Irrit. 2	H312 H301 H336 H370 (upper respiratract) H373 (blood system	•	Н312	•	dermal ATE = 1100 mg/ kg bw oral: ATE = 100 mg/ kg bw'	:

				Eye Dam. 1 Skin Sens.					
⁶¹⁷⁻⁰	0 16∔\$(10 0÷,α	-201-27	%0 -43-:	3Org.	H242	GHS02	H242		
X		ylbenzy		Perox.	H360D	GHS08	H360D	•	ı
	peroxi	de	ĺ	F	H315	GHS07	H315		ı
				Repr.	H319	GHS09			ı
				1B	H411	Dgr	H411'		ı
				Skin					ı
				Irrit.					ı
				2					ı
				Eye					ı
				Irrit.					ı
				2					ı
				Aquati					ı
				Chroni	c				ı
				2					1

⁽³⁾ the entries corresponding to index numbers 601-064-00-8 and 607-693-00-4 are deleted.

- (1) OJ L 353, 31.12.2008, p. 1.
- (2) The opinions are accessible via the following website: https://echa.europa.eu/registry-of-clh-intentions-until-outcome/-/dislist/name/-/ecNumber/-/casNumber/-/dte_receiptFrom/-/ dte_receiptTo/-/prc_public_status/Opinion+Adopted/dte_withdrawnFrom/-/dte_withdrawnTo/-/ sbm_expected_submissionFrom/-/sbm_expected_submissionTo/-/dte_finalise_deadlineFrom/-/ dte_finalise_deadlineTo/-/haz_addional_hazard/-/lec_submitter/-/dte_assessmentFrom/-/ dte_assessmentTo/-/prc_regulatory_programme/-/
- (3) Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EEC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC (OJ L 396, 30.12.2006, p. 1).
- (4) Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1).

Changes to legislation:

There are currently no known outstanding effects for the Commission Delegated Regulation (EU) 2020/1182.