XIII. MEDICINAL PRODUCTS

The EFTA Surveillance Authority may designate, according to its working procedures, two observers entitled to participate in the tasks of the Committee which are described in Article 2, first indent of the Council Decision 75/320/EEC of 20 May 1975 setting up a pharmaceutical committee.

Notwithstanding Article 101 of the Agreement, the EC Commission shall invite experts from the EFTA States according to Article 99 of the Agreement, to participate in the tasks which are described in Article 2, second indent of the Council Decision 75/320/EEC.

The EC Commission shall, in due time, inform the EFTA Surveillance Authority about the date of the meeting of the Committee and transmit the relevant documentation.

For the purposes of this Chapter and notwithstanding the provisions of Protocol 1 to this Agreement, the terms Member States and competent authorities shall be understood to include, in addition to their meaning in the relevant EU acts, the EFTA States and their competent authorities, respectively.

When decisions on the authorisation of medicinal products are taken according to the Union procedures laid down in Regulation (EC) No 726/2004 of the European Parliament and of the Council, Directive 2001/83/EC of the European Parliament and of the Council, the EFTA States shall simultaneously and within 30 days of the Union Decision take corresponding decisions on the basis of the

\[1\] For the purposes of this Chapter and notwithstanding the provisions of Protocol 1 to this Agreement, the terms Member States and competent authorities shall be understood to include, in addition to their meaning in the relevant EU acts, the EFTA States and their competent authorities, respectively.

relevant acts. The EEA Joint Committee shall be informed and shall periodically publish lists of such decisions in the EEA Supplement to the *Official Journal of the European Union*.

The EFTA Surveillance Authority shall monitor the application of the decisions taken by the EFTA States as provided for in Article 109 of the Agreement.

Where any of the relevant acts provide for Union procedures on the granting, suspension and withdrawal of a marketing authorisation as well as supervision, including pharmacovigilance, and inspections and sanctions, these and similar tasks shall be carried out by the competent authorities in the EFTA States, on the basis of the same obligations as those of the competent authorities of EU Member States.

Should any disagreement between the contracting parties arise as to the administration of these provisions, Part VII of the Agreement shall apply *mutatis mutandis*.


The EFTA States shall therefore participate in the Union contribution referred to in Article 67(3) of Regulation (EC) No 726/2004 of the European Parliament and of the Council.

For this purpose, the procedures laid down in Article 82(1)(a) and Protocol 32 of the Agreement shall apply *mutatis mutandis* with regard to the financial contribution of the EFTA States to the above-mentioned Union contribution.

The EFTA States may send observers to meetings of the Agency's Management Board.

The EFTA States shall be fully associated with the work of the Committee for Medicinal Products for Human Use (CHMP), the Committee on Orphan Medicinal Products (COMP), the Paediatric Committee, the Pharmacovigilance Risk Assessment Committee (PRAC) and the Committee on Herbal Medicinal Products (HMPC). The provisions of Title IV, Chapter 1, of Regulation (EC) No 726/2004 of the European Parliament and of the Council shall apply to the members of the Committees appointed by the EFTA States. Members of the Committees appointed by the EFTA States shall have the same rights and obligations as the members appointed by the EU Member States, except for the right to vote. The positions of the members of the EFTA States shall be recorded separately. The position of Chairman shall be reserved for a member nominated by an EU Member State.

The EFTA States shall be fully associated with the work of the Committee for Veterinary Medicinal Products (CVMP). The provisions of Article 140 of Regulation (EU) 2019/6 of the European Parliament and of the Council shall apply to the members of the Committee appointed by the EFTA States. Members of the Committee appointed by the EFTA States shall have the same rights and obligations as the members appointed by the EU Member States, except for the right to vote. The positions of the members of the EFTA States shall be recorded separately. The position of Chairman shall be reserved for a member nominated by an EU Member State.

The EFTA States shall be fully associated with the work of the coordination groups as set up by Article 27 of Directive 2001/83/EC of the European Parliament and of the Council and Article 142 of Regulation (EU) 2019/6 of the European Parliament and of the Council. The members of the EFTA States shall not participate in the voting, however, their positions shall be recorded separately. The position of Chairman shall be reserved for a member nominated by an EU Member State. An objection raised by the competent authority of an EFTA State in accordance with Articles 49(5), 52(6), 53(8) or 66(8) of Regulation (EU) 2019/6 of the European Parliament and of the Council shall initiate the review procedure referred to in Article 54 of that Regulation.

An EFTA State may request the Agency to initiate an arbitration procedure according to Title III, Chapter 4 of Directive 2001/83/EC of the European Parliament and of the Council. Such a request shall, in the first place, be addressed to the Commission which shall, where it considers that the request is of common interest, forward it to the Agency for further processing.

The EFTA States shall participate fully in the Telematic Exchange of Information on Medicinal Products (IMP) programme.

Iceland and Norway shall provide their national competent authorities and the marketing authorisation holders with the linguistic version of the marketing authorisations required to access their own market.

A centralised marketing authorisation granted for a medicinal product shall not be subject to any fees other than those referred to in Article 67(3) of Regulation (EC) No 726/2004 of the European Parliament and of the Council.
The Agency having legal personality shall enjoy in all the States of the Contracting Parties the most extensive legal capacity accorded to legal persons under their laws.

The EFTA States shall grant privileges and immunities to the Agency equivalent to those contained in the Protocol on the Privileges and Immunities of the European Union.


By way of derogation from Article 12(2)(a) of the Conditions of employment of other servants of the European Communities, nationals of the EFTA States enjoying their full rights as citizens may be engaged under contract by the Executive Director of the Agency.

ACTS REFERRED TO


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

(a) References to other acts in the Regulation shall be considered relevant to the extent and in the form that those acts are incorporated into the Agreement.

(b) An EFTA State may request the Agency to issue an opinion according to Article 9(1), first paragraph of Article 11, Article 15(1) and Article 27(2). Such a request shall, in the first place, be addressed to the Commission which shall, where it considers that the request is of common interest, forward it to the Agency for further processing.


The provisions of the Regulation shall, for the purposes of the EEA Agreement, be read with the following adaptation:

References to other acts in the Regulation shall be considered relevant to the extent and in the form that those acts are incorporated into the Agreement.


(\textsuperscript{15}) Indent added by Decision No 82/2013 (OJ L 144, 30.5.2013, p. 12 and EEA Supplement No 31, 30.5.2013, p.14), e.i.f. 2.2.2013.

(\textsuperscript{16}) Indent added by Decision No 8/2013 (OJ L 144, 30.5.2013, p. 12 and EEA Supplement No 31, 30.5.2013, p.14), e.i.f. 2.2.2013.


(\textsuperscript{22}) Indent added by Decision No 189/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.

(\textsuperscript{23}) Indent added by Decision No 189/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.

(\textsuperscript{24}) Indent added by Decision No 189/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.

(\textsuperscript{25}) Indent added by Decision No 189/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.

(\textsuperscript{26}) Indent added by Decision No 190/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.


(\textsuperscript{28}) Indent added by Decision No 80/2014 (OJ L 310, 30.10.2014, p. 32 and EEA Supplement No 63, 30.10.2014, p. 24), e.i.f. 17.5.2014.

(\textsuperscript{29}) Indent added by Decision No 80/2014 (OJ L 310, 30.10.2014, p. 32 and EEA Supplement No 63, 30.10.2014, p. 24), e.i.f. 17.5.2014.
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(65) Indent added by Decision No 177/2015 (OJ L 8, 12.1.2017, p. 5 and EEA Supplement No 3, 12.1.2017, p. 5), e.i.f. 11.7.2015.


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(81) Indent added by Decision No 22/2017 (OJ L 297, 22.11.2018, p. 28 and EEA Supplement No 78, 22.11.2018, p. 34), e.i.f. 4.2.2017.


(93) Indent added by Decision No 62/2019 (OJ L 210, 2.7.2020, p. 29 and EEA Supplement No 44, 2.7.2020, p. 31), e.i.f. 30.3.2019.
14. [ ] [105]

15. [ ] [106]


15b. [ ] [108]


[98] Indent added by Decision No 153/2020 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 24.10.2020.

[99] Indent added by Decision No 198/2020 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 12.12.2020.

[100] Indent added by Decision No 198/2020 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 12.12.2020.


[102] Indent added by Decision No 106/2021 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 20.3.2021.

[103] Indent added by Decision No 203/2021 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 20.3.2021.

[104] Indent added by Decision No 302/2021 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 10.7.2021.


15c. [ ] [199]
15d. [ ] [199]
15e. [ ] [199]
15f. [ ] [113]
15g. [ ] [113]


15j. [ ] [133]

15k. [ ] [125]


\(^{[138]}\)The provisions of the Regulation shall, for the purposes of the present Agreement, be read with the following adaptation:


\(^{[123]}\) Indent added by Decision No 178/2014 (OJ L 202, 30.7.2015, p. 28 and EEA Supplement No 43, 30.7.2015, p. 28), e.i.f. 26.9.2014.

\(^{[124]}\) Indent added by Decision No 178/2015 (OJ L 8, 12.1.2017, p. 6 and EEA Supplement No 3, 12.1.2017, p. 6), e.i.f. 11.7.2015.


\(^{[128]}\) Indent added by Decision No 25/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.

\(^{[129]}\) Indent added by Decision No 26/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.


The EFTA States shall be fully associated with the work of the Committee for Orphan Medicinal Products. The detailed arrangements of participation for the representatives of EFTA States shall be in accordance with the provisions of Article 4 of the Regulation. Such representatives will, however, not participate in the voting and their positions shall be recorded separately. The position of Chairman shall be reserved for a member nominated by a Member State of the Community. The internal rules of procedure of this committee shall be amended in order to give full effect to the EFTA States' participation.

The EFTA States shall contribute financially to the work of the Committee in accordance with Article 82(1)(a) of the Agreement.

15n.\(^{[138]}\) 32000 R 0847: Commission Regulation (EC) No 847/2000 of 27 April 2000 laying down the provisions for implementation of the criteria for designation of a medical product as an orphan medicinal product and definitions of the concepts 'similar medicinal product' and 'clinical superiority' (OJ L 103, 28.4.2000, p. 5), as amended by:


15o. \([\text{ }][\text{ }]\) \(^{[138]}\)


\(^{[138]}\) The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for Lithuania (Annex IX, Chapter I, Point 1) and Poland (Annex XII, Chapter I, Point 4), shall apply.

\(^{[139]}\) The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

Liechtenstein shall not be obliged to participate in the decentralised procedure (DCP) and in the mutual recognition procedure (MRP) and shall, therefore, not be obliged to issue the corresponding marketing...


\textsuperscript{[154]} Indent added by Decision No 158/2013 (OJ L 58, 27.2.2014, p. 10 and EEA Supplement No 13, 27.2.2014, p. 11), c.f. 28.5.2014.


The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for Cyprus (Annex VII, Chapter 1), Lithuania (Annex IX, Chapter 1, Point 2), Malta (Annex XI, Chapter 1, Point 2), Poland (Annex XII, Chapter 1, Point 5) and Slovenia (Annex XIII, Chapter 1), shall apply.

The transitional arrangements set out in the Annexes to the Act of Accession of 9 December 2011 for Croatia (Annex V, Chapter 1) shall apply.

The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptations:

(a) Liechtenstein shall not be obliged to participate in the decentralised procedure (DCP) and in the mutual recognition procedure (MRP) and shall, therefore, not be obliged to issue the corresponding marketing authorisations. Instead, Austrian marketing authorisations within the DCP and the MRP shall be valid for Liechtenstein upon request of a marketing authorisation applicant.

(b) The EFTA States may initiate the urgent Union procedure pursuant to Section 4 of Chapter 3 of Title IX of the Directive.

(c) With respect to Title IX, the obligations of Liechtenstein will be executed by Austria. Liechtenstein will however, as far as applicable to Liechtenstein:

- operate a pharmacovigilance system in accordance with Article 101(1);
- perform a regular audit of its pharmacovigilance system in accordance with Article 101(2);
- designate a competent authority for the performance of its pharmacovigilance tasks in accordance with Article 101(3),
- take all appropriate measures to encourage patients, doctors, pharmacists and other healthcare professionals to report suspected adverse reactions to the national competent authority in accordance with Article 102(a);
- facilitate patient reporting through the provision of alternative reporting formats in addition to web-based formats in accordance with Article 102(b);
- impose an obligation on a marketing authorisation holder to operate a risk management system, as referred to in Article 104(3)(c), if there are concerns about the risks affecting the risk-benefit balance of an authorised medicinal product in accordance with Article 104a(2). For the imposition of such obligation Liechtenstein will base itself on a corresponding decision of the Austrian authorities;


Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2020.

\(\text{(\text{[159]}\text{)}}\) Indent added by Decision No 371/2021 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. pending.


- set up and maintain a national medicines web-portal which shall be linked to the European medicines web-portal in accordance with Article 106;

- record all suspected adverse reactions that occur in its territory which are brought to its attention from healthcare professionals and patients and ensure that reports of such reactions may be submitted by means of the national medicines web-portals or by other means in accordance with Article 107a(1), and

- submit reports in accordance with Article 107a(4).

(d) The following subparagraph shall be added to Article 107c(5):

“A Swiss marketing authorisation for a medicinal product taking effect in Liechtenstein by virtue of Liechtenstein law on the basis of the Customs Union between the Principality of Liechtenstein and the Swiss Confederation shall not be considered as a first authorisation to place a product on the market for the purposes of this paragraph.”


15qb. 32012 D 0715: Commission Implementing Decision 2012/715/EU of 22 November 2012 establishing a list of third countries with a regulatory framework applicable to active substances for medicinal products for human use and the respective control and enforcement activities ensuring a level of protection of public health equivalent to that in the Union, in accordance with Directive 2001/83/EC of the European Parliament and of the Council (OJ L 325, 23.11.2012, p. 15), as amended by:

- 32013 D 0262: Commission Implementing Decision 2013/262/EU of 4 June 2013 (OJ L 152, 5.6.2013, p. 52),


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[155] Inserted and words "as amended by:“, inserted by Decision No 191/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

The inspections in Liechtenstein shall be performed by the Swiss inspectorate on behalf of Liechtenstein on the basis of the agreement between Swissmedic and Amt für Gesundheit (Verwaltungsvereinbarung betreffend Inspektionsen).


15r. [178]  

15s. [179]  

15t. [180]  


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[175] Indent and words “, as amended by” added by Decision No 333/2021 (OJ L [to be published]) and EEA Supplement No [to be published]), c.f.f. 11.12.2021.
\textsuperscript{-183} 32011 L 0038: Commission Implementing Directive 2011/38/EU of 11 April 2011 (OJ L 97, 12.4.2011, p. 28),


\textsuperscript{183} Indent and words „as amended by:“ added by Decision No 46/2012 (OJ L 207, 2.8.2012, p. 26 and EEA Supplement No 43, 2.8.2012, p. 30) e.i.f. 1.5.2012.
\textsuperscript{184} Indent added by Decision No 89/2015 (OJ L 211, 4.8.2016, p. 26 and EEA Supplement No 42, 4.8.2016, p. 25) e.i.f. 1.5.2015.
\textsuperscript{191} Indent added by Decision No 50/2018 (OJ L 26, 30.1.2020, p. 30 and EEA Supplement No 6, 30.1.2020, p. 23), e.i.f. 24.3.2018.
\textsuperscript{192} Indent added by Decision No 151/2019 (OJ L 291, 10.11.2022, p. 28 and EEA Supplement No 74, 10.11.2022, p. 29), e.i.f. 15.6.2019.
\textsuperscript{193} Indent added by Decision No 238/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 28.10.2022.
\textsuperscript{194} Indent added by Decision No 303/2022 (OJ L 164, 29.6.2023, p. 27 and EEA supplement No 48, 29.6.2023, p. 27), e.i.f. 10.12.2022.
The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The text of Article 11 (2) shall not apply.

(b) In Article 14a a reference is made to the Committee set up by Article 30 of Council Regulation (EC) No 111/2005. When that Committee deals with matters covered by this Regulation the EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.

15xa.


The provisions of the Delegated Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

The Regulation shall only apply to the EEA EFTA States with regard to Regulation (EC) No 273/2004.

15xb.


The provisions of the Implementing Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

The Regulation shall only apply to the EEA EFTA States with regard to Regulation (EC) No 273/2004.

15y.


The provisions of this Agreement, as amended by:

15za.

veterinary medicinal products, a list of substances essential for the treatment of equidae (OJ L 367, 22.12.2006, p. 33), as amended by:


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

The powers vested in the European Commission in relation to the infringement procedure foreseen in Article 84a, including the power to impose financial penalties on the holders of marketing authorisations, shall, in the cases where the marketing authorisation holder is established in an EFTA State, be carried out by the EFTA Surveillance Authority in close cooperation with the Commission. Before the EFTA Surveillance Authority takes a decision regarding financial penalties, the Commission shall provide it with its assessment and a proposal on how to act.


15ze. [\[15ze\]]

15zf. [\[15zf\]]


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:
The EFTA States shall be fully associated with the work of the Committee for Advanced Therapy, but without the right to vote.


and Austria laying down the technical details for Liechtenstein’s recognition of Austrian marketing authorisations within the decentralized procedure (DCP) and the mutual recognition procedure (MRP).


\[216\] Three points and text inserted by Decision No 128/2009 (OJ L 62, 11.3.2010, p. 16 and EEA Supplement No 12, 11.3.2010, p. 15), e.i.f. 23.12.2009. For Liechtenstein, Decision 128/2009 entered into force 1.1.2010 which is the day of entry into force of the Agreement between Liechtenstein and Austria laying down the technical details for Liechtenstein’s recognition of Austrian marketing authorisations within the decentralized procedure (DCP) and the mutual recognition procedure (MRP).

\[217\] Indent and words “,” as amended by: “added by Decision No 158/2013 (OJ L 58, 27.2.2014, p. 10 and EEA Supplement No 13, 27.2.2014, p. 11), e.i.f. 28.5.2014.


\[219\] Indent added by Decision No 335/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:
The powers vested in the European Commission in relation to the infringement procedure, including the power to impose financial penalties on the holders of marketing authorisations, shall, in the cases where the marketing authorisation holder is established in an EFTA State, be carried out by the EFTA Surveillance Authority in close cooperation with the Commission. Before the EFTA Surveillance Authority takes a decision regarding financial penalties, the Commission shall provide it with its assessment and a proposal on how to act.

15zk. [ ] 15zk.


- 32012 D 0067: Commission Implementing Decision 2012/67/EU of 3 February 2012 (OJ L 34, 7.2.2012, p. 5),
- 32012 D 0068: Commission Implementing Decision 2012/68/EU of 3 February 2012 (OJ L 34, 7.2.2012, p. 8),


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

This Directive shall not apply to Liechtenstein, the exception of Articles 15 and 16. The definitions in Article 3 and general provisions in Articles 17(2)(h) and 23 shall only apply to Liechtenstein as far as necessary for transposing Articles 15 and 16 of the Directive.


The provisions of the Implementing Directive shall, for the purposes of this Agreement, be read with the following adaptation:

This Implementing Directive shall not apply to Liechtenstein, with the exception of Article 7.

15zo.\[24\] 32013 R 0198: Commission Implementing Regulation (EU) No 198/2013 of 7 March 2013 on the selection of a symbol for the purpose of identifying medicinal products for human use that are subject to additional monitoring (OJ L 65, 8.3.2013, p. 17).


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\[231\] Indent added by Decision No 304/2022 (OJ L 164, 29.6.2023, p. 28 and EEA supplement No 48, 29.6.2023, p. 28), e.i.f. 10.12.2022.


\[234\] Point inserted by Decision No 179/2014 (OJ L 202, 30.7.2015, p. 29 and EEA Supplement No 43, 30.7.2015, p. 29), e.i.f. 1.6.2015.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The application of Article 36(3) shall not be made dependent on an authorisation of the medicinal product in Liechtenstein.

(b) The powers vested in the European Commission in relation to the infringement procedure foreseen in Article 49(3), including the power to impose financial penalties on the holders of marketing authorisations, shall, in the cases where the marketing authorisation holder is established in an EFTA State, be carried out by the EFTA Surveillance Authority in close cooperation with the Commission. Before the EFTA Surveillance Authority takes a decision regarding financial penalties, the Commission shall provide it with its assessment and a proposal on how to act.


The provisions of the Delegated Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

The inspections in Liechtenstein shall be carried out, on behalf of Liechtenstein, by the Swiss inspectorate.


The provisions of this Regulations shall, for the purposes of the EEA Agreement, be read with the following adaptations:

(a) The following provisions of the Regulation regarding veterinary matters shall not apply to Liechtenstein:

- Article 57
- Articles 103 to 115, included.

(b) Liechtenstein shall not be obliged to participate in the decentralised procedure (DCP) and in the mutual recognition procedure (MRP) and shall, therefore, not be obliged to issue the corresponding marketing authorisations. Instead, Austrian marketing authorisations within the DCP and the MRP shall be valid for Liechtenstein upon request of a marketing authorisation applicant.

(c) In Article 136, the powers vested in the European Commission to impose financial penalties on the holders of marketing authorisations shall, in the cases where the marketing authorisation holder is established in an EFTA State, be carried out by the EFTA Surveillance Authority in close cooperation with the Commission. Before the EFTA Surveillance Authority takes a decision regarding financial penalties, the Commission shall provide it with its assessment and a proposal on how to act.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 6(4), the words “official languages of the Union” shall be replaced by “official languages of the Contracting Parties to the EEA Agreement”.


22h.\(^{[294]}\) 32022 R 0209: Commission Implementing Regulation (EU) 2022/209 of 16 February 2022 establishing the format of the data to be collected and reported in order to determine the volume of sales and the use of antimicrobial medicinal products in animals in accordance with Regulation (EU) 2019/6 of the European Parliament and of the Council (OJ L 35, 17.2.2022, p. 7).

\(^{[287]}\) Point inserted by Decision No 5/2022 (OJ L 175, 30.6.2022, p. 9 and EEA Supplement No 42, 30.6.2022, p. 8), e.i.f pending.

\(^{[288]}\) Point inserted by Decision No 5/2022 (OJ L 175, 30.6.2022, p. 9 and EEA Supplement No 42, 30.6.2022, p. 8), e.i.f pending.

\(^{[289]}\) Point inserted by Decision No 5/2022 (OJ L 175, 30.6.2022, p. 9 and EEA Supplement No 42, 30.6.2022, p. 8), e.i.f pending.

\(^{[290]}\) Point inserted by Decision No 5/2022 (OJ L 175, 30.6.2022, p. 9 and EEA Supplement No 42, 30.6.2022, p. 8), e.i.f pending.

\(^{[291]}\) Point inserted by Decision No 11/2022 (OJ L 246, 22.9.2022, p. 60 and EEA Supplement No 61, 22.9.2022, p. 59), e.i.f pending.

\(^{[292]}\) Point inserted by Decision No 11/2022 (OJ L 246, 22.9.2022, p. 60 and EEA Supplement No 61, 22.9.2022, p. 59), e.i.f pending.

\(^{[293]}\) Point inserted by Decision No 168/2022 (OJ L 267, 13.10.2022, p. 7 and EEA Supplement No 66, 13.10.2022, p. 7), e.i.f pending.


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**

The Contracting Parties take note of the content of the following acts:


2. (290) **C/115/82/p. 5:** Commission Communication on parallel imports of proprietary medicinal products for which marketing authorisation have already been granted (OJ No C 115, 6.5.1982, p. 5).


**XIV. FERTILIZERS**(272)

**ACTS REFERRED TO**


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286 Point inserted by Decision No 278/2022 (OJ L 117, 4.5.2023, p. 3 and EEA Supplement No 35, 4.5.2023, p. 3), e.i.f. 29.10.2022.


288 Point inserted by Decision No 8/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The EFTA States shall be free to continue to apply their national limit values for cadmium in phosphate fertilisers existing at the date of entry into force of Decision of the EEA Joint Committee No 77/2023 of 28 April 2023 until such time as harmonised limit values for cadmium content in phosphate fertilisers which are equal to or lower than those limit values become applicable in the European Economic Area.

(b) In Article 1(2), the following points shall be added after point (p):

“(q) the national plant health law of the EFTA States;

(r) the national law on invasive alien species of the EFTA States.”

(c) In Article 52, as regards the EFTA States, the words “or the date of entry into force of Decision of the EEA Joint Committee No 77/2023 of 28 April 2023, whichever is the later” are inserted after the words “16 July 2023”.


\[274\] Indent added by Decision No 80/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[275\] Indent added by Decision No 80/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[276\] Indent added by Decision No 80/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[277\] Indent added by Decision No 81/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[278\] Indent added by Decision No 82/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[279\] Point inserted by Decision No 83/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**

The Contracting Parties shall take note of the following acts:


**XV. DANGEROUS SUBSTANCES**

**ACTS REFERRED TO**

1. **294**
2. **295**
3. **296**
4. **297**
5. **298**
6. **299**

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**Notes:**

- **294** Point inserted by Decision No 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
- **295** Point inserted by Decision No 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
- **296** Point inserted by Decision No 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
- **297** Point inserted by Decision No 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
7. [ (296) ]


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12a. [ ]

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12d. [ ]

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12f. [ ]


12j. [ ]


\[\text{\textsuperscript{404}}\] Indent added by Decision No 159/2014 (OJ L 15, 22.1.2015, p. 87 and EEA Supplement No 5, 22.1.2015, p. 10), e.i.f. pending; it shall apply from 9.7.2014.


\[\text{\textsuperscript{406}}\] Point 12n (Directive 98/86/EC) inserted by Decision No 32/2003 (OJ L 137, 5.6.2003, p. 33 and EEA Supplement No 29, p. 21), e.i.f. 1.3.2004, and subsequently replaced by Decision No 225/2013 (OJ L 154, 22.5.2014, p. 22 and EEA Supplement No 29, 22.5.2014, p. 21), e.i.f. 1.6.2014. For Liechtenstein, this Decision shall enter into force on the same day or on the day of entry into force of the Agreement between Liechtenstein and Switzerland laying down the cooperation in the field of authorisation procedures for biocidal products according to Regulation (EU) No 528/2012, whichever is the later. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 6.7.2018.


The provisions of the Regulation shall, for the purpose of this Agreement, be read with the following adaptations:


(b) Notwithstanding the provisions of Protocol 1 to the Agreement, the term ‘Member State(s)’ contained in Regulation (EU) No 528/2012 shall be understood to include, in addition to its meaning in that Regulation, the EFTA States.

(c) As regards the EFTA States, the Agency shall, as and when appropriate, assist the EFTA Surveillance Authority or the Standing Committee, as the case may be, in the performance of their respective tasks.

(d) The following paragraph shall be added in Article 35:

“4. The EFTA States shall be entitled to participate fully in the work of the coordination group and shall within it have the same rights and obligations as EU Member States, except the right to vote. The rules of procedures of the coordination Group shall give full effect to the EFTA States’ participation.”

(e) The following subparagraph shall be added in Article 44(5):

“When the Commission grants a Union authorisation or decides that a Union authorisation has not been granted, the EFTA States will simultaneously and within 30 days of the Commission act take corresponding decisions. The EEA Joint Committee shall be informed and shall periodically publish lists of such decisions in the EEA Supplement to the Official Journal.”

(f) The following paragraph shall be added in Article 48:

“4. If the Commission cancels or amends a Union authorisation, the EFTA States shall cancel or amend the corresponding decision.”

(g) The following subparagraph shall be added in Article 49:

“If the Commission cancels a Union authorisation, the EFTA States shall cancel the corresponding decision.”

(h) The following paragraph shall be added in Article 50:

“4. If the Commission amends a Union authorisation, the EFTA states shall amend the corresponding decision.”

(i) The following paragraph shall be added in Article 75:

“5. The EFTA States shall be entitled to participate fully in the work of the Biocidal Products Committee and shall within it have the same rights and obligations as EU Member States, except the right to vote.”

(j) The following paragraph shall be added in Article 78:

“3. The EFTA States shall, as from the entry into force of this Decision participate in the financing of the Agency. For this purpose the procedures laid down in Article 82(1)(a) and Protocol 32 of the Agreement shall apply mutatis mutandis.”

(k) Should any disagreement between the Contracting Parties arise as to the administration of these provisions, Part VII of the Agreement shall apply mutatis mutandis.


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(351) Point inserted by Decision No 218/2014 (OJ L 230, 3.9.2015, p. 16 and EEA Supplement No 52, 3.9.2015, p. 16), e.i.f. 1.11.2014.
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30.4.2014, p. 68).

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17.10.2014, p. 15).

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and of the Council as regards the rules for the renu

ralisation of bicidal products subject to mutual recog


2014 D 0397: Commission Implementing Decision 2014/397/EU of 25 June 2014 postponing the expiry date of appr


2014 D 0402: Commission Implementing Decision 2014/402/EU of 25 June 2014 regarding restrictions of au

isations of bicidal products containing IPBC notified by Germany in accordance with Directive 98/8/EC of the European Parliamen


2015 D 0646: Commission Implementing Decision (EU) 2015/646 of 23 April 2015 pursuant to Article 3(3) of Regu

EU No 528/2012 of the European Parliament and of the Council on bacterial cultures intended to reduce organic solid

d to be placed on the market for that purpose (OJ L 106, 24.4.2015, p. 79).

\[32014 R 0437: Commission Implementing Regulation (EU) No 437/2014 of 29 April 2014 approving 4,5-

Dichloro-2-octyl-2H-isothiazol-3-one as an existing active substance for use in bicidal products for product-type 21 (OJ L 128, 30.4.2014, p. 64).\]


zole as an existing active substance for use in bicidal products for product-type 8 (OJ L 128, 30.4.2014, p. 68).\]


the systematic examination of all existing active substances contained in bicidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 294, 10.10.2014, p. 1), as amended by:\]


\[32015 D 0646: Commission Implementing Decision (EU) 2015/646 of 23 April 2015 pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on bacterial cultures intended to reduce organic solids and to be placed on the market for that purpose (OJ L 106, 24.4.2015, p. 79).\]
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- [ ] [389]
The provisions of the Regulation shall, for the purposes of the present Agreement, be read with the following adaptations, in addition to the adaptations to the Directive 98/8/EC:

(a) The EFTA States and their producers shall participate in the 10-year programme of work mentioned in Article 16(2) of Directive 98/8/EC. The EFTA States may be designated under Article 7(5) as responsible for the review of active substances under this programme.

(b) The Commission shall take into account information concerning active substances already on the market in the EFTA States on 14 May 2000 submitted under Article 3 paragraph 1 before the entry into force of the Joint Committee Decision integrating this Regulation into the Agreement.

(c) The Commission shall take into account notifications concerning active substances already on the market in the EFTA States on 14 May 2000 submitted under Article 4 paragraph 1 before the entry into force of the Joint Committee Decision integrating this Regulation into the Agreement.


\[\{403\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{404\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{405\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{406\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{407\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{408\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{409\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{410\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{411\} \quad \text{Indent added by Decision No 122/2014 (OJ L 342, 27.11.2014, p. 17 and EEA Supplement No 71, 27.11.2014, p. 17), e.i.f. 28.6.2014.}\]

\[\{412\} \quad \text{Indent added by Decision 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.}\]

\[\{413\} \quad \text{Indent added by Decision 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.}\]

\[\{414\} \quad \text{Indent added by Decision 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.}\]

\[\{415\} \quad \text{Indent added by Decision 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.}\]


\[^{416}\] Indent added by Decision No 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.

\[^{417}\] Indent added by Decision No 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.

\[^{418}\] Indent added by Decision No 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.

\[^{419}\] Indent added by Decision No 181/2014 (OJ L 202, 30.7.2015, p. 31 and EEA Supplement No 43, 30.7.2015, p. 31), e.i.f. 26.9.2014.

\[^{420}\] Indent added by Decision No 179/2015 (OJ L 8, 12.1.2017, p. 7 and EEA Supplement No 3, 12.1.2017, p. 7), e.i.f. 11.7.2015.

\[^{421}\] Indent added by Decision No 179/2015 (OJ L 8, 12.1.2017, p. 7 and EEA Supplement No 3, 12.1.2017, p. 7), e.i.f. 11.7.2015.


\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), c.i.f. 27.10.2018.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]

\[\text{Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), c.i.f. 15.6.2019.}\]


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\(^{[82]}\) Indent added by Decision No 140/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 16.7.2013.


\(^{[85]}\) Point inserted by Decision No 373/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

\(^{[86]}\) Indent added by Decision No 88/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.
The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) Notwithstanding the provisions of Protocol 1 to the Agreement, and unless otherwise provided for in this Agreement, the terms Member State(s) and competent authorities shall be understood to include, in addition to their meaning in the Regulation, the EFTA States and their competent authorities, respectively.

(b) The following provisions shall not apply to the EFTA States:

(i) The fourth subparagraph of Article 4(2);

(ii) Article 12; and,

(iii) Article 13(3).

(c) In the second subparagraph of Article 13(1), as regards the EFTA States, the words “or at the reporting intervals decided by the Conference of Parties to the Stockholm Convention” shall be inserted after the words “three years”.


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

Article 17 shall not apply to Liechtenstein.


biocidal products on the market of certain substances to be examined under the 10-year work programme referred to in Article 16(2) thereof (OJ L 216, 21.8.2007, p. 17).

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\[\text{Indent added by Decision No 143/2018 (OJ L 67, 25.2.2018, p. 34 and EEA Supplement No 13, EEA Annex, p. 35, e.g. 7.7.2018.}\]

\[\text{Indent added by Decision No 11/2019 (OJ L 63, 16.7.2020, p. 20 and EEA Supplement No 48, 16.7.2020, p. 20, e.g. 9.2.2019.}\]

\[\text{Indent added by Decision No 65/2019 (OJ L 210, 27.7.2020, p. 31 and EEA Supplement No 44, 27.7.2020, p. 31, e.g. 30.3.2019.}\]

\[\text{Indent added by Decision No 66/2019 (OJ L 210, 27.7.2020, p. 32 and EEA Supplement No 44, 27.7.2020, p. 32, e.g. 30.3.2019.}\]

\[\text{Indent added by Decision No 296/2019 (OJ L 68, 5.3.2020, p. 34 and EEA Supplement No 14, 5.3.2020, p. 34, e.g. 14.12.2019.}\]

\[\text{Indent added by Decision No 30/2020 (OJ L 57, 23.2.2023, p. 1 and EEA Supplement No 16, 23.2.2023, p. 1, e.g. 21.3.2020.}\]

\[\text{Indent added by Decision No 27/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 6.2.2021.}\]

\[\text{Indent added by Decision No 28/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 6.2.2021.}\]

\[\text{Indent added by Decision No 109/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 20.3.2021.}\]

\[\text{Indent added by Decision No 136/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 24.4.2021.}\]

\[\text{Indent added by Decision No 137/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 24.4.2021.}\]

\[\text{Indent added by Decision No 139/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 24.4.2021.}\]

\[\text{Indent added by Decision No 138/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 24.4.2021.}\]

\[\text{Indent added by Decision No 285/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 30.10.2021.}\]

\[\text{Indent added by Decision No 286/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.g. 30.10.2021.}\]
The transitional arrangements set out in the Annexes to the Act of Accession of 9 December 2011 for Croatia (Annex V, Chapter 10, Section VI) shall apply.

The Provisions of the Regulation shall, for the purpose of this Agreement, be read with the following adaptations:


(b) Notwithstanding the provisions of Protocol 1 to the Agreement, the term ‘Member State(s)’ contained in the Regulation shall be understood to include, in addition to its meaning in the Regulation, the EFTA States.

(c) As regards the EFTA States, the Agency shall, as and when appropriate, assist the EFTA Surveillance Authority or the Standing Committee, as the case may be, in the performance of their respective tasks.

(d) For products covered by Council Directive 91/414/EEC, the EFTA States will be free to limit access to their markets according to the requirements of their legislation existing at the date of entry into force of this Agreement. New EC rules will be dealt with according to the procedures laid down in Articles 97 to 104 of the Agreement.

(e) Should any disagreement between the contracting parties arise as to the administration of these provisions, Part VII of the Agreement shall apply *mutatis mutandis*.

(f) In Article 3, paragraph 10, the following shall be added at the end: “or into the territory of the EFTA States”.

(g) Article 64 (8) shall be read with the following adaptation:

“When the Commission takes authorisation decisions, the EFTA States will simultaneously and within 30 days of the Community Decision, take corresponding decisions. The EEA Joint Committee
shall be informed and shall periodically publish lists of such decisions in the EEA Supplement to the Official Journal.”

(h) The following paragraph shall be added in Article 79:

“4. The EFTA States shall participate fully in the Management Board and shall within it have the same rights and obligations as EC Member States, except for the right to vote. The internal rules of procedure of the Board shall give full effect to the EFTA States’ participation.”

(i) The following paragraph shall be added in Article 85:

“10. The EFTA States shall participate fully in the Member State Committee, in the Committee for Risk Assessment and the Committee for Socio-economic Analysis, and shall have the same rights and obligations as EC Member States, except for the right to vote. The internal rules of procedure of these committees shall give full effect to the EFTA States’ participation.”

(j) The following paragraph shall be added in Article 86:

“5. EFTA States shall participate in the Forum for Exchange of Information on Enforcement. The internal rules of procedure of this Forum shall give full effect to the EFTA States’ participation.’

(k) The following paragraph shall be added in Article 89:

“National of the EFTA States shall be eligible as members, or as alternates, of the Board of Appeal.”

(l) The following paragraph shall be added in Article 96:

“12. The EFTA States shall, as from the entry into force of this Decision, participate in the financing of the Agency. For this purpose the procedures laid down in Article 82(1)(a) and Protocol 32 of the Agreement shall apply mutatis mutandis.”

(m) The following shall be added in Article 102:

“The EFTA States shall grant privileges and immunities to the Agency equivalent to those contained in the Protocol on privileges and immunities of the European Communities.”

(n) The following paragraph shall be added to Article 103:

“By way of derogation from Article 12(2)(a) of the Conditions of employment of other servants of the European Communities, nationals of the EFTA States enjoying their full rights as citizens may be engaged under contract by the Executive Director of the Agency.”

(o) The following paragraph shall be added in Article 118:


(p) The following paragraph shall be added in Article 124:

“Liechtenstein shall not be obliged to establish a national helpdesk. Instead, Liechtenstein will publish a link to the helpdesk of the German Federal Institute for Occupational Safety and Health on the homepage of the competent Liechtenstein authority for chemicals, the Office for Environmental Protection.”


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[94] Indent added by Decision No 183/2014 (OJ L 202, 30.7.2015, p. 34 and EEA Supplement No 43, 30.7.2015, p. 34), e.i.f. 26.9.2014.


\[102\] Point and adaptation text inserted by Decision No 106/2012 (OJ L 270, 4.10.2012, p. 6 and EEA Supplement No 56, 4.10.2012, p. 8), e.i.f. 1.2.2013. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 3.2.2017.

\[103\] Indent and words “, as amended by:” added by Decision No 107/2012 (OJ L 270, 4.10.2012, p. 29 and EEA Supplement No 56, 4.10.2012, p. 28), e.i.f. 1.2.2013.

\[104\] Indent and words “, as amended by:” added by Decision No 107/2012 (OJ L 270, 4.10.2012, p. 29 and EEA Supplement No 56, 4.10.2012, p. 28), e.i.f. 1.2.2013.

\[105\] Indent added by Decision No 9/2013 (OJ L 144, 30.5.2013, p. 13 and EEA Supplement No 31, 30.5.2013, p.15), e.i.f. 2.2.2013.


\[109\] Indent added by Decision No 159/2014 (OJ L 15, 22.1.2015, p. 87 and EEA Supplement No 5, 22.1.2015, p. 10), e.i.f. pending: it shall apply from 9.7.2014.

\[110\] Indent added by Decision No 159/2014 (OJ L 15, 22.1.2015, p. 87 and EEA Supplement No 5, 22.1.2015, p. 10), e.i.f. pending: it shall apply from 9.7.2014.


[^623]: Indent added by Decision No 103/2020 (OJ L 172, 6.7.2023, p. 18 and EEA Supplement No 51, 6.7.2023, p. 18), e.i.f. 15.7.2020.

[^624]: Indent added by Decision No 199/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 12.12.2020. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 10.12.2021

The Provisions of Regulation (EC) No 1272/2008 shall, for the purpose of this Agreement, be read with the following adaptations:

(a) Liechtenstein shall not be obliged to establish a national helpdesk in the meaning of Article 44 of Regulation (EC) No 1272/2008. Instead, Liechtenstein will publish a link to the helpdesk of the German Federal Institute for Occupational Safety and Health on the homepage of the competent Liechtenstein authority for chemicals, the Office of Environmental Protection.

(b) The following provisions shall not apply to Norway:

(i) Article 51, in conjunction with Articles 4 and 46(1), with respect to the requirements for the classification, labelling and/or specific concentration limits for the substances or groups of substances listed in Part 3 of Annex VI to the Regulation (EC) No 1272/2008 and shown in the following list. Norway may require the use of different classification, labelling and/or specific concentration limits for this substance:

<table>
<thead>
<tr>
<th>Name</th>
<th>CAS No</th>
<th>Index No</th>
<th>EINECS</th>
</tr>
</thead>
<tbody>
<tr>
<td>acrylamide</td>
<td>79-06-1</td>
<td>616-003-00-0</td>
<td>201-173-7</td>
</tr>
</tbody>
</table>

(ii) Article 51, in conjunction with Articles 4 and 46(1), with respect to the requirements for the classification, labelling and/or specific concentration limits for the substances or group of substances not listed in Part 3 of Annex VI to the Regulation (EC) No 1272/2008 shown in the following list. Norway may require the use of different classification, labelling and/or specific concentration limits for these substances:

<table>
<thead>
<tr>
<th>Name</th>
<th>CAS No</th>
<th>Index No</th>
<th>ELINCS</th>
</tr>
</thead>
<tbody>
<tr>
<td>methyl acrylamidoglycolate</td>
<td>77402-05-2</td>
<td>[NOR-UNN-02-91]</td>
<td>403-230-3</td>
</tr>
<tr>
<td>(containing 0,01% ≤ acrylamide &lt; 0,1%)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>


(iii) Article 51, in conjunction with Articles 4, 9 and 46(1), with respect to mixtures containing substances as defined in adaptation text (i) and (ii) above.

(iv) These derogations shall elapse as from 1 June 2012 if by that date Norway does not pursue, in accordance with Article 37(1) of Regulation (EC) No 1272/2008, with the proposals for harmonised classification and labelling that were submitted to the European Chemical Agency on 1 June 2009 to support the more stringent classification and labelling.

If the procedure for harmonisation of classification and labelling foreseen in Article 37 of Regulation (EC) No 1272/2008 is pursued, a review of the derogations shall take place before 31 December 2013. If the derogations are supported by the outcome of that procedure, the derogations may be maintained by a Decision of the EEA Joint Committee. In the absence of such a Decision before 1 July 2014, the derogations shall elapse on that date.

(c) The Icelandic and Norwegian versions of the statements referred to in Articles 21 and 22 are set out in Appendices 5 and 6, respectively.


Point inserted by Decision No 139/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 16.7.2013.

Point inserted by Decision No 139/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 16.7.2013.

Point inserted by Decision No 194/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 9.11.2013.


Point inserted by Decision No 91/2015 (OJ L 211, 4.8.2016, p. 28 and EEA Supplement No 42, 4.8.2016, p. 27), e.l.f. 1.5.2015.

Point inserted by Decision No 92/2015 (OJ L 211, 4.8.2016, p. 29 and EEA Supplement No 42, 4.8.2016, p. 28), e.l.f. 1.5.2015.

Point inserted by Decision No 92/2015 (OJ L 211, 4.8.2016, p. 29 and EEA Supplement No 42, 4.8.2016, p. 28), e.l.f. 1.5.2015.


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\(^{[69]}\) Point inserted by Decision No 84/2016 (OJ L 300, 16.11.2017, p. 28 and EEA Supplement No 73, 16.11.2017, p. 32), e.i.f. 30.4.2016.

\(^{[70]}\) Point inserted by Decision No 84/2016 (OJ L 300, 16.11.2017, p. 28 and EEA Supplement No 73, 16.11.2017, p. 32), e.i.f. 30.4.2016.


\(^{[75]}\) Point inserted by Decision No 180/2016 (OJ L 80, 22.3.2018, p. 22 and EEA Supplement No 19, 22.3.2018, p. 29), e.i.f. 24.9.2016 and subsequently replaced by Decision No 87/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.


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12zzzzzp.[79] 32018 D 1477: Commission Implementing Decision (EU) 2018/1477 of 2 October 2018 on the terms and conditions of the authorisations of biocidal products containing ethyl butylacetylamino propionate referred by

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\textsuperscript{[2]} Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 13.6.2019.


\textsuperscript{[2]} Point inserted by Decision No 68/2019 (OJ L 210, 2.7.2020, p. 34 and EEA Supplement No 44, 2.7.2020, p. 39), e.i.f. 30.3.2019.

\textsuperscript{[2]} Point inserted by Decision No 68/2019 (OJ L 210, 2.7.2020, p. 34 and EEA Supplement No 44, 2.7.2020, p. 39), e.i.f. 30.3.2019.

\textsuperscript{[2]} Point inserted by Decision No 118/2019 (OJ L 279, 27.10.2022, p. 16 and EEA Supplement No 69, 27.10.2022, p. 16), e.i.f. 1.6.2019.

\textsuperscript{[2]} Point inserted by Decision No 154/2019 (OJ L 291, 10.11.2022, p. 33 and EEA Supplement No 74, 10.11.2022, p. 34), e.i.f. 15.6.2019.


\textsuperscript{[2]} Point inserted by Decision No 228/2019 (OJ L 4, 5.1.2023, p. 44 and EEA Supplement No 3, 5.1.2023, p. 44), e.i.f. 28.9.2019.

\textsuperscript{[2]} Point inserted by Decision No 228/2019 (OJ L 4, 5.1.2023, p. 44 and EEA Supplement No 3, 5.1.2023, p. 44), e.i.f. 28.9.2019.


\(^{(40)}\) Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\(^{(41)}\) Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\(^{(42)}\) Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.


\(^{(44)}\) Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.


\(^{(46)}\) Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\(^{(47)}\) Point inserted by Decision No 154/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.10.2020.

\(^{(48)}\) Point inserted by Decision No 29/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.


\textsuperscript{[40]} Point inserted by Decision No 30/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.

\textsuperscript{[41]} Point inserted by Decision No 30/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.

\textsuperscript{[42]} Point inserted by Decision No 30/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.

\textsuperscript{[43]} Point inserted by Decision No 111/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 20.3.2021.

\textsuperscript{[44]} Point inserted by Decision No 138/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.4.2021.

\textsuperscript{[45]} Point inserted by Decision No 138/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.4.2021.

\textsuperscript{[46]} Point inserted by Decision No 206/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

\textsuperscript{[47]} Point inserted by Decision No 206/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

\textsuperscript{[48]} Point inserted by Decision No 288/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[49]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[50]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[51]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.


\textsuperscript{[74]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[75]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[76]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[77]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[78]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[79]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[80]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[81]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[82]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[83]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[84]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

\textsuperscript{[85]} Point inserted by Decision No 289/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.


(77) Point inserted by Decision No 65/2022 (OJ L 182, 7.7.2022, p. 43 and EEA Supplement No 45, 7.7.2022, p. 31), c.f. 19.3.2022.
(78) Point inserted by Decision No 65/2022 (OJ L 182, 7.7.2022, p. 43 and EEA Supplement No 45, 7.7.2022, p. 31), c.f. 19.3.2022.
(80) Point inserted by Decision No 65/2022 (OJ L 182, 7.7.2022, p. 43 and EEA Supplement No 45, 7.7.2022, p. 31), c.f. 19.3.2022.
12zzzzzz.\cite{132022 D 0146} Commission Implementing Decision (EU) 2022/146 of 1 February 2022 determining whether a product containing Alkyl (C12-16) dimethylbenzyl ammonium chloride is a biocidal product, pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 24, 3.2.2022, p. 133).


12zzzzzzza.\cite{132022 D 1388} Commission Implementing Decision (EU) 2022/1388 of 23 June 2022 on the unresolved objections regarding the terms and conditions of the authorisation of the biocidal product Pat'Appât Souricide Canadien Foudroyant referred by France and Sweden in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 208, 10.8.2022, p. 7).


12zzzzzzzd.\cite{132022 D 0874} Commission Implementing Decision (EU) 2022/874 of 1 June 2022 on the terms and conditions of the authorisation of a biocidal product containing N-(trichloromethylthio)phthalimide (Folpet) referred by the Netherlands in accordance with Article 36(1) of Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 152, 3.6.2022, p. 187).


12zzzzzzzf.\cite{132022 D 1486} Commission Implementing Decision (EU) 2022/1486 of 7 September 2022 postponing the expiry date of the approval of acrolein for use in biocidal products of product-type 12 in accordance

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\(^{785}\) Point inserted by Decision No 279/2022 (OJ L 117, 4.5.2023, p. 4 and EEA Supplement No 35, 4.5.2023, p. 4), e.i.f. 29.10.2022.

\(^{786}\) Point inserted by Decision No 279/2022 (OJ L 117, 4.5.2023, p. 4 and EEA Supplement No 35, 4.5.2023, p. 4), e.i.f. 29.10.2022.

\(^{787}\) Point inserted by Decision No 279/2022 (OJ L 117, 4.5.2023, p. 4 and EEA Supplement No 35, 4.5.2023, p. 4), e.i.f. 29.10.2022.


\(^{790}\) Point inserted by Decision No 10/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.

\(^{791}\) Point inserted by Decision No 10/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.

\(^{792}\) Point inserted by Decision No 10/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.

\(^{793}\) Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.

\(^{794}\) Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.


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[^79]: Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.F. 29.4.2023.
[^78]: Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.F. 29.4.2023.
[^77]: Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.F. 29.4.2023.
[^76]: Point inserted by Decision No 84/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.F. 29.4.2023.


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1. **Point inserted by Decision No 86/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
2. **Point inserted by Decision No 86/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
3. **Point inserted by Decision No 87/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
4. **Point inserted by Decision No 87/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
5. **Point inserted by Decision No 87/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
6. **Point inserted by Decision No 87/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 29.4.2023.**
7. **Point inserted by Decision No 149/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 14.6.2023.**
8. **Point inserted by Decision No 180/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.l.f. 6.7.2023.**
9. **Point inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.l.f. 1.6.2015.**
The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The EFTA States shall be free to limit access to their markets of plant protection products containing active substances approved in accordance with Council Directive 91/414/EEC or the Transitional measures in Article 80 of Regulation (EC) No 1107/2009.

(b) The EFTA States, with the exception of Liechtenstein, may be ‘rapporteur Member State’ and ‘co-rapporteur’.

(c) The following shall be added to Article 18:

“The allocation of evaluation of active substances to an EFTA State according to Article 18(f) is subject to consent from that State.”

(d) The following shall be added to Articles 37(4) and 42(2):

“For the EFTA States the time limit of 120 days shall at the earliest run from the date when the Act of approval of the active substances contained in the plant protection product is incorporated into the present Agreement.”

(e) The following shall be added to Article 47(3):

“For the EFTA States the time limit of 120 days shall at the earliest run from the date when the Act of approval of the active substances contained in the low-risk plant protection product is incorporated into the present Agreement.”

\[825\] Indent and words “, as amended by:” added by Decision No 18/2015 (OJ L 93, 7.4.2016, p. 30 and EEA Supplement No 21, 7.4.2016, p. 26), e.i.f. pending. It shall apply provisionally pending the entry into force of the 2014 EEA Enlargement Agreement, from the day of entry into force of Decision of the Joint Committee No 203/2014 which e.i.f. 1.6.2015.


\[817\] Indent added by Decision No 183/2018 (OJ L 75, 4.3.2021, p. 18 and EEA Supplement No 15, 4.3.2021, p. 17), e.i.f. 22.9.2018.

\[818\] Indent added by Decision No 210/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.3.2020 and subsequently corrected before publication by Corrigendum of 28.10.2022.

\[819\] Indent added by Decision No 234/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

\[820\] Indent added by Decision No 340/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

\[821\] Indent added by Decision No 89/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.

\[822\] Indent added by Decision No 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
(f) The following shall be added to Article 48:

“The EFTA States may limit access to their markets of plant protection products containing genetically modified organisms, when measures to restrict or prohibit those organisms have been taken according to Article 23 of Directive 2001/18/EC, as adapted by this Agreement.”

(g) Article 49 shall not apply to Liechtenstein.

(h) Article 80(6) shall be replaced by the following:

“Plant protection products authorized in accordance with national provisions applicable at the time of authorization may continue to be placed on the market until the plant protection product has been risk assessed according to Regulation (EU) No 1107/2009.”

(i)[82] In Article 80(8), as regards the EFTA States, the words “15 July 2019” shall read “the date of entry into force of Decision of the EEA Joint Committee No 77/2023 of 28 April 2023.”

(j) The following shall be added to “Zone A – North” in Annex I:

“Iceland, Norway”

(k) The following shall be added to “Zone B – Centre” in Annex I:

“Liechtenstein”


[82] Adaptations (i) to (j) are renumbered as adaptations (j) to (k) and adaptation text (i) inserted by Decision 77/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.I.f. pending


32012 R 0087: Commission Implementing Regulation (EU) No 87/2012 of 1 February 2012 (OJ L 30, 2.2.2012, p. 8),


\[\text{Indent added by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015 and subsequently corrected before publication by Corrigendum of 20.3.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]

\[\text{Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.}\]


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-[**] 32013 R 0017: Commission Implementing Regulation (EU) No 17/2013 of 14 January 2013 (OJ L 9, 15.1.2013, p. 5);


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**[641]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[642]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[643]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015 and subsequently deleted by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.9.2022.

**[644]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[645]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[646]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[647]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[648]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[649]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[650]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[651]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[652]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[653]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.

**[654]** Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.i.f. 1.6.2015.


(*76) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*77) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*78) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.


(*80) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*81) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*82) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*83) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*84) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*85) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*86) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*87) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*88) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

(*89) Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.


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- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.

- [**] Indent added by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), e.l.f. 1.6.2015.


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\(^{(22)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2015.

\(^{(23)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(24)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(25)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(26)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(27)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(28)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(29)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

\(^{(30)}\)
Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

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\(^{(33)}\)

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\(^{(35)}\)


32015 R 0543: Commission Implementing Regulation (EU) 2015/543 of 1 April 2015 (OJ L 90, 2.4.2015, p. 1),

32015 R 0553: Commission Implementing Regulation (EU) 2015/553 of 7 April 2015 (OJ L 92, 8.4.2015, p. 86),

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\textsuperscript{(906)} Indent added by Decision No 183/2015 (OJ L 8, 12.1.2017, p. 12 and EEA Supplement No 3, 12.1.2017, p. 11), e.i.f. 11.7.2015.
5.2.2016, p. 5

5.3.2018, p. 2

5.4.2016, p. 25

5.5.2016, p. 32


\[1058\] Indent added by Decision No 54/2018 (OJ L 26, 30.1.2020, p. 35 and EEA Supplement No 6, 30.1.2020, p. 28), c.i.f. 24.3.2018.


\[1061\] Indent added by Decision No 54/2018 (OJ L 26, 30.1.2020, p. 35 and EEA Supplement No 6, 30.1.2020, p. 28), c.i.f. 24.3.2018.


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Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.

Indent added by Decision No 69/2019 (OJ L 210, 2.7.2020, p. 36 and EEA Supplement No 44, 2.7.2020, p. 41), e.i.f. 30.3.2019.


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\[1104\] Indent added by Decision No 121/2019 (OJ L 279, 27.10.2022, p. 23 and EEA Supplement No 69, 27.10.2022, p. 23), e.i.f. 1.6.2019.

\[1105\] Indent added by Decision No 121/2019 (OJ L 279, 27.10.2022, p. 23 and EEA Supplement No 69, 27.10.2022, p. 23), e.i.f. 1.6.2019.

\[1106\] Indent added by Decision No 121/2019 (OJ L 279, 27.10.2022, p. 23 and EEA Supplement No 69, 27.10.2022, p. 23), e.i.f. 1.6.2019.

\[1107\] Indent added by Decision No 121/2019 (OJ L 279, 27.10.2022, p. 23 and EEA Supplement No 69, 27.10.2022, p. 23), e.i.f. 1.6.2019.

\[1108\] Indent added by Decision No 121/2019 (OJ L 279, 27.10.2022, p. 23 and EEA Supplement No 69, 27.10.2022, p. 23), e.i.f. 1.6.2019.


\[1110\] Indent added by Decision No 155/2019 (OJ L 291, 10.11.2022, p. 34 and EEA Supplement No 74, 10.11.2022, p. 35), e.i.f. 15.6.2019.


The document contains a list of Commission Implementing Regulations (EU) from various dates and pages. The regulations are cited in a natural text format, with page numbers and dates of publication. The text is indented and formatted to resemble a typical footnote or reference list.


\[\text{(\textsuperscript{1131}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1132}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1133}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1134}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1135}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1136}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1137}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1138}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1139}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1140}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1141}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1142}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1143}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1144}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]

\[\text{(\textsuperscript{1145}) Indent added by Decision No 33\textsuperscript{a}2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.}\]
-\footnote{3202 R 0616: Commission Implementing Regulation (EU) 2020/616 of 5 May 2020 (OJ L 143, 6.5.2020, p. 1).}

-\footnote{3202 R 0617: Commission Implementing Regulation (EU) 2020/617 of 5 May 2020 (OJ L 143, 6.5.2020, p. 6).}


-\footnote{3202 R 0646: Commission Implementing Regulation (EU) 2020/646 of 13 May 2020 (OJ L 151, 14.5.2020, p. 3).}


-\footnote{3202 R 1004: Commission Implementing Regulation (EU) 2020/1004 of 9 July 2020 (OJ L 221, 10.7.2020, p. 133).}


\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}

\footnote{\footnote{Indent added by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.}}


(1174) Indent added by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021.

(1177) Indent added by Decision No 292/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

(1179) Indent added by Decision No 293/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

(1181) Indent added by Decision No 293/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 30.10.2021.

(1188) Indent added by Decision No 340/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1182) Indent added by Decision No 340/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1183) Indent added by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1184) Indent added by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1185) Indent added by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1186) Indent added by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1187) Indent added by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1188) Indent added by Decision No 343/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.

(1189) Indent added by Decision No 672/2022 (OJ L 182, 7.7.2022, p. 46 and EEA Supplement No 45, 7.7.2022, p. 34), e.i.f. 19.3.2022.

(1190) Indent added by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), e.i.f. 30.4.2022.


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Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 73 and EEA Supplement No 61, 22.9.2022, p. 72), e.i.f. 30.4.2022.

Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 77 and EEA Supplement No 61, 22.9.2022, p. 76), e.i.f. 30.4.2022.


32022 R 0698: Commission Implementing Regulation (EU) 2022/698 of 3 May 2022 (OJ L 130, 4.5.2022, p. 3),

32022 R 0800: Commission Implementing Regulation (EU) 2022/800 of 20 May 2022 (OJ L 143, 23.5.2022, p. 4),


32022 R 0814: Commission Implementing Regulation (EU) 2022/814 of 20 May 2022 (OJ L 146, 25.5.2022, p. 6),


32022 R 1480: Commission Implementing Regulation (EU) 2022/1480 of 7 September 2022 (OJ L 233, 8.9.2022, p. 43),

32022 R 0378: Commission Implementing Regulation (EU) 2022/378 of 4 March 2022 (OJ L 72, 7.3.2022, p. 2),


32022 R 0437: Commission Implementing Regulation (EU) 2022/437 of 16 March 2022 (OJ L 89, 17.3.2022, p. 3),


32022 R 0686: Commission Implementing Regulation (EU) 2022/686 of 28 April 2022 (OJ L 126, 29.4.2022, p. 18),


\[1207\] Indent added by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1208\] Indent added by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1209\] Indent added by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1210\] Indent added by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023 and subsequently corrected before publication by Corrigendum of 17.3.2023.

\[1211\] Indent added by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1212\] Indent added by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1213\] Indent added by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 4.2.2023.

\[1214\] Indent added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1215\] Indent added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1216\] Indent added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1217\] Indent added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1218\] Indent added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1219\] Indent added by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.

\[1220\] Indent added by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 29.4.2023.
The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

The EFTA States shall be free to limit access to their markets of plant protection products containing active substances approved in accordance with Council Directive 91/414/EEC or the Transitional measures in Regulation (EC) No 1107/2009 Article 80.


[1221] Indent added by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.
[1222] Indent added by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.
[1223] Indent added by Decision No 92/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The following shall be added to the list under the title “RSh 1” in point 1.1. of Annex II:

“IS: Eftirð í snertingu við augu.

NO: Giftig ved øyekontakt.”

(b) The following shall be added to the list under the title “RSh 2” in point 1.1. of Annex II:

“IS: Getur valdið ljósnæmingu.

NO: Kan gi overfølsomhet for sollys/UV-stråling.”

(c) The following shall be added to the list under the title “RSh 3” in point 1.1. of Annex II:

“IS: Efnið brennr húð og augu í snertingu við gufu og veldur kali í snertingu við vökva.

NO: Kontakt med damp virker etsende på hud og øyne, og kontakt med væske gir frostskade.”

(d) The following shall be added to the list in point 1 of Annex III:

\(\text{125}\) Indent added by Decision No 89/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.


\(\text{125}\) Indent added by Decision No 89/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.

\(\text{125}\) Point inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.i.f. 1.6.2015.

\(\text{125}\) Indent and words “as amended by:” added by Decision No 186/2018 (OJ L 75, 4.3.2021, p. 22 and EEA Supplement No 15, 4.3.2021, p. 21), e.i.f. 22.9.2018.

\(\text{125}\) Indent added by Decision No 89/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.

\(\text{125}\) Point and adaptation text inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.i.f. 1.6.2015.
"IS: Mengið ekki vatn með efninu eða íflátí þess. (Hreinsís ekki búnað nálgött yfirborðsvatn/Koma skal í veg fyrir að mengun verði með afremsli frá hærjarlóðum og vegum.)

NO: Úngag forurensning av vannmiljøet med produktet eller emballasjen. (Ikke rengjør spredet støy nær overflatevann/úngag forurensning via avrenning fra gjørsplasser og veier)."

(e) The following shall be added to the list under the title “SPo 1” under the title “Specific Provisions” in point 2.1 of Annex III:

"IS: Ef efnið kemst í snertingu við háð skal fyrst hreinsa það af með þurrum klút og skola síðan húðina með miklu vatni.

NO: Etter kontakt med huden, fjern først produktet med en tørr klut, og vask deretter med mye vann.”

(f) The following shall be added to the list under the title “SPo 2” under the title “Specific Provisions” in point 2.1 of Annex III:

"IS: Þvoið Allan hlífðarfatnað að lokinni notkun.

NO: Vask alt personleg verneutstyr etter bruk.”

(g) The following shall be added to the list under the title “SPo 3” under the title “Specific Provisions” in point 2.1 of Annex III:

"IS: Fordi lið innöndun reyks eftir að kveikt hefur verið í efninu og yfirgefið þegar í stað svæðið sem er til meðhöndunar.

NO: Pust ikke inn røyken etter at produktet har antent, og forlat det behandlede området sikkertlig.”

(h) The following shall be added to the list under the title “SPo 4” under the title “Specific Provisions” in point 2.1 of Annex III:

"IS: Opna skal íflátí utanhús og við þurr skilyði.

NO: Beholderen skal åpnes utenårs og under tørrte forhold.”

(i) The following shall be added to the list under the title “SPo 5” under the title “Specific Provisions” in point 2.1 of Annex III:

"IS: Loftræsta skal úðuð svæði/gröðurfür (vandlega/eða í tilgreindan tíma/þar til úðinn hefur þornað) áður en farið er þangað inn aftur.

NO: De behandlede områder/veksthus ventileres (grundig/eller avgivelse av tid/inntil produktet har tøkt) før man oppholder seg der igjen.”

(j) The following shall be added to the list under the title “SPe 1” in point 2.2 of Annex III:

"IS: Til að verna grunnvatn/jarðvegslíflverur skal ekki nota þetta eða annað efní sem inniheldur (tilgreinið virkt efní eða flokk virkra efní eftir þvi sem við á) lengur eða oftar en (tilgreinið hversu lengi eða oft má nota efní).

NO: For å beskytte (grunnvannet/jordlevende organiser) må dette produktet eller andre produkter som inneholder (angi navnet på virksomt stoff eller gruppe av virksomme stoffer) kun brukes/ikke brukes mer enn (angi tidsperiode eller antall behandlinger).”

(k) The following shall be added to the list under the title “SPe 2” in point 2.2 of Annex III:

"IS: Til að verna grunnvatn/vatnalíflverur skal ekki nota þetta efní (á tilgreindar jarðvegsgerð eða við tilgreinandar aðstæður).

NO: For å beskytte (grunnvannet/vannlevende organiser) må dette produktet ikke brukes (på beskrevet jordtype eller under beskrevne forhold).”
(l) The following shall be added to the list under the title “Sp e 3” in point 2.2 of Annex III:

“IS: Til að vernda vatnafíverur/plóntur utan markhóps/líliaur utan markhóps/skordýr má ekki nota efnið nær óræðuleitlandi/yfirförðsvatni en (tilgreind breidd sveðís sem er óheimilt að úða).

NO: For å beskytte (vannlevende organiser/viltlevende planter/insekter/ledýr) må dette produktet ikke brukes nærmere enn (angi avstand) fra (overflatevann/kantvegetasjon).”

(m) The following shall be added to the list under the title “Sp e 4” in point 2.2 of Annex III:

“IS: Til að vernda vatnafíverur/plóntur utan markhóps má ekki nota efnið á malbikað, steinsteyp, hellulagðt eða malarborð yfirborð eða vegi (jarnbrautarþor) eða önnur sveði þar sem hett er við afrennsli út í umhverfið.

NO: For å beskytte (vannlevende organiser/viltlevende planter) må dette produktet ikke brukes på harde overflater som asfaltere, betong- brostein- eller gruslagte områder og veier/jernbaner, eller på andre områder med stor risiko for avrenning.”

(n) The following shall be added to the list under the title “Sp e 5” in point 2.2 of Annex III:

“IS: Til að vernda fugla/víllt spendýr verður að geta þess vandlega að efnið sé algerlega hulið jarðvegi; getið þess serstaklega að efnið sé hulið í endum ræða.

NO: For å beskytte (fugler/ville pattedyr) skal produktet innblandes i jorden. Sørg også for at produktet er helt innblanded i enden av radene.”

(o) The following shall be added to the list under the title “Sp e 6” in point 2.2 of Annex III:

“IS: Hreinsíð upp allt efní, sem hefur farið til spillis, til að vernda fugla/víllt spendýr.

NO: For å beskytte (fugler/ville pattedyr) skal alt søl fjernes.”

(p) The following shall be added to the list under the title “Sp e 7” in point 2.2 of Annex III:

“IS: Óheimilt er að nota efnið á varþíma fugla.

NO: Må ikke brukes i fuglenes hekketíð.”

(q) The following shall be added to the list under the title “Sp e 8” in point 2.2 of Annex III:

“IS: Hættulegt frævandi skordýrun/Til að vernda hýflugur og önnur frævandi skordýr er óheimilt að nota efnið á blómstrandi nýtalóntúr/Óheimilt er að nota efnið þar sem hýflugur eru í fæuletíð/fjarðlegi hýkýpur meðan meðhöðlundi með efnini fer fram eða hylji þær á meðan og í (tilgreindi tíma) að lokini meðhöðlundi/Óheimilt er að nota efnið ef blómstrandi ilgrið er til staðar/Eyða skal ilgrið áður en það blómgaði/Óheimilt er að nota efnið fyrir (tilgreindi tíma).

NO: Farlig for bier./For å beskytte hier og andre pollinerende insekter må dette produkt ikke brukes mens kulturen blomstrer./Må ikke brukes der biene søker næring./Dekk til eller flytt bikuber i behandlingsperioden og i (nevnt antall timer/dager) etter behandlingen./Må ikke brukes i nærheten av blomstrende uggress./Fjern uggresset før det blomstrer./Må ikke brukes før (tidspunkt).”

(r) The following shall be added to the list in point 2.3 of Annex III:

“IS: Til að koma í veg fyrir þolmyndun skal ekki nota þetta eða annað varmarefní sem inniheldur (tilgreindi virkt efní eða flogk virkra efní eftir því sem við á) oftar eða lengur en (tilgreinið hversu oft eða lengi má nota efnið).

NO: For å unngå utvikling av resistens må dette produkt eller andre produkter som inneholder (angi virksomt stoff eller gruppe av virksomme stoffer) kun brukes/ikke brukes mer enn (i tidsperioden eller antall ganger).”

(s) The following shall be added to the list under the title “SPr l” in point 2.4 of Annex III:
“IS: Beitu skal komið fyrrir þannig að ekki sé hættu á að önnur dýr komist í hana. Festa skal beituna tryggilega þannig að nagdýr geti ekki dregið hana í burtu.

NO: Produkten skal plASSES på en slik måte at risikoen for at andre dyr kan innta produktet minimeres. Pass på at produkt i blokkform ikke kan flyttes vegg av de gnagere som skal bekjempes.”

The following shall be added to the list under the title “SPr 2” in point 2.4 of Annex III:

“IS: Auðkennið svæðið, sem meðhöndl the stendur. Varað skal við hættunni á að verða fyrir eitrun (beinni eða öbenni) af völum storkuvarans og tilgreina skal móteitrið við honum.

NO: Det behandlede området skal merkes i behandlingsperioden. Faren for forgiftning (primær eller sekundær) ved inntak av antikoaguleringsmidler, samt motgift, skal angis på oppslag.”

The following shall be added to the list under the title “SPr 3” in point 2.4 of Annex III:


NO: Døde gnagere skal fjernes fra behandlingsområdet hver dag. Døde gnagere må ikke plasseres i åpne avfallsbeholdere.”


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(1239) Point inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), c.f.f. 1.6.2015 and subsequently replaced by Decision No 341/2021 (OJ L, to be published) and EEA Supplement No [to be published], c.f.f. pending.


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Point inserted by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), c.f.f. 1.6.2015 and subsequently corrected before publication by Corrigendum of 20.3.2015, subsequently corrected before publication by Corrigendum of 10.7.2015.

Point inserted by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. 1.6.2015.

Point inserted by Decision No 204/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. 1.6.2015.


13zr.\[1278\] 32012 D 0677: Commission Implementing Decision 2012/677/EU of 30 October 2012 allowing Member States to extend provisional authorisations granted for the new active substances ametocadbacin (initially applied for under the development code BAS 650 F) and disodium phosphate (OJ L 305, 1.11.2012, p. 27).


\[1274\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1275\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1276\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1277\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1278\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.


\[1280\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1281\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1282\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.

\[1283\] Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. l.6.2015.


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13zzze.\(^{[18]}\)2012 R 0686: Commission Implementing Regulation (EU) No 686/2012 of 26 July 2012 allocating to Member States, for the purposes of the renewal procedure, the evaluation of the active substances whose approval expires by 31 December 2018 at the latest (OJ L 200, 27.7.2012, p. 5), as amended by:


\(^{[134]}\) Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. 1.6.2015.

\(^{[135]}\) Point inserted by Decision No 205/2014 (OJ L 202, 30.7.2015, p. 73 and EEA Supplement No 43, 30.7.2015, p. 72), c.f.f. 1.6.2015.


\(^{[143]}\) Indent added by Decision No 144/2017 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 7.7.2018.


32013 D 0205: Commission Implementing Decision 2013/205/EU of 25 April 2013 allowing Member States to extend provisional authorisations granted for the new active substances acequinocyl, aminopyralid, ascorbic acid, fluibendiamide, gamma-cyhalothrin, ipconazole, metalumizone, orthosulfuron, pseudomonas sp. strain DSMZ 13134, pyridalil, pyroxasulam, spiroimesifen, thiencarbazone and topramezone (OJ L 117, 27.4.2013, p. 20).

32013 R 0485: Commission Implementing Regulation (EU) No 485/2013 of 24 May 2013 amending Implementing Regulation (EU) No 540/2011, as regards the conditions of approval of the active substances clothianidin, thiamethoxam and imidacloprid, and prohibiting the use and sale of seeds treated with plant protection products containing those active substances (OJ L 139, 25.5.2013, p. 12), as amended by:


32014 R 0108: Commission Implementing Regulation (EU) No 108/2014 of 5 February 2014 concerning the non-approval of the active substance potassium thiocyanate, in accordance with Regulation (EC) No


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\[^{1339}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1340}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2015.

\[^{1341}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1342}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1343}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1344}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1345}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1346}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.

\[^{1347}\] Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), c.i.f. 1.6.2015.


\[1345\]Indent and words “, as amended by” added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.


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Corrigendum to the EUT act subsequently taken note of by the EEA Joint Committee on 10.6.2022.


[1575] Indent added by Decision No 33/2021 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 6.2.2021.


\[1378\] Indent added by Decision No 12/2023 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 4.2.2023.


\[1381\] Point inserted by Decision No 264/2015 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 1.11.2015.

\[1382\] Point inserted by Decision No 265/2015 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 1.11.2015.

\[1383\] Point inserted by Decision No 264/2015 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 1.11.2015.

\[1384\] Point inserted by Decision No 264/2015 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 1.11.2015.


[1394] Indent and words “*, as amended by:*” added by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.


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\(^{1423}\) Point inserted by Decision No 30/2017 (OJ L 297, 22.11.2018, p. 38 and EEA Supplement No 78, 22.11.2018, p. 44), e.i.f. 4.2.2017.


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[1441] Indent and words “., as amended by:” added by Decision No 293/2021 (OJ L [to be published]) and EEA Supplement No [to be published], c.f.f. 30.10.2021.


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**Notes:**


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1507 Point inserted by Decision No 155/2019 (OJ L 291, 10.11.2022, p. 34 and EEA Supplement No 74, 10.11.2022, p. 35), e.i.f. 15.6.2019.


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1536 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1537 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1538 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1539 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1540 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.


1541 Point inserted by Decision No 31/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1542 Point inserted by Decision No 32/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1543 Point inserted by Decision No 32/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1544 Point inserted by Decision No 32/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1545 Point inserted by Decision No 32/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1546 Point inserted by Decision No 32/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1547 Point inserted by Decision No 33/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1548 Point inserted by Decision No 33/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
1549 Point inserted by Decision No 33/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.


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[1558] Point inserted by Decision No 33/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
[1559] Point inserted by Decision No 139/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.4.2021.
[1560] Point inserted by Decision No 140/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.4.2021.
[1561] Point inserted by Decision No 208/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.
[1562] Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021.
[1563] Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021.
[1564] Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021.
[1565] Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 25.9.2021.
13zzzzzzzzz.\(^{(158\text{a})}\)


13zzzzzzzzz.\(^{(158b)}\)


13zzzzzzzzz.\(^{(158c)}\)


13zzzzzzzzz.\(^{(158d)}\)


13zzzzzzzzz.\(^{(158e)}\)


13zzzzzzzzz.\(^{(158f)}\)


13zzzzzzzzz.\(^{(158g)}\)


13zzzzzzzzz.\(^{(158h)}\)


13zzzzzzzzz.\(^{(158i)}\)


13zzzzzzzzz.\(^{(158j)}\)

**2021 R 0843:** Commission Implementing Regulation (EU) 2021/843 of 26 May 2021 renewing the approval of the active substance cyazofamid, in accordance with Regulation (EC) No 1107/2009 of

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\(^{(158a)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158b)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158c)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158d)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158e)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158f)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158g)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158h)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158i)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.

\(^{(158j)}\) Point inserted by Decision No 239/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 25.9.2021.


13zzzzzzzzz.\(^{(1571)}\)32021 R 0428: Commission Implementing Regulation (EU) 2021/428 of 10 March 2021 adopting standard data formats for the submission of applications for the approval or the amendment to the conditions of approval of active substances, as provided for in Regulation (EC) No 1107/2009 of the European Parliament and of the Council (OJ L 84, 11.3.2021, p. 25).


\(^{(1568)}\) Point inserted by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 11.12.2021.

\(^{(1569)}\) Point inserted by Decision No 342/2021 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 11.12.2021.

\(^{(1570)}\) Point inserted by Decision No 343/2021 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 11.12.2021.

\(^{(1571)}\) Point inserted by Decision No 67/2022 (OJ L 182, 7.7.2022, p. 46 and EEA Supplement No 45, 7.7.2022, p. 34), c.i.f. 19.3.2022.

\(^{(1572)}\) Point inserted by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), c.i.f. 30.4.2022.

\(^{(1573)}\) Point inserted by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), c.i.f. 30.4.2022.

\(^{(1574)}\) Point inserted by Decision No 127/2022 (OJ L 246, 22.9.2022, p. 73 and EEA Supplement No 61, 22.9.2022, p. 72), c.i.f. 30.4.2022.

\(^{(1575)}\) Point inserted by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), c.i.f. 30.4.2022.

\(^{(1576)}\) Point inserted by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), c.i.f. 30.4.2022.


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**Point inserted by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.**

**Point inserted by Decision No 11/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.**

**Point inserted by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.**

**Point inserted by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.**

**Point inserted by Decision No 12/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.**

**Point inserted by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.**

**Point inserted by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.**

**Point inserted by Decision No 90/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.**

**Point inserted by Decision No 91/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.**


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1595 Point inserted by Decision No 91/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.
1596 Point inserted by Decision No 92/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 29.4.2023.
ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

The Contracting Parties take note of the content of the following acts:

1.  \{1611\}


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13.\[1623\] 32004 H 0394: Commission Recommendation 2004/394/EC of 29 April 2004 on the results of the risk evaluation and the risk reduction strategies for the substances: Acetonitrile; Acrylamide; Acrylonitrile; Acrylic acid; Butadiene; Hydrogen fluoride; Hydrogen peroxide; Methacrylic acid; Methyl methacrylate; Toluene; Trichlorobenzene (OJ L 144, 30.4.2004, p. 72), as corrected by OJ L 199, 7.6.2004, p. 41.

14.\[1624\] 32006 H 0283: Commission Recommendation of 11 April 2006 on risk reduction measures for the substances: Dibutylphthalate; 3,4-Dichloroaniline; Di-isodecyl phthalate; 1,2-Benzenedicarboxylic acid, di-C9,11-branched alkyl esters, C10-rich; Di-isononyl phthalate; 1,2-Benzenedicarboxylic acid, di-C8,10-branched alkyl esters, C9-rich; Ethylenediaminetetraacetate; Methyl acetate; Monochloroacetic acid; n-Pentane; Tetrasodium ethylenediaminetetraacetate (OJ L 104, 13.4.2006, p. 45).


XVI. COSMETICS

ACTS REFERRED TO

1. [ ]


p. 4),


[1452] Indent added by Decision No 123/2019 (OJ L 279, 27.10.2022, p. 28 and EEA Supplement No 69, 27.10.2022, p. 28), e.i.f. 1.6.2019.


32022 R 1176: Commission Regulation (EU) 2022/1176 of 7 July 2022 (OJ L 183, 8.7.2022, p. 51),
32022 R 1181: Commission Regulation (EU) 2022/1181 of 8 July 2022 (OJ L 184, 11.7.2022, p. 3),

[1662] Indent added by Decision No 34/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
[1663] Indent added by Decision No 34/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
[1664] Indent added by Decision No 34/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
[1665] Indent added by Decision No 294/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.
[1666] Indent added by Decision No 295/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.12.2022.
[1668] Indent added by Decision No 130/2022 (OJ L 246, 22.9.2022, p. 79 and EEA Supplement No 61, 22.9.2022, p. 78), e.i.f. 30.4.2022.


9. [ ] (1678)

10. [ ] (1679)

11. (1680) [ ] (1681)
12. \[1682\]  [ ]\[1682\]

13.\[1684\]  [ ]\[1684\]


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**\[1687\]

The Contracting Parties take note of the content of the following acts:


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XVII. ENVIRONMENT PROTECTION

ACTS REFERRED TO

1. [ ]

2. [ ]

3. [ ]

4. [ ]


6. [ ]


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The provisions of the Directive shall, for the purposes of the present Agreement, be read with the following adaptions:

(a) In point 4 (outermost regions) of Article 2, the words “Iceland, with regard to all of its territory,” shall be inserted after the word “departments.”.

(b) In point 1 of Article 6, the words “Article 95(10) of the Treaty” shall be replaced by reference to “Article 75 of the Agreement”.

(c) In Article 2(5), the word “Iceland” shall be added after the word “Finland” and the word “Norway” shall be added after the word “Lithuania”.

(d) In Article 3(4), the following subparagraph shall be added after the first subparagraph:

“...Iceland may permit the placing on the market, during the summer period, of petrol containing ethanol or methanol with maximum vapour pressure of 70 kPa, on condition that the ethanol used is a biofuel or that the greenhouse gas emission saving from the use of methanol fulfils the criteria specified in Article 7b(2).”

(e) Articles 7a to 7e shall not apply to Liechtenstein.

(f) Article 7b(6) shall not apply to the EFTA States.

6aa. [ ]

6ab. [ ]

6ac. [ ]

6ad. [ ]


6af. [ ]

6ag. [ ]


(1) Listed here for information purposes only; for application see Annex IV on Energy.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

By way of derogation to paragraph 8 of Protocol 1 to the EEA Agreement, Article 2 shall not apply to grasslands situated within the territories of the EFTA States.


The provisions of the Directive, shall for the purposes of this Agreement be read with the following adaptation:

References to other acts in the Directive shall apply to the extent and in the form that those acts are incorporated into this Agreement.


The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for the Czech Republic (Annex V, Chapter 7, Section A), Cyprus (Annex VII, Chapter 9, Section B), Latvia (Annex VIII, Chapter 10, Section B, point 2), Lithuania (Annex IX, Chapter 10, Section B), Hungary (Annex X, Chapter 8, Section A, point 2), Malta (Annex XI, Chapter 10, Section B, point 2), Poland (Annex XII, Chapter 13, Section B, point 2), Slovenia (Annex XIII, Chapter 9, Section A) and Slovakia (Annex XIV, Chapter 9, Section B, point 2) shall apply.

The transitional arrangements set out in the Annexes to the Act of Accession of 25 April 2005 for Bulgaria (Annex VI, Chapter 10, Section B, point 2) and Romania (Annex VII, Chapter 9, Section B, point 2), shall apply.

The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptations:

In Article 6(7), the words “, Iceland” shall be inserted after the word “Ireland” and “, the presence of rural areas and low population density” shall be inserted after the word “areas”.


Footnotes:


1742 Indent added by Decision No 296/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 30.10.2021.


[1754] The provisions of the Decision shall, for the purposes of this Agreement, be read with the following adaptation:

For the purposes of Article 6c and Annex II, Liechtenstein shall use an equivalent method to determine the weight of the municipal waste recycled.


[1759] The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for Estonia (Annex VI, Chapter 9, Section A), Latvia (Annex VIII, Chapter 10, Section A), Lithuania (Annex IX, Chapter 10, Section A), Malta (Annex XI, Chapter 10, Section A), Poland (Annex XII, Chapter 13, Section A, point 1) and Slovakia (Annex XIV, Chapter 9, Section A) shall apply.

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[1752] Indent and words ‘‘, as amended by:’’ added by Decision No 155/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.10.2020.


The transitional arrangements set out in the Annexes to the Act of Accession of 25 April 2005 for Bulgaria (Annex VI, Chapter 10, Section A, point 1) and Romania (Annex VII, Chapter 9, Section A), shall apply.

The provisions of the Directive shall, for the purposes of the present Agreement, be read with the following adaptation:

Paragraph 4 of Annex I shall not apply to existing terminals in Iceland with a throughput of less than 5,000 metric tons per year, which are serviced by ships.

9. [1764]


The provisions of the Directive shall, for the purposes of the present Agreement, be read with the following adaptation:

In Article 3(4), the words “, and for Iceland for a period of 36 months following the latter date in Annex II, to finish the Icelandic stocks” shall be inserted after the word “force”.

9a. [ ] [1764]

9aa. [ ] [1764]

9b. [1766]


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) In Article 4(2), as regards the EFTA States, the words “31 December 2016” shall read “one year after the date of entry into force of Decision of the EEA Joint Committee No 160/2019 of 14 June 2019”.

(b) In Article 5(2), as regards the EFTA States, the words “1 January 2017” shall read “one year after the date of entry into force of Decision of the EEA Joint Committee No 160/2019 of 14 June 2019”.

(c) In Article 12(3)(c), as regards the EFTA States, the words “1 January 2017” shall read “one year after the date of entry into force of Decision of the EEA Joint Committee No 160/2019 of 14 June 2019”.

(d) Articles 14 to 19 and Article 25(2) shall not apply.


9bd.\(^{(1770)}\) **32015 R 2067**: Commission Implementing Regulation (EU) 2015/2067 of 17 November 2015 establishing, pursuant to Regulation (EU) No 517/2014 of the European Parliament and of the Council, minimum requirements and the conditions for mutual recognition for the certification of natural persons as regards stationary refrigeration, air conditioning and heat pump equipment, and refrigeration units of refrigerated trucks and trailers, containing fluorinated greenhouse gases and for the certification of companies as regards stationary refrigeration, air conditioning and heat pump equipment, containing fluorinated greenhouse gases (OJ L 301, 18.11.2015, p. 28).


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The first paragraph of Article 3 shall not apply to the EFTA States.

(b) In the second paragraph of Article 3, with regard to the EFTA States, the words “Annex A, B or C to Regulation (EC) No 338/97” shall read “the relevant parts of the legislation implementing the Convention on International Trade in Endangered Species of Wild Fauna and Flora in that EFTA State”.

(c) In Article 8(3), (5) and (6), if monitoring organisations of an EFTA State are concerned and without prejudice to Protocol 1 to the Agreement, the word “Commission” shall read “the EFTA Surveillance Authority”.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

If monitoring organisations of an EFTA State are concerned and without prejudice to Protocol 1 to the Agreement, the word “Commission” shall read “the EFTA Surveillance Authority”.


The provision of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 6(2)(b), if monitoring organisations of an EFTA State are concerned and without prejudice to Protocol 1 to the Agreement, the word “Commission” shall read “the EFTA Surveillance Authority”.


9da. (179) 32021 D 0958: Commission Implementing Decision (EU) 2021/958 of 31 May 2021 laying down the format for reporting data and information on fishing gear placed on the market and waste fishing gear collected in Member States and the format for the quality check report in accordance with Articles 13(1)(d) and 13(2) of

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(178) Point and adaptation text inserted by Decision No 75/2013 (OJ L 291, 31.10.2013, p. 32 and EEA Supplement No 61, 31.10.2013, p. 36), e.i.f. 1.5.2015.

(179) Point and adaptation text inserted by Decision No 75/2013 (OJ L 291, 31.10.2013, p. 32 and EEA Supplement No 61, 31.10.2013, p. 36), e.i.f. 1.5.2015.


The provisions of the Decision shall, for the purposes of this Agreement, be read with the following adaptation: Articles 1 and 2(1) shall not apply to Liechtenstein.


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

Articles 7, 11 and 16 shall not apply to Liechtenstein.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

This Regulation shall not apply to Liechtenstein.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE (TM)


(TM2) Point inserted by Decision No 13/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 4.2.2023.

(TM3) Point and adaptation inserted by Decision No 13/2023 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 4.2.2023.


The Contracting Parties take note of the content of the following acts:


### XVIII. INFORMATION TECHNOLOGY, TELECOMMUNICATION AND DATA PROCESSING

**ACTS REFERRED TO**

1. [ ]<sup>1789</sup>

2. [ ]<sup>1790</sup>


4. [ ]<sup>1791</sup>

4a.<sup>1792</sup> [ ]<sup>1793</sup>

4b.<sup>1794</sup> [ ]<sup>1795</sup>

4c.<sup>1796</sup> [ ]

4d.<sup>1797</sup> [ ]

4e.<sup>1798</sup> [ ]

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<sup>1793</sup> Text of point 4a (Commission Decision 94/11/EC) deleted with effect from 10 July 1998 by Decision No 63/98 (OJ L 100, 15.4.1999, p. 50 and EEA Supplement No 16, 15.4.1999, p. 96), e.i.f. 5.7.1998.


<sup>1795</sup> Text of point 4b (Commission Decision 94/12/EC) deleted with effect from 10 July 1998 by Decision No 63/98 (OJ L 100, 15.4.1999, p. 50 and EEA Supplement No 16, 15.4.1999, p. 96), e.i.f. 5.7.1998.


4f.\(^{(179)}\)  

4g.\(^{(180)}\)  

4h.\(^{(181)}\)  

4i.  

4j.  

4k.  

4l.  

4m.\(^{(183)}\)  

4n.  

4o.  


\(^{(190)}\) Point inserted by Decision No 61/98 (OJ L 100, 15.4.1999, p. 46 and EEA Supplement No 16, 15.4.1999, p. 79), e.i.f. 5.7.1998.
4r. COMMISSION DECISION 97/486/EC of 9 July 1997 on a common technical regulation for the general attachment requirements for terminal equipment to interface to Open Network Provision (ONP) two-wire analogue leased lines (OJ L 208, 2.8.1997, p. 44).

4s. COMMISSION DECISION 97/487/EC of 9 July 1997 on a common technical regulation for the general attachment requirements for terminal equipment to interface to Open Network Provision (ONP) four-wire analogue leased lines (OJ L 208, 2.8.1997, p. 47).


4u. [ ]

4v. [ ]

4w. COMMISSION DECISION 97/528/EC of 9 July 1997 on a common technical regulation for the general attachment requirements for mobile stations intended to be used with Phase II public digital cellular telecommunications networks operating in the DCS 1800 band (OJ L 215, 7.8.1997, p. 60).

4x. [ ]


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(1811) Point inserted by Decision No 61/98 (OJ L 100, 15.4.1999, p. 46 and EEA Supplement No 16, 15.4.1999, p. 79), e.i.f. 5.7.1998.


4ze. [ ] 4ze2

4zf. [ ] 4zf2

4zg. [ ] 4zg2


4zu.[1862] 398 D 0482: Council Decision 98/482/EC of 20 July 1998 on a common technical regulation for the attachment requirements for connection to the analogue public switched telephone networks (PSTNs) of terminal equipment (excluding terminal equipment supporting the voice telephony justifiable case service) in which network addressing, if provided, is by means of dual tone multi-frequency (DTMF) signalling (OJ L 216, 4.8.1998, p. 8).


4zw.[1864] 398 D 0575: Commission Decision 98/575/EC of 16 September 1998 on a common technical regulation for the general attachment requirements for mobile stations intended to be used with Phase II public digital

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4zhh. Commission Decision 1999/303/EC of 12 April 1999 on a common technical regulation for connection to the analogue public switched telephone networks (PSTNs) of terminal equipment supporting the voice telephony justified case service in which network addressing, if provided, is by means of dual tone multi-frequency (DTMF) signalling (OJ L 118, 6.5.1999, p. 55).


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(1) Listed here for purposes of information only. For application, see Annex XIV on competition.

4zzq.[1862] 32013 D 0638: Commission Decision 2013/638/EU of 12 August 2013 on essential requirements relating to marine radio communication equipment which is intended to be used on non-SOLAS vessels and to participate in the Global Maritime Distress and Safety System (GMDSS) (OJ L 296, 7.11.2013, p. 22).


The provisions of the Regulation shall, for the purpose of this Agreement, be read with the following adaptation:


[1867] Indent and words “, as amended by: “added by Decision No 234/2021 (OJ L [to be published]) and EEA Supplement No [to be published]”, as E.1.F. pending.

In Annex II, the following shall be added to the abbreviations:

“Iceland (IS)
Liechtenstein (LI)
Norway (NO)”.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

The Contracting Parties take note of the content of the following acts:


XIX. GENERAL PROVISIONS IN THE FIELD OF TECHNICAL BARRIERS TO TRADE

ACTS REFERRED TO


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The second subparagraph of Article 1(1)(c) shall be replaced by the following:

“The term ‘technical specification’ also covers production methods and processes used in respect of products intended for human and animal consumption, and in medicinal products as defined in Article 1 of Directive 2001/83/EC (as incorporated into point 15 q of Chapter XIII of Annex II to the Agreement by Decision of the EEA Joint Committee No 82/2002 of 25 June 2002), as well as production methods and processes relating to other products, where these have an effect on their characteristics.”;

(b) The following shall be added to the first subparagraph of Article 5(1):

“A full text of the draft technical regulation notified shall be made available in the original language as well as in a full translation into one of the official languages of the Union.”;

(c) The following subparagraph shall be added to Article 5(1):


“The Commission on behalf of the Union, on the one side, and the EFTA Surveillance Authority or the EFTA States through the EFTA Surveillance Authority, on the other side, may ask for further information on a draft technical regulation notified.”;

(d) The following subparagraph shall be added to Article 5(2):

“The comments of the EFTA States shall be forwarded by the EFTA Surveillance Authority to the Commission in the form of a single coordinated communication and the comments of the Union shall be forwarded by the Commission to the EFTA Surveillance Authority.”;

(e) The terms “Member State” and “Commission” in Article 6(1), (2) and (7) shall be replaced respectively by the terms “EFTA State” and “EFTA Surveillance Authority”.

(f) Article 6(3), (4), (5) and (6) shall not apply.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 2(7), the words “Article 57 TFEU” shall be replaced by “Article 37 of the EEA Agreement”.

3.

3a.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The following shall be added at the end of Article 4(2):

“Liechtenstein shall also have recourse to the national accreditation body of Switzerland for the product sectors covered by the Agreement between the European Community and the Swiss Confederation on mutual recognition in relation to conformity assessment and in respect of which EU and Swiss requirements are deemed equivalent pursuant to Article 1(2) and (3) of that Agreement”.

(b) Products exported from Liechtenstein to the other Contracting Parties may be subjected to border controls according to Articles 27-29.


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(appliances burning gaseous fuels), 91/263/EEC (telecommunications terminal equipment), 92/42/EEC (new hot-water boilers fired with liquid or gaseous fuels) and 73/23/EEC (electrical equipment designed for use within certain voltage limits) (OJ No L 220, 30.8.1993, p. 1).


The provisions of the Directive shall, for the purposes of the present Agreement, be read with the following adaptations:

(a) In Annex I, point 1(a) the following shall be added to the list of written indications concerning "upper":
   IS Efri hluti
   N Overdel
   \[\text{[1883]}\]

(b) In Annex I, point 1(b) the following shall be added to the list of written indications concerning "lining and sock":
   IS Fôður og bindsóli
   N För og bindsåle
   \[\text{[1884]}\]

(c) in Annex I, point 1(c) the following shall be added to the list of written indications concerning "outer sole":
   IS Slitsóli
   N Yttersåle
   \[\text{[1885]}\]


\footnote{1883} Indent added by Decision No 159/2014 (OJ L 15, 22.1.2015, p. 87 and EEA Supplement No 5, 22.1.2015, p. 10), e.l.f. pending; it shall apply from 9.7.2014.

(d) In Annex I, point 2(a)(i) the following shall be added to the list of written indications concerning "leather":

| IS Lædur |
| N lær |

\[(1886)\]

(e) In Annex I, point 2(a)(ii) the following shall be added to the list of written indications concerning "coated leather":

| IS Húðað leður |
| N Belagt lær |

\[(1887)\]

(f) In Annex I, point 2(b) the following shall be added to the list of written indications concerning natural textile materials and synthetic or non-woven textile materials:

| IS Textilefni |
| N Tekstilmaterialer |

\[(1888)\]

(g) In Annex I, point 2(c) the following shall be added to the list of written indications concerning "all other materials":

| IS Ölj önnur efni |
| N Andre materialer |

\[(1889)\]

3f. \[(1890)\]


-\[(1892)\] 1 03 T: Act concerning the conditions of accession of the Czech Republic, the Republic of Estonia, the Republic of Cyprus, the Republic of Latvia, the Republic of Lithuania, the Republic of Hungary, the Republic of Malta, the Republic of Poland, the Republic of Slovenia and the Slovak Republic and the adjustments to the Treaties on which the European Union is founded adopted on 16 April 2003 (OJ L 236, 23.9.2003, p. 33),


3k. {1999} 32006 D 0502: Commission Decision 2006/502/EC of 11 May 2006 requiring Member States to take measures to ensure that only lighters which are child-resistant are placed on the market and to prohibit the placing on the market of novelty lighters (OJ L 198, 20.7.2006, p. 41), as amended by:


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{1986} Indent and words “, as amended by:” added by Decision No 126/2012 (OJ L 309, 8.11.2012, p. 4 and EEA Supplement No 63, 8.11.2012, p. 5), e.i.f. 1.4.2013.


{1990} Indent and words “, as amended by:” added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.

{1991} Indent and words “, as amended by:” added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.

{1992} Indent and words “, as amended by:” added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.

{1993} Indent and words “, as amended by:” added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.


3n. 32009 D 0251: Commission Decision 2009/251/EC of 17 March 2009 requiring Member States to ensure that products containing the biocide dimethylfumarate are not placed or made available on the market (OJ L 74, 20.3.2009, p. 32), as amended by:


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1990  Point and indent inserted by Decision No 118/2010 (OJ L 58, 3.3.2011, p. 75 and EEA Supplement No 12, 3.3.2011, p. 16), e.l.f. 11.11.2010.
This Directive shall not apply to Liechtenstein.

The provisions of the Directive shall, for the purposes of the present Agreement, be read with the following adaptation:

Article 11 shall not apply.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) As regards the EFTA States, this Regulation shall only apply to products covered by Article 8(3) of the EEA Agreement.

(b) The Regulation shall not apply to Liechtenstein in relation to products covered by Annex I, Chapters XII and XXVII of Annex II and Protocol 47 to the EEA Agreement, as long as the application of the Agreement between the European Community and the Swiss Confederation on trade in agricultural products is extended to Liechtenstein.

(c) The words “Article 34 TFEU” shall read “Article 11 of the EEA Agreement”.

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[1922] Indent added by Decision No 40/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 18.3.2023.
The words “Article 36 TFEU” shall read “Article 13 of the EEA Agreement”.

In Article 8, the words “the Commission” shall be replaced by the words “the EFTA Surveillance Authority” when the administrative decision in question has been taken by an authority located in an EFTA State.


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**

The Contracting Parties take note of the content of the following acts:


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*1927* Point inserted by Decision No 14/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.7.2023.


*1931* Point inserted by Decision No 64/98 (OJ L 100, 15.4.1999, p. 52 and EEA Supplement No 16, 15.4.1999, p. 113), e.i.f. 15.7.1998 and subsequently corrected by Corrigendum noted in the EEA Joint Committee Meeting on 4.10.2013.
14. {1352} **96/C 224/03:** Council Resolution of 8 July 1996 on legislative and administrative simplification in the field of the internal market (OJ C 224, 18.9.1996, p. 5).


**XX. FREE MOVEMENT OF GOODS – GENERAL**

**ACTS REFERRED TO**{1359}


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**

The Contracting Parties take note of the content of the following acts:

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{1352} Point inserted by Decision No 64/98 (OJ L 100, 15.4.1999, p. 52 and EEA Supplement No 16, 15.4.1999, p. 113), e.i.f. 15.7.1998 and subsequently corrected by Corrigendum noted in the EEA Joint Committee Meeting on 4.10.2013.


{1356} Point inserted by Decision No 14/2020 (OJ L 49, 16.2.2023, p. 32 and EEA Supplement No 13, 16.2.2023, p. 32), e.i.f. 8.2.2020.

{1357} Point inserted by Decision No 14/2020 (OJ L 49, 16.2.2023, p. 32 and EEA Supplement No 13, 16.2.2023, p. 32), e.i.f. 8.2.2020.

{1358} Point inserted by Decision No 14/2020 (OJ L 49, 16.2.2023, p. 32 and EEA Supplement No 13, 16.2.2023, p. 32), e.i.f. 8.2.2020.

2. [1948] 380 Y 1003(01): Communication from the Commission concerning the consequences of the judgment given by the Court of Justice of the European Communities on 20 February 1979 in Case 120/78 ("Cassis de Dijon") (OJ No C 256, 3.10.80, p. 2).

3. [1941] 585 PC 0310: Commission Communication on the completion of the Internal Market COM (85) 310 Final ("White Paper").


5. [1943] [


**XXL CONSTRUCTION PRODUCTS**

**ACTS REFERRED TO**


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[1945] Point inserted by Decision No 49/2010 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.5.2010.


1f. [ ]


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ln. [ ]


1zr. []


prefabricated wood-based load-bearing stressed skin panels and self-supporting composite lightweight panels (OJ L 180, 19.7.2000, p. 40), as amended by:


1zw.**


1xz.**


1zy.**


1zz.**


1zza.**


1zvb.**


1zzc.**


1zzd.**


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- 32022 D 0381: Commission Implementing Decision (EU) 2022/381 of 4 March 2022 (OJ L 75, 7.3.2022, p. 1),
- 32022 D 1517: Commission Implementing Decision (EU) 2022/1517 of 9 September 2022 (OJ L 235, 12.9.2022, p. 65),


The provisions of this Decision shall, for the purposes of the present Agreement, be read with the following adaptations:

(a) The second paragraph of chapter 3.2.2. of the Annex shall be replaced by the following:

If consensus is achieved in the Technical Board on applications according to Article 8.2a of the CDP, then the appropriate information is sent with the approval of the President of EOTA to the EEA Joint Committee to obtain authority to issue ETAs. If consensus cannot be achieved in the Technical
Board it will be passed to the Executive Commission for decision, as to whether it should be submitted to the EEA Joint Committee.

(b) In the third paragraph of chapter 3.2.2. of the Annex the term 'Commission of the EC' shall be replaced by the term 'EEA Joint Committee'.

(c) In the last sentence of chapter 3.2.5. of the Annex the words 'shall be referred to the Standing Committee on Construction (the Directive, Article 9.2) via the Commission of the EC.' shall be replaced by 'shall be referred to the EEA Joint Committee.'


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Footnotes:


2v. **2017 R 1228**: Commission Delegated Regulation (EU) 2017/1228 of 20 March 2017 on the conditions for classification, without testing, of external render and internal plasters based on organic binders covered by the harmonised standard EN 15824 and rendering and plastering mortars covered by the harmonised standard EN 998-1 with regard to their reaction to fire (OJ L 177, 8.7.2017, p. 4).


### ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**[2051]**

The Contracting Parties take note of the content of the following acts:


### XXII. PERSONAL PROTECTIVE EQUIPMENT

#### ACTS REFERRED TO

1.  


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ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE{286a}

The Contracting Parties take note of the content of the following acts:


**XXIII. TOYS**

ACTS REFERRED TO

1. [] {285}


- **32012 L 0007:** Commission Directive 2012/7/EU of 2 March 2012 (OJ L 64, 3.3.2012, p. 7),


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{288} Indent and words “as amended by:” added by Decision No 114/2013 (OJ L 318, 28.11.2013, p. 16 and EEA Supplement No 67, 28.11.2013, p. 18), e.i.f. 1.4.2014.

{289} Indent added by Decision No 85/2014 (OJ L 310, 30.10.2014, p. 38 and EEA Supplement No 63, 30.10.2014, p. 29), e.i.f. 17.5.2014.

{290} Indent added by Decision No 276/2014 (OJ L 311, 26.11.2015, p. 29 and EEA Supplement No 71, 26.11.2015, p. 28), e.i.f. 13.12.2014.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE{[897]}

The Contracting Parties take note of the content of the following acts:

2. C/87/93/p. 3: Commission Communication pursuant to Article 9(2) of Council Directive 88/378/EEC regarding the list of bodies approved by the Member States responsible for carrying out the EC type-examination referred to in Articles 8(2) and 10 of that Directive (OJ No C 87, 27.3.1993, p. 3).


XXIV. MACHINERY


{[901] Indent added by Decision No 209/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

{[902] Indent added by Decision No 209/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

{[903] Indent added by Decision No 210/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

{[904] Indent added by Decision No 210/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.

{[905] Indent added by Decision No 345/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 11.12.2021.


ACTS REFERRED TO

1. [ ]

1a. [ ]

1b. [ ]


---


{2082} Indent and words “, as amended by:”, added by Decision No 279/2014 (OJ L 311, 26.11.2015, p. 32 and EEA Supplement No 71, 26.11.2015, p. 31), c.f. 1.6.2015.


{2089} Indent added by Decision No 70/2022 (OJ L 182, 7.7.2022, p. 50 and EEA Supplement No 45, 7.7.2022, p. 38), c.f. 19.3.2022.


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**

The Contracting Parties take note of the content of the following acts:


XXV. TOBACCO

ACTS REFERRED TO

1. [2201] [ ]

2. [2202] [ ]


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptations:

(a) In the second subparagraph of Article 5(1), as regards the EFTA States, the words “by 20 November 2016” shall read “at the latest six months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

(b) The following subparagraph shall be added to Article 6(4) and Article 7(13):

“In cases concerning manufacturers and importers in the EFTA States, the EFTA Surveillance Authority shall collect any fees charged by the Commission.”.

(c) With regard to Norway, the following subparagraph shall be added to Article 12(1):

“Taking into account the specific national circumstances supported by statistics regarding the health risks related to the use and use patterns of tobacco for oral use, tobacco for oral use placed on the market in Norway may carry the following alternative health warning:

“This tobacco product increases the risk of harm to the foetus and stillbirth””.

(d) In Article 15(13), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.


(c) In Article 16(3), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

(f) The prohibition in Article 17 shall not apply to the placing on the market in Norway of the product defined in Article 2, point (8). Norway shall ban export of the product defined in Article 2, point (8), to all Contracting Parties to this Agreement, with the exception of Sweden.

(g) In Article 30, as regards the EFTA States, the words “20 May 2017” shall read “one year after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

In Article 30, points (a) and (c), as regards the EFTA States, the words “20 May 2016” shall read “the date of entry into force Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

In Article 30, point (b), as regards the EFTA States, the words “20 November 2016” shall read “six months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.


3f. [2109] 32016 D 0786: Commission Implementing Decision (EU) 2016/786 of 18 May 2016 laying down the procedure for the establishment and operation of an independent advisory panel assisting Member States and the Commission in determining whether tobacco products have a characterising flavour (OJ L 131, 20.5.2016, p. 79).

3g. [2110] 32016 D 0787: Commission Implementing Decision (EU) 2016/787 of 18 May 2016 laying down a priority list of additives contained in cigarettes and roll-your-own tobacco subject to enhanced reporting obligations (OJ L 131, 20.5.2016, p. 88).

The provisions of the Decision shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 2, as regards the EFTA States, the words “1 January 2017” shall read “the date of entry into force of Decision of the EEA Joint Committee No 7/2022 of 4 February 2022”.

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[2106] Point 3e inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

[2107] Point 3d inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

[2108] Point 3e inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

[2109] Point 3f inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

[2110] Point 3g and adaptation text inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.
3h. 32016 R 0779: Commission Implementing Regulation (EU) 2016/779 of 18 May 2016 laying down uniform rules as regards the procedures for determining whether a tobacco product has a characterising flavor (OJ L 131, 20.5.2016, p. 48).


The provisions of the Regulation shall, for the purposes of the EEA Agreement, be read with the following adaptations:

(a) In Article 7(6)(a), as regards the EFTA States, the words “20 May 2020” shall read “sixteen months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(b) In Article 7(6)(b), as regards the EFTA States, the words “20 May 2021” shall read “two years after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(c) In Article 37(1), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(d) In Article 37(2), as regards the EFTA States, the words “20 May 2024” shall read “five years after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.


The provisions of the Decision shall, for the purposes of the EEA Agreement, be read with the following adaptations:

(a) In Article 3(4), as regards the EFTA States, the words “20 September 2018” shall read “eight months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(b) In Article 4(3), as regards the EFTA States, the words “20 September 2018” shall read “eight months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(c) In Article 9(1), the words “20 May 2019” shall, as regards the EFTA States, read “sixteen months after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

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Point 3h inserted by Decision No 8/2022 (OJ L 175, 30.6.2022, p. 17 and EEA Supplement No 42, 30.6.2022, p. 15), e.l.f. pending.

Point 3i inserted by Decision No 9/2022 (OJ L 175, 30.6.2022, p. 18 and EEA Supplement No 42, 30.6.2022, p. 16), e.l.f. pending.

Point 3j and adaptation text inserted by Decision No 9/2022 (OJ L 175, 30.6.2022, p. 18 and EEA Supplement No 42, 30.6.2022, p. 16), e.l.f. pending.

Point 3k and adaptation text inserted by Decision No 9/2022 (OJ L 175, 30.6.2022, p. 18 and EEA Supplement No 42, 30.6.2022, p. 16), e.l.f. pending, and subsequently corrected [before publication] by Corrigendum of 10.6.2022.
In Article 9(1), the words “20 May 2020” shall, as regards the EFTA States, read “two years and four months after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(d) In Article 9(2), the words “20 May 2024” shall, as regards the EFTA States, read “five years after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

In Article 9(2), the words “20 May 2026” shall, as regards the EFTA States, read “seven years after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.


XXVI. ENERGY

ACTS REFERRED TO

1. [ ] [2117]

XXVII. SPIRIT DRINKS

Contracting Parties shall authorise imports and marketing of spirit drinks which are in conformity with the Community legislation as listed in this Chapter. For all other purposes, EFTA States may continue to apply their national legislation.

[2118] This Chapter shall not apply to Liechtenstein as long as the application of the Agreement between the European Community and the Swiss Confederation on trade in agricultural products is extended to Liechtenstein.

ACTS REFERRED TO:

1. [ ] [2119]

2. [ ] [2120]


3.  

4.  


5.  


6.  


7.  


8.  


9.  


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The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The provisions of the Regulation shall not prejudice the right of the EFTA States to prohibit, on a non-discriminatory basis, the placing on their national market of spirit drinks for direct human consumption which exceed an alcoholic strength of 60%.

(b) The EFTA States shall be invited to send observers to the meetings of the Committee for Spirit Drinks, as referred to in Article 47, dealing with matters which fall within acts referred to in the Agreement. The representatives of the EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.

(c) Paragraph 4(d) of Protocol 1 to the Agreement shall not apply to Chapter III of the Regulation.

9a. [ ]


\[2139\] Indent and words “; as amended by:” added by Decision No 132/2022 (OJ L 246, 22.9.2022, p. 81 and EEA Supplement No 61, 22.9.2022, p. 80), e.i.f. 30.4.2022.


\[2135\] Point inserted by Decision No 71/2022 (OJ L 182, 7.7.2022, p. 51 and EEA Supplement No 45, 7.7.2022, p. 39), e.i.f. 19.3.2022.

\[2136\] Point inserted by Decision No 71/2022 (OJ L 182, 7.7.2022, p. 51 and EEA Supplement No 45, 7.7.2022, p. 39), e.i.f. 19.3.2022.

\[2137\] Point inserted by Decision No 71/2022 (OJ L 182, 7.7.2022, p. 51 and EEA Supplement No 45, 7.7.2022, p. 39), e.i.f. 19.3.2022.

\[2138\] Point inserted by Decision No 71/2022 (OJ L 182, 7.7.2022, p. 51 and EEA Supplement No 45, 7.7.2022, p. 39), e.i.f. 19.3.2022.

9af. Article 30(2) of Regulation (EU) 2020/179 of 3 February 2020 approving amendments to the specification for a spirit drink whose name is registered as a geographical indication (Berliner Kümmel) (OJ L 37, 10.2.2020, p. 4).

9ag. Article 30(2) of Regulation (EU) 2020/623 of 30 April 2020 approving amendments to the specification for a spirit drink whose name is registered as a geographical indication (Ratafia de Champagne) (OJ L 144, 7.5.2020, p. 10).

9ah. Article 30(2) of Regulation (EU) 2020/1286 of 9 September 2020 approving amendments to the specification for a spirit drink whose name is registered as a geographical indication (Scotch Whisky) (OJ L 302, 16.9.2020, p. 4).

9ai. Article 30(2) of Regulation (EU) 2020/1287 of 9 September 2020 approving amendments to the specification for a spirit drink whose name is registered as a geographical indication (Hierbas de Mallorca) (OJ L 302, 16.9.2020, p. 6).

9aj. Article 30(2) of Regulation (EU) 2020/2079 of 8 December 2020 approving amendments to the product specification for a spirit drink whose name is registered as a geographical indication (Münchener Kümmel) (OJ L 423, 15.12.2020, p. 1).


9an. Article 30(2) of Regulation (EU) 2021/724 of 3 March 2021 laying down rules for the application of Regulation (EU) 2019/787 of the European Parliament and of the Council as regards the communications to be made by Member States to the Commission with regard to the bodies appointed to supervise ageing processes for spirit drinks and the competent authorities responsible for ensuring compliance with that Regulation (OJ L 155, 5.5.2021, p. 3).

The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 3(1), as regards the EFTA States, the words “25 August 2021” shall read “three months after the date of entry into force of the Decision of the EEA Joint Committee No 241/2022 of 23 September 2022”.


The provisions of the Regulation shall, for the purpose of this Agreement, be read with the following adaptation:

The following shall be added in Article 13:

“Notwithstanding the provisions of Protocol 1 to the Agreement, the communications made by the competent authorities of the EFTA States to the Commission pursuant to Article 13(1) shall follow the procedure set out in point (b). Point 4 of Protocol 1 shall not apply to Article 13.”


The provisions of the Regulation shall, for the purpose of this Agreement, be read with the following adaptation:

The following shall be added in Article 13:

“Notwithstanding the provisions of Protocol 1 to the Agreement, the communications made by the competent authorities of the EFTA States to the Commission pursuant to Article 13(1) shall follow the procedure set out in point (b). Point 4 of Protocol 1 shall not apply to Article 13.”


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

The EFTA States concerned shall be invited to send observers to the meetings of the Committee on aromatised wine products, as referred to in Article 34, dealing with matters which fall within the acts referred to in the Agreement. The representatives of the EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.


XXVIII. CULTURAL GOODS

ACTS REFEREDED TO

1. [ ]

The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptations:

In Article 14, the reference to “on or after 1 January 1993” shall with regard to Iceland and Norway read “on or after 1 January 1995” and with regard to Liechtenstein read “on or after 1 May 1995”.

XXIX. EXPLOSIVES FOR CIVIL USE { 2386 }

ACTS REFERRED TO


3. [2372] [ ]


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{188} ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

The Contracting Parties take note of the content of the following acts:


XXX. MEDICAL DEVICES {182}

ACTS REFERRED TO

1. [] {189}


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{275} Point inserted by Decision No 119/2010 (OJ L 58, 3.3.2011, p. 76 and EEA Supplement No 12, 3.3.2011, p. 18), e.i.f. 1.11.2012.


{277} Point inserted by Decision No 269/2014 (OJ L 311, 26.11.2015, p. 20 and EEA Supplement No 71, 26.11.2015, p. 19), e.i.f. 1.8.2015.

Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 6.7.2018.

{278} Indent and words "as amended by:" added by Decision No 109/2018 (OJ L 368, 5.11.2020, p. 10 and EEA Supplement No 71, 5.11.2020, p. 11), e.i.f. 1.6.2018.


{281} Heading, the words “The Contracting Parties take note of the content of the following acts:" and point added by Decision No 191/2018 (OJ L 75, 4.3.2021, p. 27 and EEA Supplement No 15, 4.3.2021, p. 20), e.i.f. 22.9.2018.


The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for Poland (Annex XII, Chapter I, Point 3), shall apply.


5.\[\text{[2193]}\]


7.\[\text{[2196]}\]


\[\text{[2191]}\] Indent added by Decision No 201/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 12.12.2020.

\[\text{[2192]}\] Indent added by Decision No 38/2021 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 6.2.2021.


\[\text{[2195]}\] Point inserted by Decision No 141/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 16.7.2013.


9. [ ] {2199}


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptations:

(a) The EFTA States shall participate fully in the Medical Device Coordination Group ("MDCG") established under Article 103, but shall not have the right to vote.

(b) The EFTA States shall participate in the European database on medical devices (Eudamed) set up by the Commission as referred to in Article 33.


The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

Notwithstanding the provisions of Protocol 1 to this Agreement, and unless otherwise provided for in this Agreement, the terms Member State(s) and competent authorities shall be understood to include, in addition to their meaning in the Regulation, the EFTA States and their competent authorities, respectively.


{2203} Indent and words “, as amended by:” added by Decision No 90/2020 (OJ L 78, 16.3.2023, p. 40 and EEA Supplement No 22, 16.3.2023, p. 38), e.i.f. 18.6.2020.

{2204} Indent added by Decision No 153/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

{2205} Point inserted by Decision No 39/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.2.2021.

{2206} Point inserted by Decision No 134/2022 (OJ L 246, 22.9.2022, p. 85 and EEA Supplement No 61, 22.9.2022, p. 82), e.i.f. 30.4.2022.


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[228] Point inserted by Decision No 48/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 18.3.2023.

[229] Point inserted by Decision No 48/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 18.3.2023.


[232] Indent added by Decision No 153/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

[233] Point inserted by Decision No 15/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 4.2.2023.

[234] Point inserted by Decision No 49/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 18.3.2023.

[235] Point inserted by Decision No 49/2023 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 18.3.2023.


[237] Point inserted by Decision No 144/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 24.4.2021.

[238] Point inserted by Decision No 211/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 10.7.2021.
The Contracying Parties take note of the following acts:


XXXI. RECREATIONAL CRAFT\{\[\]\}

ACTS REFERRED TO

1. [ ] \{[\[\]

2. \{[\[\]

3. \{[\[\]

XXXII. MARINE EQUIPMENT\{\[\]

ACTS REFERRED TO

1. [ ] \{[\[\]

2. \{[\[\]

   - \{[\[\]

3. \{[\[\]

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APPENDIX I\textsuperscript{2232}  

ENERGY LABELS  

SECTION I\textsuperscript{2233} [ ]


SECTION 2[2234] [ ]

SECTION 3\textsuperscript{(228)} [ ]

SECTION 4{(228)}

Commission Directive 96/60/EC

(household combined washer-driers)
<table>
<thead>
<tr>
<th>Energi</th>
<th>Kombinert vaske- og tørkemaskin</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Merke</strong></td>
<td>LOGO</td>
</tr>
<tr>
<td><strong>Modell</strong></td>
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</tr>
<tr>
<td><strong>Lavt forbruk</strong></td>
<td>B</td>
</tr>
<tr>
<td><strong>Høyt forbruk</strong></td>
<td>X.YZ</td>
</tr>
<tr>
<td>Energiforbruk</td>
<td>kWh</td>
</tr>
<tr>
<td>(Ved del C: vaskning og tøkning med full kapasitet / funksjonstilstand)</td>
<td>X.YZ</td>
</tr>
<tr>
<td>Vask &amp; centrifugering</td>
<td>kWh</td>
</tr>
<tr>
<td>Den faktiske energibruk avhenger av hvordan vaske- og tørkemaskinen brukes</td>
<td></td>
</tr>
<tr>
<td>Vaskeevne</td>
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</tr>
<tr>
<td>A: høy</td>
<td>G: lav</td>
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<tr>
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</tr>
<tr>
<td>Kapasitet</td>
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</tr>
<tr>
<td>Vasking</td>
<td>y.z</td>
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<tr>
<td>(bomull) kg</td>
<td></td>
</tr>
<tr>
<td>Tøkning</td>
<td>y.z</td>
</tr>
<tr>
<td>Vannforbruk (totalt)</td>
<td></td>
</tr>
<tr>
<td>(y.h)</td>
<td>yx</td>
</tr>
<tr>
<td>Lydnivå</td>
<td></td>
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<tr>
<td>Vasket</td>
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</tr>
<tr>
<td>dB(A) (Støy)</td>
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<td>Tøkning</td>
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Produktbeskyttede innhold.
ytterligere opplysninger.
SECTION 5\textsuperscript{(2237)} [ ]

### Energimerkning

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<th><strong>Modell</strong></th>
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<td><strong>Lavt forbruk</strong></td>
<td><strong>Oppvaskmaskin</strong></td>
</tr>
<tr>
<td>A</td>
<td>Logo</td>
</tr>
<tr>
<td>B</td>
<td>ABC</td>
</tr>
<tr>
<td>C</td>
<td>123</td>
</tr>
<tr>
<td>D</td>
<td></td>
</tr>
<tr>
<td>E</td>
<td></td>
</tr>
<tr>
<td>F</td>
<td></td>
</tr>
<tr>
<td>G</td>
<td></td>
</tr>
</tbody>
</table>

**Høyt forbruk**

<table>
<thead>
<tr>
<th>Energiforbruk kWh/oppvask</th>
<th><strong>X.YZ</strong></th>
</tr>
</thead>
</table>

(på grunnlag av testresultater for normal/program ved fastvennstillitning)

Den faktiske energibruken avhenger av hvordan maskinen brukes.

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<tr>
<th>Rengjøringssevne</th>
<th><strong>A B C D E F G</strong></th>
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<tbody>
<tr>
<td>A: høy</td>
<td>G: lav</td>
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<table>
<thead>
<tr>
<th>Tørkeevne</th>
<th><strong>A B C D E F G</strong></th>
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<tr>
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<th>Standardkuverter</th>
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<td><strong>YX</strong></td>
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<table>
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<tr>
<th>Lydnivå</th>
<th><strong>XY</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>DB(A) (Støy)</td>
<td></td>
</tr>
</tbody>
</table>

Produktbrosjyrene inneholder ytterligere opplysninger.

Europeisk standard EN 50542

Direktiv 92/77/EF om energimerkning av oppvaskmaskiner
SECTION 6\(^{(2238)}\)

*Commission Directive 2002/40/EC*

*(household electric ovens)*

Orka

Framleiðandi
Gerð

Góð nýtni

<table>
<thead>
<tr>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
<th>F</th>
<th>G</th>
</tr>
</thead>
</table>

Slæm nýtni

Orkunotkun (kWh)
Hitun: Hefðbundinn Blástursofn

Notkunarrými (lítrar)XYZ

Stærð:
Lítill Meðal Stór

Hávaði (dB(A) re 1 pW)

Nánari upplýsingar er að finna í bæklingum sem fylgja vörunum

Norm EN 50304 Rafmagnsbókunarofnarr Tískipun 2002/40/EB um orkuneringar
## Energi

### Merke

<table>
<thead>
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<th>Modell</th>
<th>Logo ABC 123</th>
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### Høyt energiforbruk

<table>
<thead>
<tr>
<th>Grade</th>
<th>Value</th>
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<tbody>
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<tr>
<td>E</td>
<td></td>
</tr>
<tr>
<td>F</td>
<td></td>
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<td>G</td>
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### Lavt energiforbruk

<table>
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<tr>
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<td>Tradisjonell oppvarming</td>
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<td>Varmluft</td>
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### Nettovolum (liter)

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<td>Stor</td>
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### Lydnivå (støy) (dB(A) re 1 pW)

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Produktbrosjyrene inneholder ytterligere opplysninger.

Standard EN 50304
Elektriske stekeovner
Direktiv 2002/40/EF om energimerking
Energi

Merke

Modell

Høy energiforbruk

A

B

C

D

E

F

G

Lav energiforbruk

Energiforbruk (kWh)

Oppvarmingsfunksjon:

Tradisjonell oppvarming

Varmluft

(basert på standardbelastning)

Nettovolum (liter)

Type:

Liten

Middels stor

Stor

Lydnivå (støy) (dB(A) re 1 pW)

Produktbrosjyrene inneholder ytterligere opplysninger

Standard EN 50304
Elektriske stekeovner
Direktiv 2002/40/EF om energimerking
SECTION 7{2239} [ ]

APPENDIX 2\footnote{2240}

ENERGY TABLES

SECTION 1\footnote{2241} [ ]
SECTION 2\(^{(224)}\) [ ]

SECT[ION 3\textsuperscript{[224]}] (1)
### Section 4{[244]}

*Commission Directive 96/60/EC*

*(household combined washer-driers)*

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<td>Energy</td>
<td>Energi</td>
<td>Orka</td>
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<td>Washer-drier</td>
<td>Kombinert vaske- og tøremaskin</td>
<td>Ítvottavel – þurkari</td>
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<td>Manufacturer</td>
<td>Merke</td>
<td>Framleiðandi</td>
</tr>
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<td>Model</td>
<td>Modell</td>
<td>Gerð</td>
</tr>
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<td>Lavt forbruk</td>
<td>Göð nýtni</td>
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<tr>
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<td>Less efficient</td>
<td>Høyt forbruk</td>
<td>Slæm nýtni</td>
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<td>3 1</td>
<td>Energy efficiency class ...... on a scale of A (more efficient) to G (less efficient)</td>
<td>Relativ energibruk ...... på skalaen A (lavt forbruk) til G (høyt forbruk)</td>
<td>Orkunýttiniflokkur ...... á kvarðanum A (göð nýtni) til G (slæm nýtni)</td>
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<td>V</td>
<td>Energy consumption</td>
<td>Energibruk</td>
<td>Orkunotkun</td>
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<tr>
<td>☐</td>
<td>kWh</td>
<td>kWh</td>
<td>kWh</td>
</tr>
<tr>
<td>5 2</td>
<td>Energy consumption for washing, spinning and drying</td>
<td>Energibruk til vasking, sentrifugering og tøring</td>
<td>Orkunotkun við þvott, þeytivindingu og þurrkun</td>
</tr>
<tr>
<td>☐</td>
<td>(To wash and dry a full capacity wash load at 60 ºC)</td>
<td>(ved 60 °C vasking og tøring med full kapasitetsutnyttelse)</td>
<td>(Til að þvo og þurrka þvott á 60 ºC- þvottalotu miðað við leyfilegt ámarksmagn taus)</td>
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<tr>
<td>VI</td>
<td>Washing (only) kWh</td>
<td>Vask og sentrifugering kWh</td>
<td>Ítvottur og þeytivindingu kWh</td>
</tr>
<tr>
<td>☐</td>
<td>Energy consumption for washing and spinning only</td>
<td>Energibruk pr vask og sentrifugering alene</td>
<td>Orkunotkun við þvott og þeytivindingu eingöngu</td>
</tr>
<tr>
<td>☐</td>
<td>Actual consumption will depend on how the appliance is used</td>
<td>Den faktiske energi-bruken avhenger av hvordan vaske- og tøremaskinen brukes</td>
<td>Raunotkun fer efir því hvernig tækið er notað</td>
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<tr>
<td>VII</td>
<td>Washing performance A (higher) G (lower)</td>
<td>Vaskeevne A (høy) G (lav)</td>
<td>Ítvottalefni A (meiri) til G (minni)</td>
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<td>☐</td>
<td>Washing performance class ...... on a scale of A (higher) to G (lower)</td>
<td>Vaskeevne ...... på skalaen fra A (høy) til G (lav)</td>
<td>Ítvottalefnið flokkur...... á kvarðanum A (meiri) til G (minni)</td>
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<td>8</td>
<td>5</td>
<td>Water remaining after spin ...% (as a</td>
<td>Restvanninhalt etter sentrifugering ...%</td>
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<td></td>
<td></td>
<td>proportion of dry weight of wash)</td>
<td>(i forhold til vekten av tørt tøy)</td>
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<tr>
<td>VIII</td>
<td>9</td>
<td>Spin speed (rpm)</td>
<td>Sentrifugerings-hastighet (omdr/min)</td>
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<tr>
<td>IX/X</td>
<td>10/11</td>
<td>Capacity (cotton) kg</td>
<td>Kapasitet (bomull) kg</td>
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<td>10</td>
<td>Washing</td>
<td>Vaskning</td>
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<td>IX</td>
<td>11</td>
<td>Drying</td>
<td>Tørkning</td>
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<td>Water consumption (total)</td>
<td>Vannforbruk (totalt)</td>
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<td>og tørkning</td>
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<td>Vannforbruk til vask-og sentrifugerings</td>
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<td>the drier (200 cycles)</td>
<td>personer som alltid tøker tøyet i</td>
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<td>Anslått årlig forbruk</td>
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<td>for a 4-person household, never using</td>
<td>for en husstand på fire</td>
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<td>the drier (200 cycles)</td>
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<td>Noise (dB(A) re 1 pW)</td>
<td>Lydnivå (dB(A) (Støy)</td>
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<td>Sentrifugering</td>
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<td>Tørkning</td>
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SECTION 5[2245] [ ]

### SECTION 6(2246)

*Commission Directive 2002/40/EC*

*(household electric ovens)*

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<td>Baking area</td>
<td>Bökunarrými</td>
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<td>Orkunotkun</td>
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<td>V 5 3 Forced air convection</td>
<td>Blástursofn</td>
<td>Varmluft</td>
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<td>V 5 3 Based on standard load</td>
<td>Miðað við staðalálag</td>
<td>Basert på standardbelastning</td>
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<td>VI 6 4 Usable volume(litres)</td>
<td>Notkunarrými (lítrar)</td>
<td>Nettovolum (liter)</td>
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<td>VII 7 5 Size</td>
<td>Stærð</td>
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<td>VII 7 5 Medium</td>
<td>Meðal</td>
<td>Middels stor</td>
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<td>Stór</td>
<td>Stor</td>
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<td>Time to cook standard load</td>
<td>Bökunartími við staðalálag</td>
<td>Koketid ved standardbelastning</td>
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<td>VIII 9 6 Noise (dB(A) re 1 pW)</td>
<td>Hávaði (dB(A) re 1 pW)</td>
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<td>Nánari upplýsingar er að finna í bæklingum sem fylgja vörunum</td>
<td>Produktbrosjyrene inneholder ytterligere opplysninger</td>
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<tr>
<td>11</td>
<td>The area of the largest baking sheet</td>
<td>Stærð stærstu bökunarplötu</td>
<td>Arealet til den største stekeplaten</td>
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SECTION 7[2247] [ ]

### HAZARD AND PRECAUTIONARY STATEMENTS IN ICELANDIC

The following shall be added to Annex III to Regulation (EC) No 1272/2008:

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<td>Óstöðugt, sprengifimt efni.</td>
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<td>Sprengifimt efni, hætta á alsprengingu.</td>
</tr>
<tr>
<td>H202</td>
<td>Sprengifimt efni, mikil hætta á sprengibroti.</td>
</tr>
<tr>
<td>H203</td>
<td>Sprengifimt efni, hætta á bruna, höggbylgju eða sprengibrotum.</td>
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<tr>
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<td>Hætta á bruna eða sprengibrotum.</td>
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<td>H205</td>
<td>Hætta á alsprengingu í bruna.</td>
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<td>Afar eldfim lofttegund.</td>
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<td>H211</td>
<td>Eldfim lofttegund.</td>
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<td>Úðabrúsi með afar eldfimum efnum.</td>
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<td>Úðabrúsi með eldfimum efnum.</td>
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<td>Eldfimt, fast efni.</td>
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<td>Eldfimt eða sprengifimt við hitun.</td>
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<td>Eldfimt við hitun.</td>
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<td>Kviknar í sjálfkrafa við snertingu við loft.</td>
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<td>H251</td>
<td>Sjálfhitandi, hætta á sjálfsvikvnikun.</td>
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<tr>
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<td>Sjálfhitandi í miklu efnismagni, hætta á sjálfsvikvnikun.</td>
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<tr>
<td>H260</td>
<td>Í snertingu við vatn myndast eldfimar lofttegundir sem er hætt við sjálfsvikvnikun.</td>
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<td>Eldfimar lofttegundir myndast við snertingu við vatn</td>
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<td>Getur valdið eða aukið bruna, eldmyndandi (oxandi).</td>
</tr>
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<td>Getur valdið bruna eða sprengingu, mjög eldmyndandi (oxandi).</td>
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<td>Getur aukið bruna, eldmyndandi (oxandi).</td>
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<td>Inniheldur lofttegund undir þýstingi, getur sprungið við hitun.</td>
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<td>Getur verið ætandi fyrir málma.</td>
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<td>Banvænt við inntóku.</td>
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<td>Eitrað við inntóku.</td>
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<tr>
<td>H302</td>
<td>Hættulegt við inntóku.</td>
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<tr>
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<td>Getur verið banvænt við inntóku ef það kemst í öndunarveg.</td>
</tr>
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<td>H350</td>
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<td>H351</td>
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H371 | Getur skaðað líffæri (eða tilgreinið öll líffæri sem verða fyrir áhrifum, ef þau eru kunn) (tilgreinið váhrifaleið ef sannað hefur verið svo öyggjandi sé að engin önnur váhrifaleið hefur þessa hættu í fór með sér).
H372 | Skaðar líffæri (tilgreinið öll líffæri sem verða fyrir áhrifum, ef þau eru kunn) við langvinn eða endurtekin váhrif (tilgreinið váhrifaleið ef sannað hefur verið svo öyggjandi sé að engin önnur váhrifaleið hefur þessa hættu í fór með sér).
H373 | Getur skaðað líffæri (tilgreinið öll líffæri sem verða fyrir áhrifum, ef þau eru kunn) við langvinn eða endurtekin váhrif (tilgreinið váhrifaleið ef sannað hefur verið svo öyggjandi sé að engin önnur váhrifaleið hefur þessa hættu í fór með sér).

H400 | Mjög eitrað lífi í vatni.
H410 | Mjög eitrað lífi í vatni, hefur langvinn áhrif.
H411 | Eitrað lífi í vatni, hefur langvinn áhrif.
H412 | Skaðlegt lífi í vatni, hefur langvinn áhrif.
H413 | Getur valdið langvinnum, skaðlegum áhrifum á líf í vatni.
H350i | Getur valdið krabbameini við innöndun.
H360F | Getur haft skaðleg áhrif á frjósemi.
H360D | Getur haft skaðleg áhrif á börn í móðurkviði.
H361f | Grunað um að hafa skaðleg áhrif á frjósemi.
H361d | Grunað um að hafa skaðleg áhrif á börn í móðurkviði.
H360FD | Getur haft skaðleg áhrif á frjósemi. Getur haft skaðleg áhrif á börn í móðurkviði.
H361fd | Grunað um að hafa skaðleg áhrif á frjósemi. Grunað um að hafa skaðleg áhrif á börn í móðurkviði.
H360Fd | Getur haft skaðleg áhrif á frjósemi. Grunað um að hafa skaðleg áhrif á börn í móðurkviði.
H360Df | Getur haft skaðleg áhrif á börn í móðurkviði. Grunað um að hafa skaðleg áhrif á frjósemi.

EUH 001 | Sprengifimt sem þurrefni.
EUH 006 | Sprengifimt með og án andrúmslofts.
EUH 014 | Hvarfast kröftuglega við vatn
EUH 018 | Getur myndað eðlifmar eða sprengifimar blöndur af efnagufu og andrúmslofti við notkun.
EUH 019 | Getur myndað sprengifim efnasambönd (peroxið).
EUH 044 | Sprengifimt við hitun í lokuðu rými.
EUH 029 | Myndar eitraða lofttegund í snertingu við vatn.
EUH 031 | Myndar eitraða lofttegund í snertingu við sýru.
EUH 032 | Myndar mjög eitraða lofttegund í snertingu við sýru.
EUH 066 | Endurtekin snerting getur valdið þurri eða sprunginni húð.
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The following shall be added to Part 2 of Annex IV of Regulation (EC) No 1272/2008:

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<tr>
<td>P382</td>
<td>Safnið upp því sem hellist niður.</td>
</tr>
<tr>
<td>P383</td>
<td>EFTIR INNTÓKU: Hringið í einrunarmíðstöð eða lækn.</td>
</tr>
<tr>
<td>P384</td>
<td>BERIST EFNIÐ Á HÚÐ: Sökkvið í kalt vatn/vefjið með blautu sáràbindi.</td>
</tr>
<tr>
<td>P385</td>
<td>BERIST EFNIÐ Á HÚÐ: Þvoið varlega með mikilli sápu og vatni.</td>
</tr>
<tr>
<td>P386</td>
<td>BERIST EFNIÐ Á HÚÐ: Þvoið með mikilli sápu og vatni</td>
</tr>
<tr>
<td>No.</td>
<td>Icelandic</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
</tr>
<tr>
<td>P304 + P341</td>
<td>EFTIR INNÖNDUN: Ef viðkomandi á erfitt með öndun skal flytja hann í ferskt loft og láta hann hvílast í stellingu sem léttir öndun.</td>
</tr>
<tr>
<td>P305 + 351 + P338</td>
<td>BERIST EFNIÐ Í AUGU: Skolið varlega með vatni í nokkrar minútur. Fjarlægðið snertilinsur ef það er auðvelt. Skolið áfram.</td>
</tr>
<tr>
<td>P306 + P360</td>
<td>EF EFNIÐ FER Á FÓT: Fót og húd, sem öhreinkast af efninu, skal skola strax með miklu vatni aður en farið er úr fótunum.</td>
</tr>
<tr>
<td>P307 + P311</td>
<td>EFum vahrif er að ræða: Hringið í EITRUNARMÍÐSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P308 + P313</td>
<td>EF um vahrif eða hugsanleg vahrif er að ræða: Leitið læknis.</td>
</tr>
<tr>
<td>P309 + P311</td>
<td>EF um vahrif er að ræða eða ef lasleika verður vart: Hringið í EITRUNARMÍÐSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P332 + P313</td>
<td>Ef efnið ertir húð: Leitið læknis.</td>
</tr>
<tr>
<td>P333 + P313</td>
<td>Ef efnið ertir húð eða útbrot koma fram: Leitið læknis.</td>
</tr>
<tr>
<td>P335 + P334</td>
<td>Dustið lausar agnið af húðinni. Sökkið í kalt vatni/vefjið með blautu sárabindi.</td>
</tr>
<tr>
<td>P337 + P313</td>
<td>Ef augnerting er viðvarandi: Leitið læknis.</td>
</tr>
<tr>
<td>P342 + P311</td>
<td>Ef vart verður einkenna frá öndunarvegi: Hringið í EITRUNARMÍÐSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P370 + P376</td>
<td>Ef eldur kemur upp: Stöðvið leka ef það er óhætt.</td>
</tr>
<tr>
<td>P370 + P378</td>
<td>Ef eldur kemur upp: Notið ... til að slökkva eldinn.</td>
</tr>
<tr>
<td>P370 + P380</td>
<td>Ef eldur kemur upp: Rýmið svæðið.</td>
</tr>
<tr>
<td>P371 + P380 + P375</td>
<td>Þegar um mikinn eld og mikil efnið er að ræða: Rýmið svæðið. Verið í fjarlægð frá eldinum við slökkvistörf vegna sprengihættu.</td>
</tr>
<tr>
<td>P401</td>
<td>Geymist ...</td>
</tr>
<tr>
<td>P402</td>
<td>Geymist á þurrum stað.</td>
</tr>
<tr>
<td>P403</td>
<td>Geymist á vel loftfræstum stað.</td>
</tr>
<tr>
<td>P404</td>
<td>Geymist í lokuðu hláti.</td>
</tr>
<tr>
<td>No.</td>
<td>Icelandic</td>
</tr>
<tr>
<td>----------</td>
<td>---------------------------------------------------------------------------</td>
</tr>
<tr>
<td>P405</td>
<td>Geymist á læstum stað.</td>
</tr>
<tr>
<td>P406</td>
<td>Geymist í tæringarþolnu/...fláti með tæringarþolnu innra lagi.</td>
</tr>
<tr>
<td>P407</td>
<td>Hafið loftibil á milli stafla/vöru Bretta.</td>
</tr>
<tr>
<td>P410</td>
<td>Hlífið við sólarljósi.</td>
</tr>
<tr>
<td>P411</td>
<td>Geymist við hitastig sem er ekki hærra en … oC/… °F.</td>
</tr>
<tr>
<td>P412</td>
<td>Setjið ekki í hærri hita en 50 oC/122 °F.</td>
</tr>
<tr>
<td>P413</td>
<td>Ef búlkavara vegur meira en ... kg/... pund skal ekki geyma hana í hærri hita en… oC/… °F.</td>
</tr>
<tr>
<td>P420</td>
<td>Má ekki geyma hjá öðru efni.</td>
</tr>
<tr>
<td>P422</td>
<td>Geymið innihald undir ...</td>
</tr>
<tr>
<td>P402 +</td>
<td>Geymist á þurrum stað. Geymist í lokuðu fláti.</td>
</tr>
<tr>
<td>P404</td>
<td></td>
</tr>
<tr>
<td>P403 +</td>
<td>Geymist á vel loftræstum stað. Ílát vera vel lukt.</td>
</tr>
<tr>
<td>P233</td>
<td></td>
</tr>
<tr>
<td>P403 +</td>
<td>Geymist á vel-loftræstum stað. Geymist á köldum stað.</td>
</tr>
<tr>
<td>P235</td>
<td></td>
</tr>
<tr>
<td>P410 +</td>
<td>Hlífið við sólarljósi. Geymist á vel loftræstum stað.</td>
</tr>
<tr>
<td>P403</td>
<td></td>
</tr>
<tr>
<td>P412</td>
<td>Hlífið við sólarljósi. Hlífið við hærri hita en 50 oC/122 °F.</td>
</tr>
<tr>
<td>P411 +</td>
<td>Geymist á köldum stað við hitastig sem er ekki hærra en … oC/… °F.</td>
</tr>
<tr>
<td>P235</td>
<td></td>
</tr>
<tr>
<td>P501</td>
<td>Fargið innihaldi/fláti hjá ...</td>
</tr>
</tbody>
</table>

**APPENDIX 6**

HAZARD AND PRECAUTIONARY STATEMENTS IN NORWEGIAN

The following shall be added to Annex III to Regulation (EC) No 1272/2008:

<table>
<thead>
<tr>
<th>No.</th>
<th>Norwegian</th>
</tr>
</thead>
<tbody>
<tr>
<td>H200</td>
<td>Ustabile eksplosive varer.</td>
</tr>
<tr>
<td>H201</td>
<td>Eksplosjonsfarlig; fare for masseeksplosjon.</td>
</tr>
<tr>
<td>H202</td>
<td>Eksplosjonsfarlig; stor fare for utkast av fragmenter.</td>
</tr>
<tr>
<td>H203</td>
<td>Eksplosjonsfarlig; fare for brann, trykkbølge eller utkast av fragmenter.</td>
</tr>
</tbody>
</table>

---

<table>
<thead>
<tr>
<th>No.</th>
<th>Norwegian</th>
</tr>
</thead>
<tbody>
<tr>
<td>H204</td>
<td>Fare for brann eller utkast av fragmenter.</td>
</tr>
<tr>
<td>H205</td>
<td>Fare for masseeksplosjon ved brann.</td>
</tr>
<tr>
<td>H220</td>
<td>Ekstremt brannfarlig gass.</td>
</tr>
<tr>
<td>H221</td>
<td>Brannfarlig gass.</td>
</tr>
<tr>
<td>H222</td>
<td>Ekstremt brannfarlig aerosol.</td>
</tr>
<tr>
<td>H223</td>
<td>Brannfarlig aerosol.</td>
</tr>
<tr>
<td>H224</td>
<td>Ekstremt brannfarlig væske og damp.</td>
</tr>
<tr>
<td>H225</td>
<td>Meget brannfarlig væske og damp.</td>
</tr>
<tr>
<td>H226</td>
<td>Brannfarlig væske og damp.</td>
</tr>
<tr>
<td>H228</td>
<td>Brannfarlig fast stoff.</td>
</tr>
<tr>
<td>H240</td>
<td>Eksplosjonsfarlig ved oppvarming.</td>
</tr>
<tr>
<td>H241</td>
<td>Brann- eller eksplosjonsfarlig ved oppvarming.</td>
</tr>
<tr>
<td>H242</td>
<td>Brannfarlig ved oppvarming.</td>
</tr>
<tr>
<td>H250</td>
<td>Selvantenner ved kontakt med luft.</td>
</tr>
<tr>
<td>H251</td>
<td>Selvopphetende; kan selvantenne.</td>
</tr>
<tr>
<td>H252</td>
<td>Selvopphetende i store mengder; kan selvantenne.</td>
</tr>
<tr>
<td>H260</td>
<td>Ved kontakt med vann utvikles brannfarlige gasser som kan selvantenne.</td>
</tr>
<tr>
<td>H261</td>
<td>Ved kontakt med vann utvikles brannfarlige gasser.</td>
</tr>
<tr>
<td>H270</td>
<td>Kan forårsake eller forsterke brann; oksiderende.</td>
</tr>
<tr>
<td>H271</td>
<td>Kan forårsake brann eller eksplosjon; sterkt oksiderende.</td>
</tr>
<tr>
<td>H272</td>
<td>Kan forsterke brann; oksiderende.</td>
</tr>
<tr>
<td>H280</td>
<td>Inneholder gass under trykk; kan eksploedere ved oppvarming.</td>
</tr>
<tr>
<td>H281</td>
<td>Inneholder nedkjølt gass; kan forårsake alvorlige forfrysninger.</td>
</tr>
<tr>
<td>H290</td>
<td>Kan være etsende for metaller.</td>
</tr>
<tr>
<td>H300</td>
<td>Dødelig ved svelging.</td>
</tr>
<tr>
<td>H301</td>
<td>Giftig ved svelging.</td>
</tr>
<tr>
<td>H302</td>
<td>Farlig ved svelging.</td>
</tr>
<tr>
<td>H304</td>
<td>Kan være dødelig ved svelging om det kommer ned i luftveiene.</td>
</tr>
<tr>
<td>H310</td>
<td>Dødelig ved hudkontakt.</td>
</tr>
<tr>
<td>H311</td>
<td>Giftig ved hudkontakt.</td>
</tr>
<tr>
<td>H312</td>
<td>Farlig ved hudkontakt.</td>
</tr>
<tr>
<td>H314</td>
<td>Gir alvorlige etseskader på hud og øyne.</td>
</tr>
<tr>
<td>H315</td>
<td>Irriterer huden.</td>
</tr>
<tr>
<td>H317</td>
<td>Kan utløse en allergisk hudreaksjon.</td>
</tr>
<tr>
<td>H318</td>
<td>Gir alvorlig øyeskade.</td>
</tr>
<tr>
<td>H319</td>
<td>Gir alvorlig øyeirritasjon.</td>
</tr>
<tr>
<td>H330</td>
<td>Dødelig ved innånding.</td>
</tr>
<tr>
<td>H331</td>
<td>Giftig ved innånding.</td>
</tr>
<tr>
<td>H332</td>
<td>Farlig ved innånding.</td>
</tr>
<tr>
<td>H334</td>
<td>Kan gi allergi eller astmasymptomer eller pustevansker ved innånding.</td>
</tr>
<tr>
<td>H335</td>
<td>Kan forårsake irritasjon av luftveiene.</td>
</tr>
<tr>
<td>H336</td>
<td>Kan forårsake døsighet eller svimmelhet.</td>
</tr>
<tr>
<td>No.</td>
<td>Norwegian</td>
</tr>
<tr>
<td>------</td>
<td>--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>H340</td>
<td>Kan gi genetiske skader &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H341</td>
<td>Mistenkes å kunne gi genetiske skader &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H350</td>
<td>Kan forårsake kreft &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H351</td>
<td>Mistenkes for å kunne forårsake kreft &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H360</td>
<td>Kan skade forplantningsevnen eller gi fosterskader &lt;Angi særlige virkninger dersom disse er kjent.&gt; &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H361</td>
<td>Mistenkes for å kunne skade forplantningsevnen eller gi fosterskader &lt;Angi særlige virkninger dersom disse er kjent.&gt; &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;</td>
</tr>
<tr>
<td>H362</td>
<td>Kan skade barn som ammes.</td>
</tr>
<tr>
<td>H370</td>
<td>Forårsaker organskader &lt;eller angi alle organer som påvirkes dersom disse er kjent.&gt; &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H371</td>
<td>Kan forårsake organskader &lt;eller angi alle organer som påvirkes dersom disse er kjent.&gt; &lt;Angi opptaksvei dersom det med sikkerhet er fastlått at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H372</td>
<td>Forårsaker organskader &lt;eller angi alle organer som påvirkes dersom disse er kjent.&gt; ved langvarig eller gjentatt eksponering &lt;Angi opptaksvei dersom det med sikkerhet er at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H373</td>
<td>Kan forårsake organskader &lt;eller angi alle organer som påvirkes dersom disse er kjent.&gt; ved langvarig eller gjentatt eksponering &lt;Angi opptaksvei dersom det med sikkerhet er at ingen andre opptaksveier er årsak til faren&gt;.</td>
</tr>
<tr>
<td>H400</td>
<td>Meget giftig for liv i vann.</td>
</tr>
<tr>
<td>H410</td>
<td>Meget giftig, med langtidsvirkning, for liv i vann.</td>
</tr>
<tr>
<td>H411</td>
<td>Giftig, med langtidsvirkning, for liv i vann.</td>
</tr>
<tr>
<td>H412</td>
<td>Skadelig, med langtidsvirkning, for liv i vann.</td>
</tr>
<tr>
<td>H413</td>
<td>Kan forårsake skadelige langtidsvirkninger for liv i vann.</td>
</tr>
<tr>
<td>H350i</td>
<td>Kan forårsake kreft ved innånding.</td>
</tr>
<tr>
<td>H360F</td>
<td>Kan skade forplantningsevnen.</td>
</tr>
<tr>
<td>H360D</td>
<td>Kan gi fosterskader.</td>
</tr>
<tr>
<td>H361f</td>
<td>Mistenkes for å kunne skade forplantningsevnen.</td>
</tr>
<tr>
<td>H361d</td>
<td>Mistenkes for å kunne gi fosterskader.</td>
</tr>
<tr>
<td>H360FD</td>
<td>Kan skade forplantningsevnen. Kan gi fosterskader.</td>
</tr>
<tr>
<td>H361fd</td>
<td>Mistenkes for å kunne skade forplantningsevnen. Mistenkes for å kunne gi fosterskader.</td>
</tr>
<tr>
<td>H360Fd</td>
<td>Kan skade forplantningsevnen. Mistenkes for å kunne gi fosterskader.</td>
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<tr>
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</tr>
<tr>
<td>H360Df</td>
<td>Kan gi fosterskader. Mistenkes for å kunne skade forplantningsevnen.</td>
</tr>
<tr>
<td>EUH 001</td>
<td>Eksplosjonsfarlig i tørr tilstand.</td>
</tr>
<tr>
<td>EUH 006</td>
<td>Eksplosjonsfarlig ved og uten kontakt med luft.</td>
</tr>
<tr>
<td>EUH 014</td>
<td>Reagerer voldsomt med vann.</td>
</tr>
<tr>
<td>EUH 018</td>
<td>Ved bruk kan brennbart damp/eksplosive damp-luft-blandinger dannes.</td>
</tr>
<tr>
<td>EUH 019</td>
<td>Kan danne eksplosive peroksider.</td>
</tr>
<tr>
<td>EUH 044</td>
<td>Eksplosjonsfarlig ved oppvarming i lukket rom.</td>
</tr>
<tr>
<td>EUH 029</td>
<td>Ved kontakt med vann utvikles giftig gass.</td>
</tr>
<tr>
<td>EUH 031</td>
<td>Ved kontakt med syrer utvikles giftig gass.</td>
</tr>
<tr>
<td>EUH 032</td>
<td>Ved kontakt med syrer utvikles meget giftig gass.</td>
</tr>
<tr>
<td>EUH 066</td>
<td>Gjentatt ekspansjonering kan gi tørre eller sprukket hud.</td>
</tr>
<tr>
<td>EUH 070</td>
<td>Giftig ved øyekontakt.</td>
</tr>
<tr>
<td>EUH 071</td>
<td>Etsende for luftveiene.</td>
</tr>
<tr>
<td>EUH 059</td>
<td>Farlig for ozonlaget.</td>
</tr>
<tr>
<td>EUH 203</td>
<td>Inneholder krom (VI). Kan gi en allergisk reaksjon.</td>
</tr>
<tr>
<td>EUH 204</td>
<td>Inneholder isocyanater. Kan gi en allergisk reaksjon.</td>
</tr>
<tr>
<td>EUH 205</td>
<td>Inneholder epoksyforbindelser. Kan gi en allergisk reaksjon.</td>
</tr>
<tr>
<td>EUH 206</td>
<td>Advarsel! Må ikke brukes sammen med andre produkter. Kan frigjøre farlige gasser (klor).</td>
</tr>
<tr>
<td>EUH 208</td>
<td>Inneholder &lt;navn på sensibiliserende stoff&gt;. Kan gi en allergisk reaksjon.</td>
</tr>
<tr>
<td>EUH 210</td>
<td>Sikkerhetsdatablad er tilgjengelig på anmodning.</td>
</tr>
<tr>
<td>EUH 401</td>
<td>Bruksanvisningen må følges, slik at man unngår risiko for menneskers helse og miljøet.</td>
</tr>
</tbody>
</table>

The following shall be added to Part 2 of Annex IV to Regulation (EC) No 1272/2008:

<table>
<thead>
<tr>
<th>No.</th>
<th>Norwegian</th>
</tr>
</thead>
<tbody>
<tr>
<td>P101</td>
<td>Dersom det er nødvendig med legehjelp, ha produsents beholder eller etikett for hånden.</td>
</tr>
<tr>
<td>P102</td>
<td>Oppbevares utilgjengelig for barn.</td>
</tr>
<tr>
<td>P103</td>
<td>Les etiketten før bruk.</td>
</tr>
<tr>
<td>P201</td>
<td>Innhent særskilt instruks før bruk.</td>
</tr>
<tr>
<td>P202</td>
<td>Skal ikke håndteres før alle advarsler er lest og oppfattet.</td>
</tr>
<tr>
<td>No.</td>
<td>Norwegian</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
</tr>
<tr>
<td>P210</td>
<td>Holdes vekk fra varme/gnister/åpen flamme/varme overflater. — Røyking forbudt.</td>
</tr>
<tr>
<td>P211</td>
<td>Ikke spray mot åpen flamme eller annen tennkilde.</td>
</tr>
<tr>
<td>P220</td>
<td>Må ikke brukes/oppbevares i nærheten av tøy /.../ brennbare materialer.</td>
</tr>
<tr>
<td>P221</td>
<td>Må ikke blandes med brennbare stoffer.</td>
</tr>
<tr>
<td>P222</td>
<td>Unngå kontakt med luft.</td>
</tr>
<tr>
<td>P223</td>
<td>Unngå all kontakt med vann, på grunn av fare for voldsom reaksjon og eksplosjonsaktig brann.</td>
</tr>
<tr>
<td>P230</td>
<td>Holdes fuktet med …</td>
</tr>
<tr>
<td>P231</td>
<td>Håndteres under inertgass.</td>
</tr>
<tr>
<td>P232</td>
<td>Beskyttes mot fuktighet.</td>
</tr>
<tr>
<td>P233</td>
<td>Hold beholderen tett lukket.</td>
</tr>
<tr>
<td>P234</td>
<td>Oppbevares bare i originalbeholder.</td>
</tr>
<tr>
<td>P235</td>
<td>Oppbevares kjølig.</td>
</tr>
<tr>
<td>P240</td>
<td>Beholder og mottakstyr jordes/potensialutlignes.</td>
</tr>
<tr>
<td>P241</td>
<td>Bruk elektrisk materiell /ventilasjonsmaterei/belysningsmateriell som er eksplosjonssikkert.</td>
</tr>
<tr>
<td>P242</td>
<td>Bruk bare verktøy som ikke avger gnister.</td>
</tr>
<tr>
<td>P243</td>
<td>Treff tiltak mot statisk elektrisitet.</td>
</tr>
<tr>
<td>P244</td>
<td>Reduksjonsventiler skal holdes fri for fett og olje.</td>
</tr>
<tr>
<td>P250</td>
<td>Må ikke utsettes for sliping/støt/…/friksjon.</td>
</tr>
<tr>
<td>P251</td>
<td>Beholder under trykk: Må ikke punkteres eller brennes, selv ikke etter bruk.</td>
</tr>
<tr>
<td>P260</td>
<td>Ikke innånd støv/røyk/gass/tåke/damp/aerosoler.</td>
</tr>
<tr>
<td>P261</td>
<td>Unngå innånding av støv/røyk/gass/tåke/damp/aerosoler.</td>
</tr>
<tr>
<td>P262</td>
<td>Må ikke komme i kontakt med øyne, huden eller klær.</td>
</tr>
<tr>
<td>P263</td>
<td>Unngå kontakt under graviditet/amming.</td>
</tr>
<tr>
<td>P264</td>
<td>Vask … grundig etter bruk.</td>
</tr>
<tr>
<td>P270</td>
<td>Ikke spis, drikk eller røyk ved bruk av produktet.</td>
</tr>
<tr>
<td>P271</td>
<td>Brukes bare utendørs eller i et godt ventilert område.</td>
</tr>
<tr>
<td>P272</td>
<td>Tilsøtde arbeidsklær må ikke fjernes fra arbeidsplassen.</td>
</tr>
<tr>
<td>P273</td>
<td>Unngå utslipp til miljøet.</td>
</tr>
<tr>
<td>P280</td>
<td>Benytt vernehansker /verneklær/vernebriller/ansiktsskjerm.</td>
</tr>
<tr>
<td>P281</td>
<td>Bruk påkrevd personlig verneutstyr.</td>
</tr>
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<td>------</td>
<td>---------------------------------------------------------------------------</td>
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<tr>
<td>P282</td>
<td>Bruk kuldeisolerende hansker /visir/øyevern.</td>
</tr>
<tr>
<td>P283</td>
<td>Benytt brannbestandige/flammehemmende klær.</td>
</tr>
<tr>
<td>P284</td>
<td>Bruk åndedrettsvern.</td>
</tr>
<tr>
<td>P285</td>
<td>Ved utilstrekkelig ventilasjon skal åndedrettsvern benyttes.</td>
</tr>
<tr>
<td>P301</td>
<td>VED SVELGING:</td>
</tr>
<tr>
<td>P302</td>
<td>VED HUDKONTAKT:</td>
</tr>
<tr>
<td>P303</td>
<td>VED HUDKONTAKT (eller håret):</td>
</tr>
<tr>
<td>P304</td>
<td>VED INNÅNDING:</td>
</tr>
<tr>
<td>P305</td>
<td>VED KONTAKT MED ØYNENE</td>
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<tr>
<td>P306</td>
<td>VED KONTAKT MED KLÆR:</td>
</tr>
<tr>
<td>P307</td>
<td>Ved eksponering:</td>
</tr>
<tr>
<td>P308</td>
<td>Ved eksponering eller mistanke om eksponering:</td>
</tr>
<tr>
<td>P309</td>
<td>Ved eksponering eller ubehag:</td>
</tr>
<tr>
<td>P310</td>
<td>Kontakt umiddelbart et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P311</td>
<td>Kontakt et GIFTINFORMASJONSSENTER eller lege.</td>
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<tr>
<td>P312</td>
<td>Kontakt et GIFTINFORMASJONSSENTER eller lege ved ubehag.</td>
</tr>
<tr>
<td>P313</td>
<td>Søk legehjelp.</td>
</tr>
<tr>
<td>P314</td>
<td>Søk legehjelp ved ubehag.</td>
</tr>
<tr>
<td>P315</td>
<td>Søk legehjelp umiddelbart.</td>
</tr>
<tr>
<td>P320</td>
<td>Særlig behandling kreves umiddelbart (se … på etiketten).</td>
</tr>
<tr>
<td>P321</td>
<td>Særlig behandling (se … på etiketten).</td>
</tr>
<tr>
<td>P322</td>
<td>Særlige tiltak (se … på etiketten).</td>
</tr>
<tr>
<td>P330</td>
<td>Skyll munnen.</td>
</tr>
<tr>
<td>P331</td>
<td>IKKE framkall brekning.</td>
</tr>
<tr>
<td>P332</td>
<td>Ved hudirritasjon:</td>
</tr>
<tr>
<td>P333</td>
<td>Ved hudirritasjon eller utstlett:</td>
</tr>
<tr>
<td>P334</td>
<td>Skyll i kaldt vann / anvend våt kompress.</td>
</tr>
<tr>
<td>P335</td>
<td>Børst bort løse partikler fra huden.</td>
</tr>
<tr>
<td>P336</td>
<td>Varm opp frostskadede legemidler med lunnent vann. Ikke gni på det skadede området.</td>
</tr>
<tr>
<td>P337</td>
<td>Ved vedvarende øyeirritasjon:</td>
</tr>
<tr>
<td>No.</td>
<td>Norwegian</td>
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<td>-----</td>
<td>-----------</td>
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<tr>
<td>P338</td>
<td>Fjern eventuelle kontaktlinser dersom dette enkelt lar seg gjøre. Fortsett skyllingen.</td>
</tr>
<tr>
<td>P340</td>
<td>Flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P341</td>
<td>Ved pustevansker, flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P342</td>
<td>Ved symptomer i luftveiene:</td>
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<tr>
<td>P350</td>
<td>Vask forsiktig med mye såpe og vann.</td>
</tr>
<tr>
<td>P351</td>
<td>Skyll forsiktig med vann i flere minutter.</td>
</tr>
<tr>
<td>P352</td>
<td>Vask med mye såpe og vann.</td>
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<tr>
<td>P353</td>
<td>Skyll/dusj huden med vann.</td>
</tr>
<tr>
<td>P360</td>
<td>Skyll umiddelbart tilsølte klær og hud med mye vann før klærnes fjernes.</td>
</tr>
<tr>
<td>P361</td>
<td>Tilsølte klær må fjernes straks.</td>
</tr>
<tr>
<td>P362</td>
<td>Tilsølte klær må fjernes og vaskes før de brukes på nytt.</td>
</tr>
<tr>
<td>P363</td>
<td>Tilsølte klær må vaskes før de brukes på nytt.</td>
</tr>
<tr>
<td>P370</td>
<td>Ved brann:</td>
</tr>
<tr>
<td>P371</td>
<td>Ved større brann og store mengder:</td>
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<tr>
<td>P372</td>
<td>Eksplosjonsfare ved brann.</td>
</tr>
<tr>
<td>P373</td>
<td>IKKE bekjemp brannen når den når eksplosive varer.</td>
</tr>
<tr>
<td>P374</td>
<td>Bekjemp brannen med normal forsiktighet på behørig avstand.</td>
</tr>
<tr>
<td>P375</td>
<td>Bekjemp brannen på avstand på grunn av eksplosjonsfare.</td>
</tr>
<tr>
<td>P376</td>
<td>Stopp lekkasje dersom dette kan gjøres på en sikker måte.</td>
</tr>
<tr>
<td>P377</td>
<td>Brann ved gasslekkasje: Ikke slukk med mindre lekkasjen kan stannes på en sikker måte.</td>
</tr>
<tr>
<td>P378</td>
<td>Slukk med:....</td>
</tr>
<tr>
<td>P380</td>
<td>Evakuér området.</td>
</tr>
<tr>
<td>P381</td>
<td>Fjern alle tennkilder dersom dette kan gjøres på en sikker måte.</td>
</tr>
<tr>
<td>P390</td>
<td>Absorber spill for å hindre materiell skade.</td>
</tr>
<tr>
<td>P391</td>
<td>Samle opp spill.</td>
</tr>
<tr>
<td>P301</td>
<td>VED SVELGING: Kontakt umiddelbart et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P310</td>
<td>VED SVELGING: Kontakt umiddelbart et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P301</td>
<td>VED SVELGING: Kontakt et GIFTINFORMASJONSSENTER eller lege ved ubehag.</td>
</tr>
<tr>
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</tr>
<tr>
<td>P301 + P330 + P331</td>
<td>VED SVELGING: Skyll munnen. IKKE framkall brekning.</td>
</tr>
<tr>
<td>P302 + P334</td>
<td>VED HUDKONTAKT: Skyll i kaldt vann / anvend våt kompress.</td>
</tr>
<tr>
<td>P302 + P350</td>
<td>VED HUDKONTAKT: Vask forsiktig med mye såpe og vann.</td>
</tr>
<tr>
<td>P302 + P352</td>
<td>VED HUDKONTAKT: Vask med mye såpe og vann.</td>
</tr>
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<td>P304 + P340</td>
<td>VED INNÅNDING: Flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P306 + P360</td>
<td>VED KONTAKT MED KLÆR: Skyll umiddelbart tilsølte klær og hud med mye vann før klærne fjernes.</td>
</tr>
<tr>
<td>P308 + P313</td>
<td>Ved eksponering eller mistanke om eksponering: Søk legehjelp.</td>
</tr>
<tr>
<td>P309 + P311</td>
<td>Ved eksponering eller ubehag: Kontakt et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P332 + P313</td>
<td>Ved hudirritasjon: Søk legehjelp.</td>
</tr>
<tr>
<td>P335 + P334</td>
<td>Børst bort løse partikler fra huden. Skyll i kaldt vann / anvend våt kompress.</td>
</tr>
<tr>
<td>P337 + P313</td>
<td>Ved vedvarende øyeirritasjon: Søk legehjelp.</td>
</tr>
<tr>
<td>P342 + P311</td>
<td>Ved symptomer i luftveiene: Kontakt et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P370 + P376</td>
<td>Ved brann: Stopp lekkasje dersom dette kan gjøres på en sikker måte.</td>
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<tr>
<td>P370 + P378</td>
<td>Ved brann: Slukk med …</td>
</tr>
<tr>
<td>P401</td>
<td>Oppbevares …</td>
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<tr>
<td>P402</td>
<td>Oppbevares tørt.</td>
</tr>
<tr>
<td>P403</td>
<td>Oppbevares på et godt ventilert sted.</td>
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<tr>
<td>P404</td>
<td>Oppbevares i lukket beholder.</td>
</tr>
<tr>
<td>P405</td>
<td>Oppbevares innelåst.</td>
</tr>
<tr>
<td>P406</td>
<td>Oppbevares i korrosjonsbestandig/… beholder med korrosjonsbestandig indre belegg.</td>
</tr>
<tr>
<td>P407</td>
<td>Se til at det er luft mellom stabler/paller.</td>
</tr>
<tr>
<td>P410</td>
<td>Beskyttes mot sollys.</td>
</tr>
<tr>
<td>P411</td>
<td>Oppbevares ved en temperatur som ikke er høyere enn …°C /… °F.</td>
</tr>
<tr>
<td>P412</td>
<td>Må ikke utsettes for temperaturer høyere enn 50 °C /122 °F.</td>
</tr>
<tr>
<td>P413</td>
<td>Bulkmengder på over …kg/…lbs oppbevares ved en temperatur som ikke er høyere enn …°C /… °F.</td>
</tr>
<tr>
<td>P420</td>
<td>Må oppbevares adskilt fra andre materialer.</td>
</tr>
<tr>
<td>P422</td>
<td>Oppbevar innholdet under …</td>
</tr>
<tr>
<td>P402 + P404</td>
<td>Oppbevares tørt. Oppbevares i lukket beholder.</td>
</tr>
<tr>
<td>P403 + P233</td>
<td>Oppbevares på et godt ventilert sted. Hold beholderen tett lukket.</td>
</tr>
<tr>
<td>P403 + P235</td>
<td>Oppbevares på et godt ventilert sted. Oppbevares kjølig.</td>
</tr>
<tr>
<td>P410 + P403</td>
<td>Beskyttes mot sollys. Oppbevares på et godt ventilert sted.</td>
</tr>
<tr>
<td>P410 + P412</td>
<td>Beskyttes mot sollys. Må ikke utsettes for temperaturer høyere enn 50 °C /122 °F.</td>
</tr>
<tr>
<td>P411 + P235</td>
<td>Oppbevares ved en temperatur som ikke er høyere enn …°C /… °F. Oppbevares kjølig.</td>
</tr>
<tr>
<td>P501</td>
<td>Innhold/beholder leveres til …</td>
</tr>
</tbody>
</table>