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Acts whose titles are printed in light type are those relating to day-to-day management of agricultural matters, and are generally valid for a limited period.

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II

(Non-legislative acts)

INTERNATIONAL AGREEMENTS

Information relating to the entry into force of the Protocol between the European Union, the Swiss Confederation and the Principality of Liechtenstein to the Agreement between the European Community and the Swiss Confederation concerning the criteria and mechanisms for establishing the State responsible for examining a request for asylum lodged in a Member State or in Switzerland regarding access to Eurodac for law enforcement purposes

The Protocol between the European Union, the Swiss Confederation and the Principality of Liechtenstein to the Agreement between the European Community and the Swiss Confederation concerning the criteria and mechanisms for establishing the State responsible for examining a request for asylum lodged in a Member State or in Switzerland regarding access to Eurodac for law enforcement purposes, signed at Brussels on 27 June 2019 ⁽¹⁾, has entered into force on 1 May 2022, in accordance with Article 4(2) thereof, the procedures necessary for the entry into force having been completed on 28 April 2022.

⁽¹⁾ OJ L 32, 4.2.2020, p. 3.

Information relating to the entry into force of the Protocol between the European Union, Iceland and the Kingdom of Norway to the Agreement between the European Community and the Republic of Iceland and the Kingdom of Norway concerning the criteria and mechanisms for establishing the State responsible for examining a request for asylum lodged in a Member State or in Iceland or Norway regarding access to Eurodac for law enforcement purposes

The Protocol between the European Union, Iceland and the Kingdom of Norway to the Agreement between the European Community and the Republic of Iceland and the Kingdom of Norway concerning the criteria and mechanisms for establishing the State responsible for examining a request for asylum lodged in a Member State or in Iceland or Norway regarding access to Eurodac for law enforcement purposes, signed at Brussels on 24 October 2019 ⁽¹⁾, has entered into force on 1 September 2022, in accordance with Article 4(2) thereof, the procedures necessary for the entry into force having been completed on 9 August 2022.

⁽¹⁾ OJ L 64, 3.3.2020, p. 3.

REGULATIONS

COMMISSION REGULATION (EU) 2022/1531

of 15 September 2022

amending Regulation (EC) No 1223/2009 of the European Parliament and of the Council as regards the use in cosmetic products of certain substances classified as carcinogenic, mutagenic or toxic for reproduction and correcting that Regulation

(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 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products ⁽¹⁾, and in particular Article 15(1) and Article 15(2), fourth subparagraph, thereof,

Whereas:

- (1) Regulation (EC) No 1272/2008 of the European Parliament and of the Council ⁽²⁾ provides for a harmonised classification of substances as carcinogenic, mutagenic or toxic for reproduction (CMR) based on a scientific assessment by the Risk Assessment Committee of the European Chemicals Agency. The substances are classified as CMR substances of category 1A, CMR substances of category 1B or CMR substances of category 2 depending on the level of evidence of their CMR properties.
- (2) Article 15 of Regulation (EC) No 1223/2009 provides that substances which have been classified as CMR substances of category 1A, category 1B or category 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 (CMR substances) are to be prohibited from use in cosmetic products. A CMR substance may however be used in cosmetic products where the conditions laid down in Article 15(1), second sentence, of Regulation (EC) No 1223/2009 or in Article 15(2), second subparagraph, of that Regulation are fulfilled.
- (3) In order to uniformly implement the prohibition of CMR substances within the internal market, to ensure legal certainty in particular for economic operators and national competent authorities, and to ensure a high level of protection of human health, all CMR substances should be included in the list of prohibited substances in Annex II to Regulation (EC) No 1223/2009 and, where relevant, deleted from the lists of restricted or allowed substances in Annexes III to VI to that Regulation. Where the conditions laid down in Article 15(1), second sentence, of Regulation (EC) No 1223/2009 or in Article 15(2), second subparagraph, of that Regulation are fulfilled, the lists of restricted or allowed substances in Annexes III to VI to that Regulation should be amended accordingly.
- (4) This Regulation covers substances classified as CMR substances of categories 1A, 1B or 2 by Commission Delegated Regulation (EU) 2021/849 ⁽³⁾, which will apply from 17 December 2022.

⁽¹⁾ OJ L 342, 22.12.2009, p. 59.

⁽²⁾ 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 (OJ L 353, 31.12.2008, p. 1).

⁽³⁾ Commission Delegated Regulation (EU) 2021/849 of 11 March 2021 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 (OJ L 188, 28.5.2021, p. 27).

- (5) With regard to the substance 'methyl 2-hydroxybenzoate' (CAS No 119-36-8), with International Nomenclature Cosmetics of Ingredient (INCI) name 'Methyl Salicylate', which has been classified as a CMR substance of category 2 (Toxic for Reproduction), a request for the application of Article 15(1), second sentence, of Regulation (EC) No 1223/2009 was submitted on 25 May 2021 concerning use of that substance as a fragrance ingredient in various cosmetic products.
- (6) Methyl Salicylate is used as a fragrance ingredient, flavouring agent and soothing agent in various cosmetic products and is currently not listed in the Annexes to Regulation (EC) No 1223/2009.
- (7) In accordance with Article 15(1), second sentence, of Regulation (EC) No 1223/2009, CMR substances of category 2 may be used in cosmetic products where the substance has been evaluated by the Scientific Committee on Consumer Safety (SCCS) and found safe for use in such products.
- (8) The SCCS concluded in its opinion of 26–27 October 2021 ⁽⁴⁾ that Methyl Salicylate can be considered safe as an ingredient in cosmetic products up to the maximum concentrations provided by the applicant. In light of the classification of Methyl Salicylate as a CMR substance of category 2 and the SCCS opinion in its final version, Methyl Salicylate should be added to the list of substances restricted in cosmetic products in Annex III to Regulation (EC) No 1223/2009.
- (9) With regard to all substances other than Methyl Salicylate, which were classified as CMR substances pursuant to Regulation (EC) No 1272/2008 by Delegated Regulation (EU) 2021/849, no request for use in cosmetic products by way of exception has been submitted. Consequently, the CMR substances that are not already listed in Annex II to Regulation (EC) No 1223/2009 should be added to the list of substances prohibited in cosmetic products in that Annex.
- (10) The substance Sodium N-(hydroxymethyl)glycinate (CAS No 70161-44-3) was classified as carcinogenic of category 1B and mutagenic of category 2 by Commission Delegated Regulation (EU) 2020/1182 ⁽⁵⁾. It follows from notes 8 and 9 to that classification that it is only applicable if it cannot be shown that the maximum theoretical concentration of releasable formaldehyde, irrespective of the source, in the mixture as placed on the market is less than 0,1 %. In Commission Regulation (EU) 2021/1902 ⁽⁶⁾, 'Sodium N-(hydroxymethyl)glycinate' was erroneously added as entry 1669 in the list of prohibited substances in Annex II to Regulation (EC) No 1223/2009, despite the fact that it was already listed in entry 51 of Annex V to that Regulation under the chemical name/INN 'Sodium hydroxymethylamino acetate' as a preservative allowed in cosmetic products under certain conditions. A substance should not be listed in both Annex II and Annex V to Regulation (EC) No 1223/2009 and entry 1669 should therefore be deleted from Annex II to that Regulation.
- (11) The additional condition that was introduced in column h of entry 51 of Annex V to Regulation (EC) No 1223/2009 by Regulation (EU) 2021/1902 concerning the maximum theoretical concentration of formaldehyde was erroneously given a slightly different wording than the condition set out in notes 8 and 9 to the CMR classification of 'Sodium N-(hydroxymethyl)glycinate'. In order to correctly reflect the prohibition of that substance in cosmetic products on the basis of the CMR classification, the wording of the conditions should be the aligned and entry 51 should be adapted accordingly.
- (12) Entry 51 of Annex V to Regulation (EC) No 1223/2009 also contains an error in column b with regard to the chemical name of the substance. The correct name of the substance is 'Sodium N-(hydroxymethyl)glycinate', as referred to in Delegated Regulation (EU) 2020/1182.

⁽⁴⁾ SCCS (Scientific Committee on Consumer Safety), Opinion on methyl salicylate (methyl 2-hydroxybenzoate), preliminary version of 24–25 June, final version of 26–27 October 2021, SCCS/1633/21.

⁽⁵⁾ 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 (OJ L 261, 11.8.2020, p. 2).

⁽⁶⁾ Commission Regulation (EU) 2021/1902 of 29 October 2021 amending Annexes II, III and V to Regulation (EC) No 1223/2009 of the European Parliament and of the Council as regards the use in cosmetic products of certain substances classified as carcinogenic, mutagenic or toxic for reproduction (OJ L 387, 3.11.2021, p. 120).

- (13) Regulation (EC) No 1223/2009 should therefore be amended and corrected accordingly.
- (14) The amendments to Regulation (EC) No 1223/2009 that are based on the classifications of the relevant substances as CMR substances by Delegated Regulation (EU) 2021/849 should apply from the same date as those classifications.
- (15) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Cosmetic Products,

HAS ADOPTED THIS REGULATION:

Article 1

Annexes II and III to Regulation (EC) No 1223/2009 are amended in accordance with Annex I to this Regulation.

Article 2

Annexes II and V to Regulation (EC) No 1223/2009 are corrected in accordance with Annex II to this Regulation.

Article 3

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

Article 1 shall apply from 17 December 2022.

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

Done at Brussels, 15 September 2022.

For the Commission
The President
Ursula VON DER LEYEN

Annexes II and III to Regulation (EC) No 1223/2009 are amended as follows:

(1) in Annex II, the following entries are added:

Reference number	Substance identification		
	Chemical name/INN	CAS number	EC number
a	b	c	d
1681	Tetrafluoroethylene	116-14-3	204-126-9
1682	6,6'-di-tert-butyl-2,2'-methylenedi-p-cresol;[DBMC]	119-47-1	204-327-1
1683	(5-chloro-2-methoxy-4-methyl-3-pyridyl)(4,5,6-trimethoxy-o-tolyl)methanone; pyriofenone	688046-61-9	692-456-8
1684	(RS)-1-{1-ethyl-4-[4-mesyl-3-(2-methoxyethoxy)-o-toluoyl]pyrazol-5-yloxy}ethyl methyl carbonate; tolpyralate	1101132-67-5	701-225-3
1685	Azamethiphos (ISO); S-[(6-chloro-2-oxooxazolo[4,5-b]pyridin-3(2H)-yl)methyl] O,O-dimethyl thiophosphate	35575-96-3	252-626-0
1686	3-methylpyrazole	1453-58-3	215-925-7
1687	N-methoxy-N-[1-methyl-2-(2,4,6-trichlorophenyl)-ethyl]-3-(difluoromethyl)-1-methylpyrazole-4-carboxamide; pydiflumetofen	1228284-64-7	817-852-1
1688	N-{2-[[1,1'-bi(cyclopropyl)-2-yl]phenyl]-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamide; sedaxane	874967-67-6	688-331-2
1689	4-methylpentan-2-one; isobutyl methyl ketone (MIBK)	108-10-1	203-550-1
1690	Dimethomorph (ISO); (E,Z)-4-(3-(4-chlorophenyl)-3-(3,4-dimethoxyphenyl)acryloyl)morpholine	110488-70-5	404-200-2
1691	Imazamox (ISO); (RS)-2-(4-isopropyl-4-methyl-5-oxo-2-imidazolin-2-yl)-5-methoxymethylnicotinic acid	114311-32-9	601-305-7
1692	Thiamethoxam (ISO); 3-(2-chloro-thiazol-5-ylmethyl)-5-methyl[1,3,5]oxadiazinan-4-ylidene-N-nitroamine	153719-23-4	428-650-4
1693	Triticonazole (ISO); (RS)-(E)-5-(4-chlorobenzylidene)-2,2-dimethyl-1-(1H-1,2,4-triazol-1-methyl)cyclopentanol	138182-18-0	-
1694	Desmedipham (ISO); ethyl 3-phenylcarbamoyloxyphenylcarbamate	13684-56-5	237-198-5'

(2) in Annex III, the following entry is added:

Reference number	Substance identification				Restrictions			Wording of conditions of use and warnings
	Chemical Name/INN	Name of Common Ingredients Glossary	CAS number	EC number	Product type, Body parts	Maximum concentration in ready for use preparation	Other	
a	b	c	d	e	f	g	h	i
'324	Methyl 2-hydroxybenzoate	Methyl Salicylate	119-36-8	204-317-7	(a) Leave-on skin products (except face makeup, spray/aerosol body lotion, spray/aerosol deodorant and hydroalcoholic-based fragrances) and leave on hair products (except spray/aerosol products) (b) Face makeup (except lip products, eye makeup and makeup remover) (c) Eye makeup and makeup remover (d) Leave-on hair products (spray/aerosol) (e) Deodorant spray/aerosol (f) Body lotion spray/aerosol (g) Rinse-off skin products (except hand wash) and rinse-off hair products (h) Hand wash (i) Hydroalcoholic-based fragrances (j) Lip products (k) Toothpaste (l) Mouthwash intended for children aged 6–10 years (m) Mouthwash intended for children above 10 years of age and adults (n) Mouth spray	(a) 0,06 % (b) 0,05 % (c) 0,002 % (d) 0,009 % (e) 0,003 % (f) 0,04 % (g) 0,06 % (h) 0,6 % (i) 0,6 % (j) 0,03 % (k) 2,52 % (l) 0,1 % (m) 0,6 % (n) 0,65 %	Not to be used in preparations for children under 6 years of age, with the exception of (k) "Toothpaste".	

ANNEX II

Annexes II and V to Regulation (EC) No 1223/2009 are corrected as follows:

- (1) in Annex II, entry 1669 is deleted;
 (2) in Annex V, entry 51 is replaced by the following:

Reference number	Substance identification				Conditions			Wording of conditions of use and warnings
	Chemical name/INN	Name of Common Ingredients Glossary	CAS number	EC number	Product type, Body parts	Maximum concentration in ready for use preparation	Other	
a	b	c	d	e	f	g	h	i
'51	Sodium N-(hydroxymethyl) glycinate	Sodium Hydroxymethylglycinate	70161-44-3	274-357-8		0,5 %	Not to be used unless it can be shown that the maximum theoretical concentration of releasable formaldehyde, irrespective of the source, in the mixture as placed on the market is < 0,1 % w/w'	

DECISIONS

COUNCIL DECISION (EU) 2022/1532

of 24 January 2022

appointing a member proposed by Hungary, of the Committee of the Regions

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on the Functioning of the European Union, and in particular Article 305 thereof,

Having regard to Council Decision (EU) 2019/852 of 21 May 2019 determining the composition of the Committee of the Regions ⁽¹⁾,

Having regard to the proposal of the Hungarian Government,

Whereas:

- (1) Pursuant to Article 300(3) of the Treaty, the Committee of the Regions is to consist of representatives of regional and local bodies who either hold a regional or local authority electoral mandate or are politically accountable to an elected assembly.
- (2) On 20 January 2020 the Council adopted Decision (EU) 2020/102 ⁽²⁾ appointing the members and alternate members of the Committee of the Regions for the period from 26 January 2020 to 25 January 2025.
- (3) A member's seat on the Committee of the Regions has become vacant following the resignation of Mr János Ádám KARÁCSONY from the national mandate on the basis of which he was proposed for appointment.
- (4) The Hungarian Government has proposed, on the basis of a different national mandate, Mr János Ádám KARÁCSONY, representative of a regional body who holds a regional authority electoral mandate (*Pest Megyei Közgyűlés tagja* – Member of the County Council of Pest Megye), as a member of the Committee of the Regions for the remainder of the current term of office, which runs until 25 January 2025,

HAS ADOPTED THIS DECISION:

Article 1

Mr János Ádám KARÁCSONY, representative of a regional body who holds an electoral mandate, *Pest Megyei Közgyűlés tagja* (Member of the County Council of Pest Megye) (change of mandate) is hereby appointed as a member of the Committee of the Regions for the remainder of the current term of office, which runs until 25 January 2025.

Article 2

This Decision shall enter into force on the date of its adoption.

⁽¹⁾ OJ L 139, 27.5.2019, p. 13.

⁽²⁾ Council Decision (EU) 2020/102 of 20 January 2020 appointing the members and alternate members of the Committee of the Regions for the period from 26 January 2020 to 25 January 2025 (OJ L 20, 24.1.2020, p. 2).

Done at Brussels, 24 January 2022.

For the Council

The President

J. BORRELL FONTELLES

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