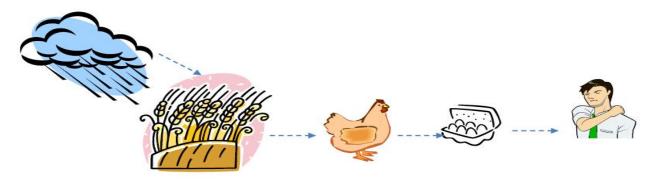


Consumer products HCT-202510-07

EU POPs Annex I restricted substances list --(Updated on October 29, 2025)

POPs is short for Persistant Organic Pollutants . The persistent organic pollutants are the natural or artificially synthesized organic pollutants that possess toxic properties, resist degradation, bioaccumulate and are transported, through air, water and migratory species, across international boundaries and deposited far from their place of release, where they accumulate in terrestrial and aquatic ecosystems.



November 2004, EU approved the STOCKHOLM CONVENTION ON PERSISTENT ORGANIC POLLUTANTS. From 20 May 2004, EU REGULATION (EC) No 850/2004 came into force, it bans the production, placing on the market and use of the listed substances.

25 June 2019, European Union published new regulation (EU) 2019/1021 on persistent organic pollutants (POPs), replaced previous (EC) No 850/2004. This new regulation was effective on 15 July 2019.

Annex I List of Restricted Substances

No.	SUBSTANCE	CAS No	EC No	SPECIFIC EXEMPTION ON INTERMEDIATE USE OR OTHER SPECIFICATION
1	Tetrabromodiphen yl ether $C_{12}H_6Br_4O$	40088-47-9 and others	254-787-2 and others	 For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of Tetrabromodiphenyl ether equal to or below 10 mg/kg (0,001 % by weight) where it is present in substances. For the purposes of the entries on tetra-, penta-, hexa-, hepta- and decaBDE, point (b) of Article 4(1) shall apply to the following sum of the concentrations of those substances: (a) 10 mg/kg upon entry into force of this Regulation where

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				they are present in mixtures or articles, except for food contact materials subject to Regulation (EC) No 1935/2004; (b) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, 350 mg/kg as of 30 December 2025 and 200 mg/kg as of 30 December 2025 where they are present in mixtures or articles containing or made of recovered material containing tetra-, penta-, hexa-, hepta- or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004; (c) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, 350 mg/kg as of 30 December 2025 and 10 mg/kg as of 17 May 2027, where they are present in toys subject to Directive 2009/48/EC, or in any product facilitating children's seating, sleep, relaxation, hygiene, changing and general body care, feeding, sucking, transportation and protection, containing or made of recovered material containing tetra-, penta-, hexa-, hepta-or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004. 3. By way of derogation, the manufacturing, placing on the market and use of the following shall be allowed: electrical and electronic equipment within the scope of Directive 2011/65/EC of the European Parliament and of the Council. 4. Use of articles already in use in the Union before 25 August 2010 containing Tetrabromodiphenyl ether shall be allowed. Article 4(2), third and fourth subparagraphs shall apply in relation to such articles.
2	Pentabromodiphen yl ether	32534-81-9	251-084-2	1. For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of pentabromodiphenyl ether equal to or below 10 mg/kg (0,001 % by weight) where it is present in substances.
	C ₁₂ H₅Br₅O	and others	and others	2. For the purposes of the entries on tetra-, penta-, hexa-, hepta- and decaBDE, point (b) of Article 4(1) shall apply to the following sum of the concentrations of those substances:

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				 (a) 10 mg/kg upon entry into force of this Regulation where they are present in mixtures or articles, except for food contact materials subject to Regulation (EC) No 1935/2004; (b) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, 350 mg/kg as of 30 December 2025 and 200 mg/kg as of 30 December 2027 where they are present in mixtures or articles containing or made of recovered material containing tetra-, penta-, hexa-, hepta- or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004; (c) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, 350 mg/kg as of 30 December 2025 and 10 mg/kg as of 17 May 2027, where they are present in toys subject to Directive 2009/48/EC, or in any product facilitating children 's seating, sleep, relaxation, hygiene, changing and general body care, feeding, sucking, transportation and protection, containing or made of recovered material containing tetra-, penta-, hexa-, hepta-or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004. 3. By way of derogation, the manufacturing, placing on the market and use of the following shall be allowed: electrical and electronic equipment within the scope of Directive 2011/65/EC. 4. Use of articles already in use in the Union before 25 August 2010 containing Pentabromodiphenyl ether shall be allowed. Article 4(2), third and fourth subparagraphs shall apply in relation to such articles. 1. For the purposes of this entry, point (b) of Article 4(1) shall
3	Hexabromodiphen yl ether $C_{12}H_4Br_6O$	36483-60-0 and others	253-058-6 and others	apply to concentrations of hexabromodiphenyl ether equal to or below 10 mg/kg (0,001 % by weight) where it is present in substances. 2. For the purposes of the entries on tetra-, penta-, hexa-, hepta- and decaBDE, point (b) of Article 4(1) shall apply to
				the following sum of the concentrations of those substances:

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	T			
				(a) 10 mg/kg upon entry into force of this Regulation where
				they are present in mixtures or articles, except for food
				contact materials subject to Regulation (EC) No 1935/2004;
				(b) by way of derogation from point (a), 500 mg/kg upon
				entry into force of this Regulation, 350 mg/kg as of 30
				December 2025 and 200 mg/kg as of 30 December 2027
				where they are present in mixtures or articles containing or
				made of recovered material containing tetra-, penta-, hexa-,
				hepta- or decaBDE, except for food contact materials subject
				to Regulation (EC) No 1935/2004;
				(c) by way of derogation from point (a), 500 mg/kg upon
				entry into force of this Regulation, 350 mg/kg as of 30
				December 2025 and 10 mg/kg as of 17 May 2027, where they
				are present in toys subject to Directive 2009/48/EC, or in any
				product facilitating children's seating, sleep, relaxation,
				hygiene, changing and general body care, feeding, sucking,
				transportation and protection, containing or made of
				recovered material containing tetra-, penta-, hexa-, hepta-
				or decaBDE, except for food contact materials subject to
				Regulation (EC) No 1935/2004.
				3. By way of derogation, the manufacturing, placing on the
				market and use of the following shall be allowed:
				electrical and electronic equipment within the scope of
				Directive 2011/65/EC.
				4. Use of articles already in use in the Union before 25 August
				2010 containing Hexabromodiphenyl ether shall be
				allowed. Article 4(2), third and fourth subparagraphs shall
				apply in relation to such articles.
				1. For the purposes of this entry, point (b) of Article 4(1) shall
				apply to concentrations of heptabromodiphenyl ether equal
	 Heptabromodiphe			to or below 10 mg/kg (0,001 % by weight) where it is
	nyl ether	68928-80-3	273-031-2	present in substances.
4	$C_{12}H_3Br_7O$	and others	and others	2. For the purposes of the entries on tetra-, penta-, hexa-,
	- 125 / -			hepta- and decaBDE, point (b) of Article 4(1) shall apply to
				the following sum of the concentrations of those
				substances:
	1	<u> </u>	<u> </u>	200210110021

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				(a) 10 mg/kg upon entry into force of this Regulation where
				they are present in mixtures or articles, except for food
				contact materials subject to Regulation (EC) No 1935/2004;
				(b) by way of derogation from point (a), 500 mg/kg upon
				entry into force of this Regulation, 350 mg/kg as of 30
				December 2025 and 200 mg/kg as of 30 December 2027
				where they are present in mixtures or articles containing or
				made of recovered material containing tetra-, penta-, hexa-,
				hepta- or decaBDE, except for food contact materials subject
				to Regulation (EC) No 1935/2004;
				(c) by way of derogation from point (a), 500 mg/kg upon
				entry into force of this Regulation, 350 mg/kg as of 30
				December 2025 and 10 mg/kg as of 17 May 2027, where they
				are present in toys subject to Directive 2009/48/EC, or in any
				product facilitating children's seating, sleep, relaxation,
				hygiene, changing and general body care, feeding, sucking,
				transportation and protection, containing or made of
				recovered material containing tetra-, penta-, hexa-, hepta-
				or decaBDE, except for food contact materials subject to
				Regulation (EC) No 1935/2004.
				3. By way of derogation, the manufacturing, placing on the
				market and use of the following shall be allowed: electrical
				and electronic equipment within the scope of Directive
				2011/65/EC.
				4. Use of articles already in use in the Union before 25 August
				2010 containing Heptabromodiphenyl ether shall be
				allowed. Article 4(2), third and fourth subparagraphs shall
				apply in relation to such articles.
				1. For the purposes of this entry, point (b) of Article 4(1) shall
				apply to concentrations of decaBDE equal to or below 10
	Bis(pentabromoph			mg/kg (0,001 % by weight) where it is present in
	enyl) ether			substances.
5	(decabromodiphen	1163-19-5	214-604-9	2. For the purposes of the entries on tetra-, penta-, hexa-,
	yl ether; decaBDE)			hepta- and decaBDE, point (b) of Article 4(1) shall apply to
	,,,,			the following sum of the concentrations of those
				substances:
	1	<u> </u>	I	

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(a) 10 mg/kg upon entry into force of this Regulation where they are present in mixtures or articles, except for food contact materials subject to Regulation (EC) No 1935/2004; (b) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, 350 mg/kg as of 30 December 2025 and 200 mg/kg as of 30 December 2027 where they are present in mixtures or articles containing or made of recovered material containing tetra-, penta-, hexa-, hepta- or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004;

(c) by way of derogation from point (a), 500 mg/kg upon entry into force of this Regulation, **350 mg/kg** as of 30 December 2025 and **10 mg/kg** as of 17 May 2027, where they are present in toys subject to Directive 2009/48/EC, or in any product facilitating children 's seating, sleep, relaxation, hygiene, changing and general body care, feeding, sucking, transportation and protection, containing or made of recovered material containing tetra-, penta-, hexa-, hepta-or decaBDE, except for food contact materials subject to Regulation (EC) No 1935/2004.

- 3. By way of derogation, the manufacturing, placing on the market and use of decaBDE shall be allowed for the following purposes, provided that Member States report to the Commission by December 2019 in accordance with the Convention:
 - (a) in the manufacturing of an aircraft, for which type approval has been applied for before 2 March 2019 and has been received before December 2022, until 18 December 2023, or, in cases where the continuing need is justified, until 2 March 2027;
 - (b) in the manufacturing of spare parts for either of the following:
 - (i) an aircraft, for which type approval has been applied for before 2 March 2019 and has been received before December 2022, produced before 18 December 2023, or, in cases where the continuing need is justified, produced

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before 2 March 2027, until the end of service life of th	at
aircraft;	

- (ii) motor vehicles within the scope of Directive 2007/46/EC of the European Parliament and of the Council, produced before 15 July 2019, either until 2036 or the end of service life of those motor vehicles, whichever date comes earlier;
- (c) electric and electronic equipment within the scope of Directive 2011/65/EC.
- 4. The specific exemptions for spare parts for use in motor vehicles referred to in point 2(b)(ii) shall apply for the manufacturing and use of commercial decaBDE falling into one or more of the following categories:
 - (a) powertrain and under-hood applications such as battery mass wires, battery interconnection wires, mobile air condition (MAC) pipes, powertrains, exhaust manifold bushings, under-hood insulation, wiring and harness under-hood (engine wiling, etc.), speed sensors, hoses, fan modules and knock sensors;
 - (b) fuel system applications such as fuel hoses, fuel tanks and fuel tanks under body;
 - (c) pyrotechnical devices and applications affected by pyrotechnical devices such as airbag ignition cables, seat covers/fabrics, only if airbag relevant and airbags (front and side).
- 5. Use of articles already in use before 15 July 2019 in the Union containing decaBDE shall be allowed. Article 4(2), third and fourth subparagraphs shall apply in relation to such articles.
- 6. Without prejudice to the application of other Union provisions on the classification, packaging and labelling of substances and mixtures, articles in which decaBDE is used shall be identifiable by labelling or other means throughout its life cycle.
- 7. The placing on the market and use of articles containing decaBDE imported for the purposes of the specific

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				exemptions in point 2 shall be allowed until the expiry of those exemptions. Point 6 shall apply as if such articles were produced pursuant to the exemption in point 2. Such articles already in use by the date of expiry of the relevant exemption may continue to be used. 8. For the purposes of this entry 'aircraft' means the following: (a) a civil aircraft produced in accordance with a type certificate issued under Regulation (EC) No 216/2008 of the European Parliament and of the Council (3) or with a design approval issued under the national regulations of a contracting state of ICAO, or for which a certificate of airworthiness has been issued by an ICAO Contracting State under Annex 8 to the Convention on International Civil Aviation;
6	Hexabromocyclodo decane 'Hexabromocyclodo odecane' means: hexabromocyclodo decane, 1,2,5,6,9,10-hexab romocyclododecan e and its main diastereoisomers: alpha-hexabromocyclododecane; beta-hexabromocyclododecane; and gamma-hexabrom ocyclododecane	25637-99-4 , 3194-55-6, 134237-50- 6, 134237-51- 7, 134237-52- 8	247-148-4, 221-695-9	 (b) a military aircraft. For the purposes of this entry, Article 4(1), point (b), shall apply to concentrations of hexabromocyclododecane equal to or below 75 mg/kg (0,0075 % by weight) where it is present in substances, mixtures, articles or as constituents of the flame-retarded articles. For the use of recycled polystyrene in the production of EPS and XPS insulation material for use in buildings or civil engineering works, that point (b) shall apply to concentrations of hexabromocyclododecane equal to or below 100 mg/kg (0,01 % by weight). The exemptions laid down in this point (1) shall be reviewed and assessed by the Commission by 1 January 2026. Expanded polystyrene articles containing hexabromocyclododecane already in use in buildings before 21 February 2018 in accordance with Commission Regulation (EU) 2016/293 (5) and Commission Implementing Decision No 2016/C 12/06 (6), and extruded polystyrene articles containing hexabromocyclododecane already in use in buildings before 23 June 2016 may continue to be used. Article 4(2), third and fourth

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7	Hexabromobiphen yl	36355-01-8	252-994-2	subparagraphs shall apply to such articles. 3. Without prejudice to the application of other Union provisions on the classification, packaging and labelling of substances and mixtures, expanded polystyrene placed on the market after 23 March 2016 in which hexabromocyclododecane was used shall be identifiable by labelling or other means throughout its life cycle.
8	Toxaphene	8001-35-2	232-283-3	1
9	Mirex	2385-85-5	219-196-6	1
10	Aldrin	309-00-2	206-215-8	1
11	Heptachlor	76-44-8	200-962-3	/
12	Dieldrin	60-57-1	200-484-5	/
13	Endrin	72-20-8	200-775-7	/
14	DDT (1,1,1-trichloro-2,2 -bis(4-chloropheny l)ethane)	50-29-3	200-024-3	
15	Chlordane	57-74-9	200-349-0	/
16	Hexachlorocyclohe xanes, including lindane	58-89-9; 319-84-6; 319-85-7; 608-73-1	200-401-2; 206-270-8; 206-271-3; 210-168-9	/
17	Chlordecone	143-50-0	205-601-3	/
18	Endosulfan	115-29-7 959-98-8 33213-65-9	204-079-4	 Placing on the market and use of articles already in use before or on 10 July 2012 containing endosulfan shall be allowed. Article 4(2), third and fourth subparagraphs shall apply to articles referred to in point 1.
19	Dicofol	115-32-2	204-082-0	None
20	Polychlorinated Biphenyls (PCB)	1336-36-3 and others	215-648-1 and others	Without prejudice to Directive 96/59/EC, articles already in use at the time of the entry into force of this Regulation are allowed to be used. Member States shall identify and remove from use equipment (e.g. transformers, capacitors or other receptacles containing

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				liquid stocks) containing more than 0,005 % PCBs and volumes
				greater than 0,05 dm³, as soon as possible but no later than 31
				December 2025.
				For the purposes of this entry, Article 4(1), point (b), shall apply
21	Hexachlorobenzen	110 74 1	204 272 0	to concentrations of hexachlorobenzene equal to or below 10
21	е	118-74-1	204-273-9	mg/kg (0,001 % by weight) where it is present in substances,
				mixtures or articles.
	Pentachlorobenzen	600.03.5	210 172 0	,
22	е	608-93-5	210-172-0	/
				1. Placing on the market and use of articles already in use
	in a ditable to the P			before or on 10 July 2012 containing hexachlorobutadiene
23	Hexachlorobutadi	87-68-3	201-765-5	shall be allowed.
	ene			2. Article 4(2), third and fourth subparagraphs shall apply to
				articles referred to in point 1.
				1. Placing on the market and use of articles already in use
	Polychlorinated naphthalenes			before or on 10 July 2012 containing polychlorinated
24		70776-03-3 and others	274-864-4 and others	naphthalenes shall be allowed.
				2. Article 4(2), third and fourth subparagraphs shall apply to
				articles referred to in point 1.
				For the purposes of this entry, point (b) of Article 4(1) shall
	Pentachlorophenol	87-86-5	201-778-6	apply to concentrations of pentachlorophenol and its salts and
25	and its salts and	and others	and others	esters equal to or below 5 mg/kg (0,0005 % by weight) where
	esters			they are present in substances, mixtures or articles.
				By way of derogation, the manufacturing, placing on the
				market and use of substances or mixtures containing SCCPs
				in concentrations lower than 1 % by weight or articles
				containing SCCPs in concentrations lower than 0,15 % by
				weight shall be allowed.
	Alkanes C ₁₀ -C ₁₃ ,			2. Use shall be allowed in respect of:
26	chloro (short-chain	85535-84-8	287-476-5	(a) conveyor belts in the mining industry and dam sealants
20	chlorinated	and others	20, 4,0 3	containing SCCPs already in use before or on 4
	paraffins) (SCCPs			December 2015; and
				(b) articles containing SCCPs other than those referred to in
				point (a) already in use before or on 10 July 2012.
				3. The third and fourth subparagraphs of Article 4(2) shall
				apply to the articles referred to in point 2.

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27	Perfluorooctane sulfonic acid (PFOS), its salts and PFOS-related compounds $C_8F_{17}SO_2X$ (X = OH, Metal salt (O-M+), halide, amide, and other related compounds including polymers)	2795-39-3 29457-72-5 29081-56-9 70225-14-8 56773-42-3 251099-16-8 4151-50-2 31506-32-8 1691-99-2 24448-09-7 307-35-7 and others	217-179-8 220-527-1 249-644-6 249-415-0 274-460-8 260-375-3 223-980-3 250-665-8 216-887-4 246-262-1 206-200-6 and others	 For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of PFOS or any of its salts equal to or below 0,025 mg/kg (0,0000025 % by weight) where they are present in substances, mixtures or in articles. For the purposes of this entry, Article 4(1), point (b), shall apply to the sum of concentrations of all PFOS-related compounds equal to or below 1 mg/kg (0,0001 % by weight) where they are present in substances, mixtures or articles. Use of articles already in use in the Union before 25 August2010 containing PFOS shall be allowed. Article 4(2), third and fourth subparagraphs shall apply in relation to such articles. Deleted. For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of PFOA or any of its salts equal to
28	Perfluorooctanoic acid (PFOA), its salts and PFOA-related compounds	335-67-1 and others	206-397-9 and others	or below 0,025 mg/kg (0,0000025 % by weight) where they are present in substances, mixtures or articles. 2. For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of any individual PFOA-related compound or a combination of PFOA-related compounds equal to or below 1 mg/kg (0,0001 % by weight) where they are present in substances, mixtures or articles. 3. For the purposes of this entry, point (b) of Article 4(1) shall apply to concentrations of PFOA-related compounds equal to or below 20 mg/kg (0,002 % by weight) where they are present in a substance to be used as a transported isolated intermediate within the meaning of Article 3 point 15(c) of Regulation (EC) No 1907/2006 and fulfilling the strictly controlled conditions set out in Article 18(4)(a) to (f) of that Regulation for the production of fluorochemicals with a carbon chain equal to or shorter than 6 atoms. 4. For the purposes of this entry, Article 4(1), point (b), shall apply to concentrations of PFOA and its salts equal to or below 1 mg/kg (0,0001 % by weight) where they are present in polytetrafluoroethylene (PTFE) micropowders

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produced by ionising irradiation or by thermal degradation as well as in mixtures and articles for industrial and professional uses containing PTFE micropowders until 18 August 2023. All emissions of PFOA during the manufacture and use of PTFE micropowders shall be avoided and, if not possible, reduced as far as possible. The limit of 1 mg/kg (0,0001 % by weight) shall apply only to manufacture, placing on the market and use of PFOA and its salts where they are present in PTFE micropowders that are transported or treated for the purpose to reduce the concentration of PFOA and its salts below the limit of 0,025 mg/kg (0,0000025 % by weight).

4a. For the purposes of this entry, Article 4(1), point (b), shall apply to concentrations of PFOA or any of its salts equal to or below 1 mg/kg (0,0001 % by weight) and to concentrations of any individual PFOA-related compound or combination of PFOA-related compounds equal to or below 10 mg/kg (0,001 % by weight) where they are present in fire-fighting foam for liquid fuel vapour suppression and liquid fuel fire (Class B fires) already installed in systems. This limit value shall apply until 3 August 2028;

4b. For the purposes of this entry, Article 4(1), point (b), shall apply to the sum of the concentration of PFOA, its salts and PFOA-related compounds equal to or below 10 mg/kg (0,001 % by weight) where they are present in fluorine-free fire-fighting foam and originate from fire-fighting equipment which has undergone cleaning in accordance with the best available techniques.

- 5. By way of derogation, the manufacturing, placing on the market and use of PFOA, its salts and PFOA-related compounds shall be allowed for the following purposes:
 - (a) photolithography or etch processes in semiconductor manufacturing, until 4 July 2025;
 - (b) photographic coatings applied to films, until 4 July 2025;
 - (c) textiles for oil- and water-repellency for the protection of workers from dangerous liquids that comprise risks to

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their health and safety, u	until 4 Jul	2023
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- (d) invasive and implantable medical devices, until 4 July 2025;
- 6. By way of derogation, the use of PFOA, its salts and PFOA-related compounds shall be allowed in fire-fighting foam for liquid fuel vapour suppression and liquid fuel fire (Class B fires) already installed in systems, including both mobile and fixed systems, until 3 December 2025, subject to the following conditions:
 - (a) fire-fighting foam that contains or may contain PFOA, its salts and/or PFOA-related compounds shall not be used for training;
 - (b) fire-fighting foam that contains or may contain PFOA, its salts and/or PFOA-related compounds shall not be used for testing unless all releases are contained;
 - (c) as from 1 January 2023, uses of fire-fighting foam that contains or may contain PFOA, its salts and/or PFOA-related compounds shall only be allowed in sites where all releases can be contained;
 - (d) fire-fighting foam stockpiles that contain or may contain PFOA, its salts and/or PFOA-related compounds shall be managed in accordance with Article 5.

"firefighting foam" means any mixture to fight fires with foam and includes but is not limited to firefighting foam concentrates and firefighting foam solutions to produce the foam.

- 7. By way of derogation, the use of perfluoroctyl bromide containing perfluoroctyl iodide for the purpose of producing pharmaceutical products shall be allowed, subject to review and assessment by the Commission by 31 December 2026, every four years thereafter and by 31 December 2036.
- 8. Use of articles already in use in the Union before 4 July 2020 containing PFOA, its salts and/or PFOA-related compounds shall be allowed. Article 4(2), third and fourth subparagraphs shall apply in relation to such articles.
- 9. By way of derogation, the use of PFOA, its salts and/or

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				PFOA-related compounds shall be allowed until 3
				December 2020 in the following articles:
				(a) medical devices other than implantable ones, within the
				scope of Regulation (EU) 2017/745 ;
				(b) latex printing inks;
				(c) plasma nano-coatings.
				10. For the purposes of this entry, point (b) of Article 4(1) shall
				apply to concentrations of PFOA and its salts and/or
				PFOA-related compounds equal to or below 2 mg/kg (0,0002 %
				by weight) where they are present in medical devices other
				than invasive devices and implantable devices.
				11. Articles containing PFOA, its salts or PFOA-related
				compounds already in use in the Union before or on the date of
				expiry of the relevant exemption laid down in points 5(a) to (d)
L				may continue to be used.
	Perfluorohexane			1. For the purposes of this entry, Article 4(1), point (b), shall
	sulfonic acid			apply to concentrations of PFHxS or any of its salts equal to or
	(PFHxS), its salts		below 0,025 mg/kg (0,0000025 % by weight) where they are	
	and PFHxS-related			present in substances, mixtures or articles.
	compounds			2. For the purposes of this entry, Article 4(1), point (b), shall
	"Perfluorohexane			apply to the sum of concentrations of all PFHxS-related
	sulfonic acid			compounds equal to or below 1 mg/kg (0,0001 % by weight)
	(PFHxS), its salts		206-587-1	where they are present in substances, mixtures or articles.
	and PFHxS-related			3. For the purposes of this entry, Article 4(1), point (b), shall
	compounds"			apply to concentrations of PFHxS, its salts and PFHxS-related
29	means the	355-46-4		compounds equal to or below 0,1 mg/kg (0,00001 % by weight)
	following:	and others	and others	where it is present in concentrated firefighting foam mixtures
	(i)			that are to be used or are used in the production of other
	perfluorohexane			firefighting foam mixtures. This exemption shall be reviewed
	sulfonic acid,			and assessed by the Commission no later than 28 August 2026.'
	including any of			,
	its branched			
	isomers;			
	(ii) its salts;			
	(iii) PFHxS-related			
	compounds			
	Compounds	I	<u>I</u>	1

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	which, for the purposes of the Convention, are any substance that contains the chemical moiety $C_6F_{13}S$ - as one of its structural elements and that degrades to PFHxS.			
30	Methoxychlor 'Methoxychlor' refers to any possible isomer of dimethoxydiphenyl trichloroethane or any combination thereof.	72-43-5 30667-99-3 76733-77-2 255065-25- 9 255065-26- 0 59424-81-6 1348358-72 -4 and others	200-779-9	For the purposes of this entry, Article 4(1), point (b), shall apply to concentrations of methoxychlor equal to or below 0,01 mg/kg (0,000001 % by weight) where they are present in substances, mixtures or articles.
31	2-(2H-benzotriazol -2-yl)-4,6- Ditertpentylphenol (UV-328)	25973-55-1	247-384-8	 For the purposes of this entry, Article 4(1), point (b), shall apply to concentrations of UV-328 equal to or below: (a) 100 mg/kg (0,01 % by weight) from 4 August 2025; (b) 10 mg/kg (0,001 % by weight) from 4 August 2027; (c) 1 mg/kg (0,0001 % by weight) from 4 August 2029; where they are present in substances, mixtures or articles. By way of derogation, the placing on the market of UV-328 present in articles and the use of such articles shall be allowed for the following purposes: (a) in land-based motor vehicles, until 4 August 2030; (b) in industrial coating for land-based motor vehicles, engineering machines, rail transportation vehicles, and in heavy-duty coatings for large steel structures, until 4 August 2030;

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			1	
				(c) in mechanical separators in blood collection tubes, until
				4 August 2030;
				(d) in triacetyl cellulose film in polarisers, until 4 August 2030;
				(e) in photographic paper, until 4 August 2030;
				(f) in civilian and military aircrafts, until 4 August 2030;
				(g) in spare parts for any of the following:
				(i) land-based motor vehicles;
				(ii) stationary industrial machines for use in agriculture,
				forestry and construction;
				(iii) liquid crystal displays in instruments for analysis,
				measurements, control, monitoring, testing, production and
				inspection, other than for medical applications;
				where UV-328 was initially used in their production, until the
				end of their service life or 31 December 2043, whichever comes
				earlier;
				(h) in spare parts for any of the following:
				(i) liquid crystal displays in devices within the scope of
				Regulation (EU) 2017/745, and within the scope of Regulation
				(EU) 2017/746 of the European Parliament and of the Council;
				(ii) liquid crystal displays in instruments for analysis,
				measurements, control, testing, production and inspection;
				where UV-328 was initially used in their production, until the
				end of their service life;
				(i) in spare parts for civilian and military aircrafts where UV-328
				was initially used in their production, until 31 December 2030.
				3. Articles containing UV-328 already in use in the Union before
				or on the date of expiry of the relevant exemption laid down in
				point 2(a) to (i) may continue to be used.
	'Dechlorane Plus	<mark>13560-89-9</mark>		1.For the purposes of this entry, Article 4(1), point (b), shall
				apply to concentrations of Dechlorane Plus:
<mark>32</mark>	"Dechlorane Plus"	<mark>135821-03-</mark>	236-948-9	(a) equal to or below 1 000 mg/kg (0,1 % by weight) where
	includes its	<mark>3</mark>		they are present in substances, mixtures or articles until 15
	syn-isomer and			April 2028;
	<mark>anti-isomer</mark>	<mark>135821-74-</mark>		(b) equal to or below 1 mg/kg (0,0001 % by weight) where they

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2028. 2.By way of derogation, the placing on the market and use of Dechlorane Plus shall be allowed for the following purposes; (a) aerospace, space and defence applications, until 26 February 2030; (b) medical imaging applications, until 26 February 2030; (c) radiotherapy devices and installations, until 26 February 2030; (d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of	8	are present in substances, mixtures or articles after 15 April
Dechlorane Plus shall be allowed for the following purposes: (a) aerospace, space and defence applications, until 26 February 2030; (b) medical imaging applications, until 26 February 2030; (c) radiotherapy devices and installations, until 26 February 2030; (d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is		
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February 2030; (b) medical imaging applications, until 26 February 2030; (c) radiotherapy devices and installations, until 26 February 2030; (d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph.		Dechlorane Plus shall be allowed for the following purposes:
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(c) radiotherapy devices and installations, until 26 February 2030; (d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph.		February 2030;
(d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of p		(b) medical imaging applications, until 26 February 2030;
(d) spare parts for, and the repair of, any of the following: (i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph.		(c) radiotherapy devices and installations, until 26 February
(i) land-based motor vehicles; (ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemptions in points (a), (b) and (c) of paragraph in the specific exemption in points (a), (b) and (c) of paragra		<mark>2030;</mark>
(ii) stationary industrial machines for use in agriculture, forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is the service of the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in points (a), (b) and (c) of paragraph is the specific exemptions in paragraph is the specific exemption in pa		(d) spare parts for, and the repair of, any of the following:
forestry and construction; (iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is		(i) land-based motor vehicles;
(iii) marine, garden, forestry and outdoor power equipment other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is		(ii) stationary industrial machines for use in agriculture,
other than that covered under (ii); (iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph a		forestry and construction;
(iv) aerospace, space and defence applications; (v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is		(iii) marine, garden, forestry and outdoor power equipment
(v) instruments for analysis, measurements, control, monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph is		other than that covered under (ii);
monitoring, testing, production and inspection; where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(iv) aerospace, space and defence applications;
where Dechlorane Plus was initially used in their production, until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(v) instruments for analysis, measurements, control,
until the end of their service life or until 31 December 2043, whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		monitoring, testing, production and inspection;
whichever comes earlier: (e) spare parts for, and the repair of, any of the following: (i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		where Dechlorane Plus was initially used in their production,
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(i) medical devices and accessories for medical devices within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		whichever comes earlier:
within the scope of Regulation (EU) 2017/745; (ii) in vitro diagnostic medical devices and accessories for invitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(e) spare parts for, and the repair of, any of the following:
(ii) in vitro diagnostic medical devices and accessories for in vitro diagnostic medical devices within the scope of Regulation (EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(i) medical devices and accessories for medical devices
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(EU) 2017/746; where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(ii) in vitro diagnostic medical devices and accessories for in
where Dechlorane Plus was initially used in their production, until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		vitro diagnostic medical devices within the scope of Regulation
until the end of their service life. 3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		(EU) 2017/746;
3. The Commission shall assess the need for a prolongation of the specific exemptions in points (a), (b) and (c) of paragraph 2		where Dechlorane Plus was initially used in their production,
the specific exemptions in points (a), (b) and (c) of paragraph 2		until the end of their service life.
		3. The Commission shall assess the need for a prolongation of
at the latest by 1 April 2028.		the specific exemptions in points (a), (b) and (c) of paragraph 2
		at the latest by 1 April 2028.
4. Articles containing Dechlorane Plus already in use in the		4. Articles containing Dechlorane Plus already in use in the
Union before or on the date of expiry of the relevant exemption		Union before or on the date of expiry of the relevant exemption

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	laid down in paragraph 2(a) to (d) may continue to be used.
	5. Placing on the market and use of spare parts containing
	Dechlorane Plus referred to in paragraph 2, point (d)(iv) that
	are present in the territory of the Union before or on 31
	December 2043 shall be allowed.'

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