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.


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 Community 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 EC 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 Community 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 Community 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 for Medicinal Products for Veterinary Use (CVMP), the Committee on Orphan Medicinal Products (COMP), the Paediatric Committee [1], the Pharmacovigilance Risk Assessment Committee (PRAC) [1] and the Committee on Herbal Medicinal Products (HMPC). The detailed arrangements of participation for the representatives of EFTA States shall be in accordance with the provisions of Title IV, chapter 1, of Regulation (EC) No 726/2004 of the European Parliament and of the Council. Such representatives shall, 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 an EC Member State.


The EFTA States shall fully participate 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 marketing authorisation granted for a medicinal product following an opinion adopted by the competent EMEA scientific Committee in accordance with Article 9 or Article 34 of Regulation (EC) No 726/2004 of the European Parliament and of the Council shall not be subject to any fees other than those referred to in Article 67(3) and Article 70 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 apply to the Agency the Protocol of Privileges and Immunities of the European Communities.


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

1.\(^{(1)}\) [ ]
2.\(^{(1)}\) [ ]
3.\(^{(1)}\) [ ]
5.\(^{(1)}\) [ ]
6.\(^{(1)}\) [ ]
10.\(^{(1)}\) [ ]
11.\(^{(1)}\) [ ]


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.


12d.\textsuperscript{18} Commission Implementing Regulation (EU) 2018/470 of 21 March 2018 on detailed rules on the maximum residue limit to be considered for control purposes for foodstuffs derived from animals which have been treated in the EU under Article 11 of Directive 2001/82/EC (OJ L 79, 22.3.2018, p. 16).


\textsuperscript{17} Point inserted by Decision No 236/2018 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 6.12.2018.

\textsuperscript{18} Point inserted by Decision No 61/2019 (OJ L 210, 2.7.2020, p. 28 and EEA Supplement No 44, 2.7.2020, p. 30), c.i.f. 30.3.2019.


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*(f)* 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.

*(g)* 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.

*(h)* 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.

*(i)* 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.

*(j)* 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.


(61) Indent added by Decision No 50/2015 (OJ L 129, 19.5.2016, p. 31 and EEA Supplement No 29, 19.5.2016, p. 32), e.i.f. 21.3.2015.


(66) 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.


(72) 32015 R 1079: Commission Implementing Regulation (EU) 2015/1079 of 3 July 2015 (OJ L 175, 4.7.2015, p. 8),

(73) 32015 R 1080: Commission Implementing Regulation (EU) 2015/1080 of 3 July 2015 (OJ L 175, 4.7.2015, p. 11),


(83) 32016 R 0710: Commission Implementing Regulation (EU) 2016/710 of 12 May 2016 (OJ L 125, 13.5.2016, p. 6),


(82) 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.

\(^{(13)}\) Indent added by Decision No 237/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.12.2018.
\(^{(14)}\) Indent added by Decision No 237/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.12.2018.
\(^{(15)}\) 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.
\(^{(16)}\) 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.
\(^{(17)}\) Indent added by Decision No 63/2019 (OJ L 210, 2.7.2020, p. 30 and EEA Supplement No 44, 2.7.2020, p. 33), e.i.f. 30.3.2019.
\(^{(18)}\) Indent added by Decision No 115/2019 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 1.6.2019.
\(^{(19)}\) Indent added by Decision No 224/2019 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 28.9.2019.

15b. [105]

15c. [103]

15d. [104]

15e. [105]

15f. [106]

15g. [106]


Footnotes:


151.\{121\} 395 R 0540: Commission Regulation (EC) No 540/95 of 10 March 1995 laying down the arrangements for reporting suspected unexpected adverse reactions which are not serious, whether arising in the Community or in a third country, to medicinal products for human or veterinary use authorised in accordance with the provisions of Council Regulation (EEC) No 2309/93 (OJ L 55, 11.3.1995, p. 5).

15j. [ ] \{122\}

15k. [ ] \{123\}


\{128\}The provisions of the Regulation shall, for the purposes of the present Agreement, be read with the following adaptation:

\{112\} Indent added by Decision No 72/2011 (OJ L 262, 6.10.2011, p. 28 and EEA Supplement No 54, 6.10.2011, p. 38), e.i.f. 2.7.2011.


\{117\} 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.


\{120\} Indent added by Decision No 141/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.


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. (127) 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. [ ] (129)


(135) The transitional arrangements set out in the Annexes to the Act of Accession of 16 April 2003 for Lithuania (Annex IX, Chapter 1, Point 1) and Poland (Annex XII, Chapter 1, Point 4), shall apply.

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

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Liechtenstein shall not be obliged to participate in the decentralized 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.


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\[154\] For Liechtenstein, Decision 61/2009 entered into force 1.12.2010 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).


\[156\] Indent added 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 1.12.2010 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).

\[157\] Indent added by Decision No 72/2011 (OJ L 262, 6.10.2011, p. 28 and EEA Supplement No 54, 6.10.2011, p. 38), e.i.f. 2.7.2011.

\[158\] Indent added by Decision No 72/2011 (OJ L 262, 6.10.2011, p. 28 and EEA Supplement No 54, 6.10.2011, p. 38), e.i.f. 2.7.2011.

\[159\] 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), e.i.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;
- 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.[156] 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:


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[157] Indent 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.
[161] Indent added by Decision No 102/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.7.2020.

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 Inspektionein).


15r. \[164\]

15s. \[165\]


\[165\] Point inserted by Decision No 116/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.


\[165\] For Liechtenstein, Decision 128/2009 entered into force 1.12.2010 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).


\[165\] For Liechtenstein, Decision 128/2009 entered into force 1.12.2010 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).


\[165\] For Liechtenstein, Decision 128/2009 entered into force 1.12.2010 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).


\[170\] 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


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.


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.


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.


\(^{(186)}\) Indent and words “,as amended by:” added by Decision No 14/2014 (OJ L 211, 17.7.2014, p. 21 and EEA Supplement No 42, 17.7.2014, p. 19), e.i.f. 15.2.2014.


For Liechtenstein, Decision 61/2009 entered into force 1.12.2010 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). The adaptation text is replaced by Decision No 92/2017 (OJ L 36, 7.2.2017, p. 41 and EEA Supplement No 11, 7.2.2019, p. 49), e.i.f. 1.6.2018.


For Liechtenstein, Decision 128/2009 entered into force 1.12.2010 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).


\(^{(190)}\) 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), e.i.f. 28.5.2014.
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 84(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.


[196] Indent added by Decision No 163/2013 (OJ L 58, 27.2.2014, p. 16 and EEA Supplement No 13, 27.2.2014, p. 18), e.i.f. 28.5.2014.

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

The EFSA States shall be fully associated with the work of the Committee for Advanced Therapy, but without the right to vote.


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 EFSA Surveillance Authority in close cooperation with the Commission. Before the EFSA 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 Directive shall, for the purposes of this Agreement, be read with the following adaptation:

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


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, with the exception of Articles 15 and 16. The definitions in Article 3 and general provisions in Articles 17(2)(b) 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.{25}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).


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.


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Footnotes:

[222] Indent and words „, as amended by:“ added by Decision No 87/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
[223] Indent added by Decision No 239/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.12.2018.


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.


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

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


2.\(^{[229]}\) **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\(^{[231]}\)**

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\(^{[228]}\) Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 3.2.2017.

\(^{[229]}\) Point inserted by Decision No 88/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.4.2018.


\(^{[228]}\) Point inserted by Decision No 114/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.8.2020.


ACTS REFERRED TO


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

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[244] Indent added by Decision No 51/2015 (OJ L 129, 19.5.2016, p. 32 and EEA Supplement No 29, 19.5.2016, p. 33), e.i.f. 21.3.2015.


[246] Indent added by Decision No 75/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.
(a) The EFTA States will be free to limit access to their market according to the requirements of their legislation existing at the date of entry into force of this Agreement concerning cadmium in fertilizers. The Contracting Parties shall jointly review the situation in 2009.

(b) In Annex I, A.2, the following shall be added to the text in brackets of No. 1, column 6, third paragraph:

“Iceland, Liechtenstein, Norway”.

(c) In Annex I, B.1.1, B.2.1 and B.4, the following shall be added to the text in brackets after (6b) in column 5, point 3, second paragraph first indent:

“Iceland, Liechtenstein, Norway”.


XV. DANGEROUS SUBSTANCES

ACTS REFERRED TO

1. [ ] [249]
2. [ ] [249]
3. [ ] [249]
4. [ ] [249]


5. [ ]
6. [ ]
7. [ ]


12f. [ ]


12j. [ ]


\(^{275}\) 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), c.i.f. pending; it shall apply from 9.7.2014.


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


\textsuperscript{277} Point 12n (Directive 98/8/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, 22.5.2013, 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.

\textsuperscript{278} Indent and words ‘as amended by’ added by Decision No 15/2014 (OJ L 211, 17.7.2014, p. 22 and EEA Supplement No 42, 17.7.2014, p. 20), e.i.f. 1.6.2014.


\textsuperscript{280} Indent added by Decision No 218/2014 (OJ L 236, 3.9.2015, p. 16 and EEA Supplement No 52, 3.9.2015, p. 16), e.i.f. 1.11.2014.

\textsuperscript{281} Indent added by Decision No 8/2020 (OJ L 230, 3.9.2015, p. 16 and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{282} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{283} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{284} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{285} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{286} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\textsuperscript{287} Indent added by Decision No 8/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.
(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.


12nnd.[(38)] 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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[35] Indent added by Decision No 117/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2019.
[36] Indent added by Decision No 152/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[38] Point inserted by Decision No 52/2015 (OJ L 129, 19.5.2016, p. 33 and EEA Supplement No 29, 19.5.2016, p. 34), e.i.f. 21.3.2015.


\(^{[28]}\) Commission Implementing Decision (EU) 2015/1751 of 29 September 2015 on the terms and conditions of the authorisation of a biocidal product containing bromadiolone referred by the United Kingdom in accordance with Article 36 of Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 265, 1.10.2015, p. 15).


- [ ] [346]

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:

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(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.


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[^388]: Indent added by Decision No 210/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.10.2018.
[^389]: Indent added by Decision No 210/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.10.2018.
[^390]: Indent added by Decision No 210/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.10.2018.
[^391]: Indent added by Decision No 210/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.10.2018.
[^392]: Indent added by Decision No 210/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.10.2018.
[^393]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^394]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^395]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^396]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^397]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^398]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^399]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^400]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
[^401]: Indent added by Decision No 153/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
12r. [ ] [411]

12s. [ ] [412]


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[404] Indent added by Decision No 9/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

[405] Indent added by Decision No 9/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

[406] Indent added by Decision No 76/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.

[407] Indent added by Decision No 76/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.

[408] Indent added by Decision No 76/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.

[409] Indent added by Decision No 76/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.

[410] Indent added by Decision No 76/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.


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

\[418\] 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.
\[425\] Indent added by Decision No 90/2015 (OJ L 211, 4.8.2016, p. 27 and EEA Supplement No 42, 4.8.2016, p. 26), e.i.f. 1.5.2015.
\[429\] Indent added by Decision No 225/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.
Notwithstanding the provisions of Protocol 1 to the Agreement, the term ‘customs territory of the Community’ contained in Article 2 shall be understood to include the territory of the EFTA States.


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.


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\(^{440}\) Indent and words "as, amended by:"


\(^{442}\) Indent added by Decision No 216/2015 (OJ L 85, 30.3.2017, p. 36 and EEA Supplement No 19, 30.3.2017, p. 35), c.i.f. 1.11.2015.


\(^{448}\) Indent and words "as, amended by:"


\[\text{\footnotesize \textsuperscript{(44)}}\] Indent added by Decision No 8/2010 (OJ L 19, 22.4.2010, p. 16 and EEA Supplement No19, 22.4.2010, p. 16), e.i.f. 30.1.2010.


\[\text{\footnotesize \textsuperscript{(46)}}\] Indent added by Decision No 74/2011 (OJ L 262, 6.10.2011, p. 31 and EEA Supplement No 54, 6.10.2011, p. 42), e.i.f. 2.7.2011.

\[\text{\footnotesize \textsuperscript{(47)}}\] Indent added by Decision No 74/2011 (OJ L 262, 6.10.2011, p. 31 and EEA Supplement No 54, 6.10.2011, p. 42), e.i.f. 2.7.2011.


\[\text{\footnotesize \textsuperscript{(51)}}\] Indent added by Decision No 146/2012 (OJ L 270, 4.10.2012, p. 6 and EEA Supplement No 56, 4.10.2012, p. 8), e.i.f. 1.2.2013.

\[\text{\footnotesize \textsuperscript{(52)}}\] Indent added by Decision No 146/2012 (OJ L 270, 4.10.2012, p. 6 and EEA Supplement No 56, 4.10.2012, p. 8), e.i.f. 1.2.2013.

\[\text{\footnotesize \textsuperscript{(53)}}\] Indent 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.

\[\text{\footnotesize \textsuperscript{(54)}}\] Indent added by Decision No 125/2012 (OJ L 309, 8.11.2012, p. 3 and EEA Supplement No 63, 8.11.2012, p. 4), e.i.f. 14.7.2012.


\[\text{\footnotesize \textsuperscript{(57)}}\] Indent added by Decision No 195/2012 (OJ L 21, 24.1.2013, p. 43 and EEA Supplement No 6, 24.1.2013, p. 9), e.i.f. 1.1.2012.

\[\text{\footnotesize \textsuperscript{(58)}}\] Indent added by Decision No 70/2013 (OJ L 291, 31.10.2013, p. 27 and EEA Supplement No 61, 31.10.2013, p. 31), e.i.f. 4.5.2013.

\[\text{\footnotesize \textsuperscript{(59)}}\] Indent added by Decision No 71/2013 (OJ L 291, 31.10.2013, p. 28 and EEA Supplement No 61, 31.10.2013, p. 32), e.i.f. 4.5.2013.

\[\text{\footnotesize \textsuperscript{(60)}}\] Indent added by Decision No 72/2013 (OJ L 291, 31.10.2013, p. 29 and EEA Supplement No 61, 31.10.2013, p. 33), e.i.f. 4.5.2013.


[footnotes]


[457] Indent added by Decision No 192/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.

[458] Indent added by Decision No 193/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.


[461] 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.


- [ ]


\[\text{(473)}\]

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\[\text{(499)}\]
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

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490 Indent added by Decision No 66/2019 (OJ L 210, 2.7.2020, p. 32 and EEA Supplement No 44, 2.7.2020, p. 36), e.i.f. 30.3.2019.
492 Indent added by Decision No 30/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
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:

“Nationals 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.”


12ze. [ ] 495


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12zt. [ ]


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*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.*

*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.*

*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.*


*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.*

*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.*


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 (containing 0.01% ≤ acrylamide &lt; 0.1%)</td>
<td>77402-05-2</td>
<td>[NOR-UNN-02-91]</td>
<td>403-230-3</td>
</tr>
<tr>
<td>methyl acrylamidomethoxyacetate (containing 0.01% ≤ acrylamide &lt; 0.1%)</td>
<td>77402-03-0</td>
<td>[NOR-UNN-03-01]</td>
<td>401-890-7</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.

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{544} Indent added by Decision No 180/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.

{545} Indent added by Decision No 67/2019 (OJ L 210, 2.7.2020, p. 33 and EEA Supplement No 44, 2.7.2020, p. 37), e.i.f. 30.3.2019.

{546} Indent added by Decision No 226/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.

{547} Indent added by Decision No 103/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.7.2020.
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.


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

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

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


12zzzf.\(^{[87]}\) **32016 D 0109**: Commission Implementing Decision (EU) 2016/109 of 27 January 2016 not to approve PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-types 1, 6 and 9 (OJ L 21, 28.1.2016, p. 84).


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**Points inserted:**


\(^{[86]}\) 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.

\(^{[87]}\) 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.


\(^{[89]}\) 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.

\(^{[90]}\) 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.


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[(67)] Point inserted by Decision No 182/2018 (OJ L [to be published] and EEA Supplement No [to be published]), c.f. 22.9.2018.

[(68)] Point inserted by Decision No 182/2018 (OJ L [to be published] and EEA Supplement No [to be published]), c.f. 22.9.2018.

[(69)] Point inserted by Decision No 182/2018 (OJ L [to be published] and EEA Supplement No [to be published]), c.f. 22.9.2018.


[(75)] Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 13.6.2019.


[^47]: Point inserted by Decision No 118/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 1.6.2019.
[^48]: Point inserted by Decision No 154/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 15.6.2019.
[^49]: Point inserted by Decision No 227/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 28.9.2019.
[^50]: Point inserted by Decision No 227/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 28.9.2019.
[^51]: Point inserted by Decision No 228/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 28.9.2019.
[^52]: Point inserted by Decision No 228/2019 (OJ L [to be published] and EEA supplement No [to be published], e.i.f. 28.9.2019.
[^54]: Point inserted by Decision No 11/2020 (OJ L [to be published] and EEA Supplement No [to be published], e.i.f. 8.2.2020.
[^55]: Point inserted by Decision No 60/2020 (OJ L [to be published] and EEA Supplement No [to be published], e.i.f. 1.5.2020.


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.”

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

“Iceland, Norway”

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

“Liechtenstein”


\(^{66}\) Point and indent 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.

\(^{66}\) 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.

\(^{66}\) 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.

\(^{66}\) 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.

\(^{66}\) 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.


Notes:

- 32011 R 0704: 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.
- 32011 R 0705: 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.
- 32011 R 0706: 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.
- 32011 R 0736: 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.
- 32011 R 0740: 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.
- 32011 R 0786: 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.
- 32011 R 0787: 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.
- 32011 R 0788: 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.


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


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

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

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

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

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

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

\[\text{Indent added by Decision No 205/2014 (OJ L 205, 30.7.2015, p. 64 and EEA Supplement No 43, 30.7.2015, p. 63), e.i.f. 1.6.2015 and subsequently before publication by Corrigendum of 20.3.2015.}\]

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

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

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

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

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

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


(745) 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.

(746) 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.

(747) 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.

(748) 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.

(749) 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.

(750) 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.

(751) 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.


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

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

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

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

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

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

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

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

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


\[45\] 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.

\[46\] 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.

\[47\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[48\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[49\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[50\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[51\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[52\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[53\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[54\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[55\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[56\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[57\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[58\] Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.

\[\text{Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]
\[\text{Indent added by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]


(785) Indent added by Decision No 270/2014 (OJ L 311, 26.11.2015, p. 21 and EEA Supplement No 71, 26.11.2015, p. 20), e.i.f. 1.6.2015.

(786) Indent added by Decision No 270/2014 (OJ L 311, 26.11.2015, p. 21 and EEA Supplement No 71, 26.11.2015, p. 20), e.i.f. 1.6.2015.


(789) Indent added by Decision No 271/2014 (OJ L 311, 26.11.2015, p. 22 and EEA Supplement No 71, 26.11.2015, p. 21), e.i.f. 1.6.2015.


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

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


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.

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.


(296) 32015 R 2047: Commission Implementing Regulation (EU) 2015/2047 of 16 November 2015 (OJ L 300, 17.11.2015, p. 8),


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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), e.i.f. 24.3.2018.

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), e.i.f. 24.3.2018.

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), e.i.f. 24.3.2018.

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), e.i.f. 24.3.2018.

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), e.i.f. 24.3.2018.


(925) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(926) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(927) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(928) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(929) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(930) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(931) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(932) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(933) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(934) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(935) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(936) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(937) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(938) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(939) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(940) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(941) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(942) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(943) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(944) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(945) Indent added by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.


- (33) 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.
- (34) 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.
- (37) Indent added by Decision No 119/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2019.
- (38) Indent added by Decision No 120/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2019.
- (39) Indent added by Decision No 120/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2019.

(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
(2018) Indent added by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.


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

\(^{*}^*\) Indent added by Decision No 32/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 35/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

\(^{*}^*\) Indent added by Decision No 77/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.

\(^{*}^*\) Indent added by Decision No 77/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.
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.

13a.\[^{[84]}\] 32012 R 0823: Commission Regulation (EU) No 823/2012 of 14 September 2012 derogating from Implementing Regulation (EU) No 540/2011 as regards the expiry dates of the approval of the active substances 2,4-DB, benzoic acid, beta-cyfluthrin, carfentrazone ethyl, Coniothyrium minitans Strain CON/M/91-08 (DSM 9660), cyzofamid, cyfluthrin, deltamethrin, dimethenamid-P, ethofumesate, ethoxyfural, fenamidone, flazasulfuron, flufenacet, flurtamone, foramsulfuron, fosthiazate, imazamox, iodosulfuron, iprodione, isoxaflutole, linuron, maleic hydrazide, mecoprop, mecoprop-P, mesosulfuron, mesotrione, oxadiargyl, oxasulfuron, pendimethalin, picoxystrobin, propiconazole, propineb, propoxycarbazine, propyzamide, pyraclostrobin, silthiofam, trifloxystrobin, warfarin and zoxamide (OJ L 250, 15.9.2012, p. 13), as amended by:


\[^{[84]}\] 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. 1.6.2015.


\[^{[92]}\] Indent and words “, as amended by:” added by Decision No 186/2018 (OJ L [to be published] and EEA Supplement No [to be published]), c.f. 22.9.2018.
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: Eitrað 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 ljósnaðingu.

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 brennir 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:

“IS: Mengið ekki vatn með efninu eða íláti þess. (Hreinsið ekki búnað nálægt yfirboðsvatni/Koma skal í veg fyrir að mengun verði með afremsli frá þeinarhliðum og vegum.)

NO: Unngå forurensning av vannmiljøet med produktet eller emballasjen. (Ikke rengjør spredeutstyr nær overflatevann/unngå forurensning via avrenning fra gårdsplasser og veler).”

(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 húð 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: Ívöði allan hlífðarfatnnað að lokinni notkun.

NO: Vask alt personlig 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: Forðist innöndun reyks eftir að hver ljósnað þegar í stað svæðið sem er til meðhöndlunar.

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

(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 ílátið utanhús og við þurr skilyrði.

NO: Beholderen skal åpnes utendørs og under tørre 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: Loftvæsta skal úðuð svæði/gróðurhús (vandlega/eða í tilgreindan tíma/þar til úðinn hefur þornað) aður en farið er þangað inn aftur.

NO: De behandlede områder/veksthus ventileres (grundig/eller angivelse av tid/inntil produktet har tørt) 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 vernda grunnvatn/jarðvægslífverur skal ekki nota þetta eða annað efnir sem inniheldur (tílgreinið virkt efnir eða flokk virkra efnir eftir þvi sem við a) lengur eða oftaer en (tílgreinið hversu lengi eða oft má nota efníð).

NO: For å beskytte (grunnvannet/jordlevende organismer) 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 vernda grunnvatn/vatnslífverur skal ekki nota þetta efnir (á tilgreinda jarðvægserð eða við tilgreindar aðstæðar).

NO: For å beskytte (grunnvannet/vannlevende organismer) må dette produktet ikke brukes (på beskrevet jordtype eller under beskrevne forhold)."

(l) The following shall be added to the list under the title “SPE 3” in point 2.2 of Annex III:

"IS: Til að vernda vatnslífverur/plöntur utan markhóps/löðþyr utan markhóps/skordyr má ekki nota efníð nær öræktuð landi/yfirborðsvatni en (tílgreind breidd svæðís sem er óheimilt að úða).

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

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

"IS: Til að vernda vatnslífverur/plöntur utan markhóps má ekki nota efníð á malbikað, steinsteyp, hellulaðið eða malarborð þjóðborð eða vegi (jarnbrautarspor) eða önnur svæði þar sem hætt er við afrensnið í umhverfið.

NO: For å beskytte (vanlevende organismer/viltlevende planter) må dette produktet ikke brukes på harde overflater som asphalt, betong- brostein- eller gruslagte områder og veier/jernbane, eller på andre områder med stor risiko for avrenning)."

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

"IS: Til að vernda fugla/viltl. spendýr verður að geta þess vandlega að efníð sé algerlega hulið jarðvegi; getið þess séstaklega að efníð sé hulið í endum raða.

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

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

"IS: Hreinsl. upp allt efní, sem hefur farið til spillis, til að vernda fugla/viltl. spendýr.

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

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

"IS: Óheimilt er að nota efníð á varftpíma fugla.
The following shall be added to the list under the title “SPe 8” in point 2.2 of Annex III:

“IS: Hættulegt frævandi skordýrum/Til að vernda býflugur og önnur fæðuleit/Fjarlægtið bykúpur meðan meðhöndlun með efninu fer fram eða hyljið þær á meðan og í (tilgreinið tíma) að lokinni meðhöndlun/Öheimilt er að nota efninn ef blómstrandi illgresi er til staðar/Öheimilt er að nota efninn fyrrir (tilgreinið tíma).”

NO: Farlig for bier./For å beskytte bier 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 (nevn antall timer/dager) ett er behandlingen./Må ikke brukes i nærheten av blomstrende ugress./Fjern ugresset før det bloms trer./Må ikke brukes før (tidspunkt).”

The following shall be added to the list under the title “SPr 1” in point 2.4 of Annex III:

“IS: Beitu skal komið fyrir þannig að ekki sé hætt á að önnur dýr komist í hana. Festu skal beituna tryggilega þannig að nagðýr geti ekki dreigið hana í burtu.

NO: Produktet skal plasseres på en slik måte at risikoen for at andre dyr kan innta produktet minimeres. Pass på at produkt i blokkform ikke kan flyttes vekk 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 meðhöndla á, meðan á meðhöndlun stendur. Varað skal við hættunni á að verða fyrir eitrun (beinni eða óbeinni) af völdum 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:

“IS: Hræ nagðýra skulu fjarlægð daglega af meðhöndlalaða svæðinu meðan meðhöndlun stendur yfir. Ekki skal setja hræin í opin soppilat.

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


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[999] Indent and words”, as amended by:” added by Decision No 121/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.ï.f. 1.6.2019.


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13zr. Commission Implementing Decision 2012/677/EU of 30 October 2012 allowing Member States to extend provisional authorisations granted for the new active substances ametocrina (initially applied for under the development code BAS 650 F) and disodium phosphonate (OJ L 305, 1.11.2012, p. 27).

\[1825\] 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.

\[1826\] 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.

\[1827\] 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.

\[1828\] 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.

\[1829\] 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.

\[1830\] 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.

\[1831\] 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.

\[1832\] 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.

\[1833\] 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.

\[1834\] 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.


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**32015** | 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), e.i.f. 1.6.2015.

**32018** | Indent and words “32015 R 1106:” added by Decision No 264/2015 (OJ L [to be published]) and EEA Supplement No [to be published], e.i.f. 1.11.2015.

**32017** | 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), e.i.f. 1.6.2015.

**32018** | 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), e.i.f. 1.6.2015.

**32019** | 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), e.i.f. 1.6.2015.

**32020** | 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), e.i.f. 1.6.2015.

**32021** | Point inserted by Decision No 34/2020 (OJ L [to be published]) and EEA Supplement No [to be published], e.i.f. 21.3.2020.

**32040** | 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), e.i.f. 1.6.2015.

**32041** | 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), e.i.f. 1.6.2015.


13zzz[\textsuperscript{1075}] 32012 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:


13zzz[\textsuperscript{1084}] 32013 D 0038: Commission Implementing Decision 2013/38/EU of 18 January 2013 allowing Member States to extend provisional authorisations granted for the new active substances emamectin and maltodextrin (OJ L 18, 22.1.2013, p. 17).


13zzzh.[\textsuperscript{1086}] 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 acacenocyl, aminopyralid, ascorbic acid, flubeniamide, gamma-cyhalothrin, iponazole, metaflumizone, orthosulfamuron, Pseudomonas sp. strain DSMZ 13134, pyridalil, pyroxsulam, spiromesifen, thiencarbazone and topramezone (OJ L 117, 27.4.2013, p. 20).

13zzzi.[\textsuperscript{1087}] 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:

\[\textsuperscript{1075}\text{Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1076}\text{Indent and words "as, amended by:" added by Decision No 273/2014 (OJ L 311, 26.11.2015, p. 26 and EEA Supplement No 71, 26.11.2015, p. 25), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1077}\text{Indent added by Decision No 95/2015 (OJ L 211, 4.8.2016, p. 33 and EEA Supplement No 42, 4.8.2016, p. 32), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1078}\text{Indent added by Decision No 149/2016 (OJ L 73, 15.3.2018, p. 22 and EEA Supplement No 16, 15.3.2018, p. 25), e.i.f. 9.7.2016.}\]
\[\textsuperscript{1079}\text{Indent added by Decision No 133/2017 (OJ L 128, 16.5.2019, p. 26 and EEA Supplement No 40, 16.5.2019, p. 27), e.i.f. 8.7.2017.}\]
\[\textsuperscript{1080}\text{Indent added by Decision No 144/2017 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.}\]
\[\textsuperscript{1081}\text{Indent added by Decision No 229/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.}\]
\[\textsuperscript{1082}\text{Indent added by Decision No 297/2019 (OJ L 68, 5.3.2020, p. 35 and EEA Supplement No 14, 5.3.2020, p. 40), e.i.f. 14.12.2019.}\]
\[\textsuperscript{1083}\text{Indent added by Decision No 298/2019 (OJ L 68, 5.3.2020, p. 38 and EEA Supplement No 14, 5.3.2020, p. 43), e.i.f. 14.12.2019.}\]
\[\textsuperscript{1084}\text{Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1085}\text{Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1086}\text{Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]
\[\textsuperscript{1087}\text{Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]


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\[\text{(188) Indent and words “, as amended by:” inserted by Decision No 119/2019 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 1.6.2019.}\]

\[\text{(189) Indent added by Decision No 119/2019 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 1.6.2019.}\]

\[\text{(190) Indent added by Decision No 119/2019 (OJ L [to be published] and EEA supplement No [to be published]), e.i.f. 1.6.2019.}\]

\[\text{(191) Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]

\[\text{(192) Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]

\[\text{(193) Point inserted by Decision No 206/2014 (OJ L 202, 30.7.2015, p. 87 and EEA Supplement No 43, 30.7.2015, p. 86), e.i.f. 1.6.2015.}\]

\[\text{(194) Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]

\[\text{(195) Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]

\[\text{(196) Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]

\[\text{(197) Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2015.}\]

\[\text{(198) Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]

\[\text{(199) Point inserted by Decision No 207/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.}\]


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\(^{1119}\) Point inserted by Decision No 274/2014 (OJ L 311, 26.11.2015, p. 27 and EEA Supplement No 71, 26.11.2015, p. 26), c.i.f. 1.6.2015.


\(^{1121}\) Point inserted by Decision No 95/2015 (OJ L 211, 4.8.2016, p. 33 and EEA Supplement No 42, 4.8.2016, p. 32), c.i.f. 1.6.2015.


\(^{1123}\) Point inserted by Decision No 95/2015 (OJ L 211, 4.8.2016, p. 33 and EEA Supplement No 42, 4.8.2016, p. 32), c.i.f. 1.6.2015.

\(^{1124}\) Point inserted by Decision No 274/2014 (OJ L 311, 26.11.2015, p. 27 and EEA Supplement No 71, 26.11.2015, p. 26), c.i.f. 1.6.2015.

\(^{1125}\) Point inserted by Decision No 183/2015 (OJ L 8, 12.1.2017, p. 12 and EEA Supplement No 3, 12.1.2017, p. 11), c.i.f. 11.7.2015.

\(^{1126}\) Point inserted by Decision No 183/2015 (OJ L 8, 12.1.2017, p. 12 and EEA Supplement No 3, 12.1.2017, p. 11), c.i.f. 11.7.2015.

Council concerning the placing of plant protection products on the market and establishing a list of candidates for substitution (OJ L 67, 12.3.2015, p. 18), as amended by:


{1128} Indent and words “as amended by” added by Decision No 89/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.4.2018.

{1129} Indent added by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.

{1130} Point inserted by Decision No 262/2015 (OJ L 161, 22.6.2017, p. 44 and EEA Supplement No 38, 22.6.2017, p. 44), e.i.f. 1.11.2015.


{1132} 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.

{1133} 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.

{1134} Point inserted by Decision No 266/2015 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.11.2015.

{1135} Point inserted by Decision No 267/2015 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.11.2015.


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**Notes:**


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13zzzzzzzzd.\[1218]\textbf{32017 R 1531}: Commission Implementing Regulation (EU) 2017/1531 of 7 September 2017 renewing the approval of the active substance imazamox, as a candidate for substitution, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of

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\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.
\[129\] Point inserted by Decision No 89/2018 (OJ L [to be published]) and EEA Supplement No [to be published]), e.i.f. 28.4.2018.


^[128] Point inserted by Decision No 144/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

^[129] Point inserted by Decision No 184/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.

^[129] Point inserted by Decision No 184/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.


Point inserted by Decision No 120/2019 (OJ L [to be published] and EEA supplement No [to be published]), c.f.f. 1.6.2019.

Point inserted by Decision No 120/2019 (OJ L [to be published] and EEA supplement No [to be published]), c.f.f. 1.6.2019.

Point inserted by Decision No 120/2019 (OJ L [to be published] and EEA supplement No [to be published]), c.f.f. 1.6.2019.


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[124] Point inserted by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[127] Point inserted by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[128] Point inserted by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[129] Point inserted by Decision No 120/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[125] Point inserted by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[122] Point inserted by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[125] Point inserted by Decision No 121/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.
[124] Point inserted by Decision No 122/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 1.6.2019.


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{1255} Point inserted by Decision No 155/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 15.6.2019.
{1256} Point inserted by Decision No 229/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 28.9.2019.
{1257} Point inserted by Decision No 229/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 28.9.2019.
{1258} Point inserted by Decision No 229/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 28.9.2019.
{1259} Point inserted by Decision No 229/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 28.9.2019.
{1260} Point inserted by Decision No 229/2019 (OJ L [to be published]) and EEA supplement No [to be published]), e.i.f. 28.9.2019.


\(^{(128)}\) Point inserted by Decision No 32/2020 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 21.3.2020.

\(^{(129)}\) Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 21.3.2020.

\(^{(130)}\) Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 21.3.2020.

\(^{(131)}\) Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 21.3.2020.


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1272 Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1273 Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1274 Point inserted by Decision No 33/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1275 Point inserted by Decision No 34/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1276 Point inserted by Decision No 34/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1277 Point inserted by Decision No 35/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
1278 Point inserted by Decision No 77/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.
1279 Point inserted by Decision No 77/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2020.


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

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

1. \[1\](1284) \[1\](1285)\[1\]  

2. \[1284\] **C/79/82/p. 3**: Communication concerning Commission Decision 81/437/EEC of 11 May 1981 laying down the criteria in accordance with which information relating to the inventory of chemical substances is supplied by the Member States to the Commission (OJ C 79, 31.3.1982, p. 3).

3. \[1285\] **C/146/90/p. 4**: Publication of the EINECS inventory (OJ C 146, 15.6.1990, p. 4).

4.\[1286\] **C/1/93/p. 3**: The European Chemical Bureau. Commission communication to the Council and the European Parliament (OJ C 1, 5.1.1993, p. 3).


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13. 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. 32006 H 0283: Commission Recommendation of 11 April 2006 on risk reduction measures for the substances: Dipentylphthalate; 3,4-Dichloroaniline; Di-isodecyl phthalate; 1,2-Benzenedicarboxylic acid, di-C9-11-branched alkyl esters, C10-rich; Di-isonylonyl phthalate; 1,2-Benzenedicarboxylic acid, di-C6-10-branched alkyl esters, C9-rich; Ethyleneendiaminetetraacetate; Methyl acetate; Monochloroacetic acid; n-Pentane; Tetrasodium ethylenediaminetetraacetate (OJ L 104, 13.4.2006, p. 45).

XVI. COSMETICS

ACTS REFERRED TO

1. [ ] 1307


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1309 Indent and words “, as amended by: “, added by Decision No 195/2013 (OJ L [to be published] and EEA Supplement No [to be published]), c.e.f. 9.11.2013.

1310 Indent added by Decision No 196/2013 (OJ L [to be published] and EEA Supplement No [to be published]), c.e.f. 9.11.2013.


\[(1333)\] Indent added by Decision No 187/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.

\[(1334)\] Indent added by Decision No 188/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.

\[(1335)\] Indent added by Decision No 123/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 16.6.2019.

\[(1336)\] Indent added by Decision No 230/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.

\[(1337)\] Indent added by Decision No 230/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.

\[(1338)\] Indent added by Decision No 231/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.

\[(1339)\] Indent added by Decision No 223/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.


\[(1341)\] Indent added by Decision No 12/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.

\[(1342)\] Indent added by Decision No 13/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 8.2.2020.


9. [ ]

10. [ ]

11. [ ]

12. [ ]
13.\{1355\} [ ]\{1356\}

ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE\{1358\}

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


\{1357\}  Point inserted by Decision No 231/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.9.2019.


XVII. ENVIRONMENT PROTECTION

ACTS REFERRED TO

1. [ ]

2. [ ]

3. [ ]

4. [ ]


6. [ ]


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

(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. [ ] 1379

6ab. [ ] 1379

6ac. [ ] 1379

6ad. [ ] 1379


6af. [ ] 1379

6ag. [ ] 1380

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1372 Indent added by Decision No 74/2019 (OJ L 210, 2.7.2020, p. 44 and EEA Supplement No 44, 2.7.2020, p. 59), e.i.f. 1.8.2020.

1373 Adaptation texts c, d, e and f added by Decision No 270/2015 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.11.2015.


6ah. [ ]

6ai. [ ]

6aj. [ ]

6ak. [ ]

6al. [ ]


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1398 Point inserted by Decision No 156/2019 (OJ L [to be published] and EEA Supplement No [to be published]), c.f.f. 15.6.2019.


(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.


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[1402] Point inserted by Decision No 37/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 21.3.2020.
[1403] Point inserted by Decision No 104/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.7.2020.


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”.


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.

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.


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”.

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{1421} Indent and words “, as amended by:” added by Decision No 40/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 28.1.2020.


{1428} Indent and words “, as amended by:” added 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.
9a. [ ]

9aa. [ ]


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 the Decision of the EEA Joint Committee No 160/2019 of 13 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 the Decision of the EEA Joint Committee No 160/2019 of 13 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 the Decision of the EEA Joint Committee No 160/2019 of 13 June 2019”.

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


9bd. \textsuperscript{[1436]} 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).


conditions for mutual recognition for the certification of companies and personnel as regards stationary fire protection systems and fire extinguishers containing certain fluorinated greenhouse gases (OJ L 92, 3.4.2008, p. 12).


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”.


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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”.

**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE** \{\[1445\]}

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


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

**ACTS REFERRED TO**

1. [ ]\{\[1446\]}
2. [ ]\{\[1447\]}
4. [ ]\{\[1448\]}
4a. [ ]\{\[1449\]}

\[1444\] 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.


4n. [ ] {1463}

4o. [ ] {1464}


4u. [ ] {1472}

4v. [ ] {1473}


4x. [ ] {1475}

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{1464} Text of point 4m (Commission Decision 96/731/EC) deleted with effect from 10 July 1998 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.


{1467} 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.

{1468} 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.

{1469} Point inserted by Decision No 62/98 (OJ L 100, 15.4.1999, p. 48 and EEA Supplement No 16, 15.4.1999, p. 86), e.i.f. 5.7.1998.

{1470} Point inserted by Decision No 62/98 (OJ L 100, 15.4.1999, p. 48 and EEA Supplement No 16, 15.4.1999, p. 86), e.i.f. 5.7.1998.

{1471} Point inserted by Decision No 62/98 (OJ L 100, 15.4.1999, p. 48 and EEA Supplement No 16, 15.4.1999, p. 86), e.i.f. 5.7.1998.


4ze. 4zf. 4zg. \[^{1482}\]


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ANNEX II – p. 264


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 justified 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).


398 D 0576: Commission Decision 98/576/EC of 16 September 1998 on a common technical regulation for the attachment requirements for terminal equipment to connect to public switched telephone networks (PSTNs) and incorporating an analogue handset function (OJ L 278, 15.10.1998, p. 40).


399 D 0303: 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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Footnotes:


4zzm.**[1517]**

4zzn.**[1518]**


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

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:

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


(4) Point and adaptation inserted by Decision No 145/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

(5) Point inserted by Decision No 158/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.
1. 32015 L 1535: Directive (EU) 2015/1535 of the European Parliament and of the Council of 9 September 2015 laying down a procedure for the provision of information in the field of technical regulations and of rules on Information Society services (OJ L 241, 17.9.2015, p. 1). 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:

“... ‘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 [1526]), 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.


- (\textsuperscript{1555}) \textbf{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),


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":

\textbf{IS Efri hluti}

N Overdel


\textsuperscript{1557} 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.
(b) In Annex I, point 1(b) the following shall be added to the list of written indications concerning "lining and sock":
IS Fóður og bindsvöli
N Förr og bindsåle

(c) in Annex I, point 1(c) the following shall be added to the list of written indications concerning "outer sole":
IS Slitsöli
N Ytersål

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

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

(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 Textílefni
N Tekstilmaterialer

(g) In Annex I, point 2(c) the following shall be added to the list of written indications concerning "all other materials":
IS Öll önnur efni
N Andre materialer

3f.  


\[\text{103 T: Act concerning the conditions of accession of the Czech Republic, the Republic of Estonia, the Republic of Cyprus, the Republic of Latvia, the Republic of Lithuania, the Republic of Hungary, the Republic of Malta, the Republic of Poland, the Republic of Slovenia and the Slovak Republic and the adjustments to the Treaties on which the European Union is founded adopted on 16 April 2003 (OJ L 236, 23.9.2003, p. 33),}\]

\[\text{194 N: Act concerning the conditions of accession of the Republic of Austria, the Republic of Finland and the Kingdom of Sweden and the adjustments to the Treaties on which the European Union is founded (OJ C 241, 29.8.1994, p. 21 as amended by OJ L 1, 1.1.1995, p. 1),}\]


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


\[\text{\{1545\} Point 3f (Decision No 3052/95/EC) inserted by Decision No 16/97 (OJ L 182, 10.7.1997, p. 49 and EEA Supplement No 29, 10.7.1997, p. 70), e.i.f. 11.12.1998 and subsequently replaced by Decision No 126/2012 (OJ L 309, 8.11.2012, p. 4 and EEA Supplement No 63, 8.11.2012, p. 5), and subsequently point 3f (Regulation (EC) 764/2008) deleted by Decision 105/2020 (OJ L \{to be published\}) and EEA Supplement No \{to be published\}, e.i.f. pending.}\]


\[\text{\{1547\} Indent and words ‘‘, as amended by:’’ above, added by the 2004 EEA Enlargement Agreement (OJ L 130, 29.4.2004, p. 3 and EEA Supplement No 23, 29.4.2004, p. 1), e.i.f. 1.5.2004.}\]


\[\text{\{1550\} Point inserted by Decision No 9/2003 (OJ L 94, 10.4.2003, p. 59 and EEA Supplement No 19, 10.4.2003, p. 12), e.i.f. 1.3.2004.}\]

\[\text{\{1551\} Indent and words ‘‘, as amended by:’’ added by Decision No 126/2012 (OJ L 309, 8.11.2012, p. 4 and EEA Supplement No 63, 8.11.2012, p. 5), e.i.f. 1.4.2013.}\]

\[\text{\{1552\} Point inserted by Decision No 7/2006 (OJ L 92, 30.3.2006, p. 28 and EEA Supplement No 17, 30.3.2006, p. 8), e.i.f. 28.1.2006.}\]


\[\text{\{1555\} Indent and words ‘‘, as amended by:’’ added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.}\]

\[\text{\{1556\} Indent and words ‘‘, as amended by:’’ added by Decision No 117/2010 (OJ L 58, 3.3.2011, p. 74 and EEA Supplement No 12, 3.3.2011, p. 14), e.i.f. 11.11.2010.}\]


3n. [1565] 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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[1557] 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.

[1558] 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.


[1565] 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.i.f. 11.11.2010.


The safety equipment requirements, as laid down in Commission Decision 2011/477/EU (OJ L 128, 16.5.2011), e.i.f., together with the following:


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.

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**[1571]** Indent added by Decision No 185/2014 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 26.9.2014.


**[1575]** Indent added by Decision No 146/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018.

**[1576]** Indent added by Decision No 159/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 15.6.2019.


**[1578]** Point inserted by Decision No 84/2014 (OJ L 310, 30.10.2014, p. 37 and EEA Supplement No 63, 30.10.2014, p. 28), e.i.f. 17.5.2014.


**[1580]** Point and adaptation inserted by Decision No 105/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.
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 in extended to Liechtenstein.

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

(d) The words “Article 36 TFEU” shall read “Article 13 of the EEA Agreement”.

(e) 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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13.\footnote{1584} \textit{96/C 224/02}: Council Resolution of 8 July 1996 on cooperation between administrations for the enforcement of legislation on the internal market (OJ C 224, 1.8.1996, p. 3).

14.\footnote{1585} \textit{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, 1.8.1996, p. 5).


XX. FREE MOVEMENT OF GOODS – GENERAL

ACTS REFERRED TO\footnote{1592}

ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

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

2. [1994] 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).


5. [1995] 1


XXL CONSTRUCTION PRODUCTS

ACTS REFERRED TO


[1578] Point inserted by Decision No 49/2010 (OJ L [to be published] and EEA Supplement No [to be published]), c.1.f. 1.5.2010.


1f. [ ]


1n. [ ] [145]


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1zr.  


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1zzn. \textsuperscript{[1667]} 32018 D 0771: Commission Delegated Decision (EU) 2018/771 of 25 January 2018 on the applicable system to assess and verify constancy of performance of anchor devices used for construction works and intended to


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.'


Decisions on certain construction products as regards wood flooring and solid wood panelling and cladding (OJ L 79, 16.3.2006, p. 27).


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2v. 32017 R 1228: Commission Delegated Regulation (EU) 2017/1228 of 20 March 2017 on the conditions for classification, without testing, of external renders 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**

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


**XXII. PERSONAL PROTECTIVE EQUIPMENT**

**ACTS REFERRED TO**


1. [ ]


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

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


**XXIII. TOYS**

**ACTS REFERRED TO**

1. [ ]


---


1700 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.

1701 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.

ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE{[en]}

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

ACTS REFERRED TO

{[en]} Indent added by Decision No 190/2018 (OJ [to be published] and EEA Supplement No [to be published]), e.i.f. 22.9.2018.
1. [ ] [\textsuperscript{1714}]

1a. [ ] [\textsuperscript{1715}]

1b. [ ] [\textsuperscript{1716}]


\textsuperscript{1717} Point inserted by Decision No 6/2007 (OJ L 209, 9.8.2007, p.11), e.i.f. 28.4.2007.

\textsuperscript{1718} 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), e.i.f. 1.6.2015.

\textsuperscript{1719} Indent added by Decision No 9/2015 (OJ L 93, 7.4.2016, p. 15 and EEA Supplement No 21, 7.4.2016, p. 13), e.i.f. 1.1.2016.


\textsuperscript{1721} Point inserted by Decision No 60/2018 (OJ L 26, 30.1.2020, p. 46 and EEA Supplement No 6, 30.1.2020, p. 37), e.i.f. 24.3.2018.


\textsuperscript{1723} Indent added by Decision No 211/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.8.2020 and subsequently corrected [before publication] by Corrigendum of 13.6.2019.

\textsuperscript{1724} Indent added by Decision No 38/2020 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.8.2020.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE{1730}

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


XXV. TOBACCO

ACTS REFERRED TO

1.[1732] [...]
2. [deleted]


The provisions of Directive (EU) 2015/1139 shall, for the purposes of this Agreement, be read with the following adaptations:

In Article 1(1) the words “20 May 2016” shall, as regards the EFTA States, read “the date of the entry into force of the EEA Joint Committee Decision incorporating Directive 2014/40/EU of the European Parliament and of the Council into the EEA Agreement”.

In Article 1(2) the words “20 May 2017” shall, as regards the EFTA States, read “one year after the date of the entry into force of the EEA Joint Committee Decision incorporating Directive 2014/40/EU of the European Parliament and of the Council into the EEA Agreement”.


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

(a) The prohibition in Article 8 shall not apply to the placing on the market in Norway of the product defined in Article 2(4). However, this derogation shall not apply to the prohibition of sales of “snus” in forms resembling food products. Furthermore, Norway shall apply an export ban on the product defined in Article 2(4) to all Contracting Parties to the present Agreement, with the exception of Sweden.


XXVI. ENERGY

ACTS REFERRED TO

1. 32014 D 1842:

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.

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. 32014 D 1842:

2. 32014 D 1842:

3. 32014 D 1842:

Footnotes:


(Footnotes continued on next page.)


- \{\textsuperscript{1757}\} I 2012 J003: Act concerning the conditions of accession of the Republic of Croatia and the adjustments to the Treaty on European Union, the Treaty on the Functioning of the European Union


\footnotesize{\textsuperscript{1758} Indent and words “as amended by;” above added by Decision No 44/2003 (OJ L 193, 31.7.2003, p. 12 and EEA Supplement No 39, 31.7.2003, p. 9), e.i.f. 17.5.2003.}

\footnotesize{\textsuperscript{1756} Indent added by Decision No 57/2017 (OJ L 305, 29.11.2018, p. 24 and EEA Supplement No 81, 29.11.2018, p. 28), e.i.f. 18.3.2017.}

\footnotesize{\textsuperscript{1755} Point, indent and adaptation text inserted by Decision No 90/2012 (OJ L 248, 13.9.2012, p. 27 and EEA Supplement No 50, 13.9.2012, p. 32), e.i.f. 1.5.2012.}

\footnotesize{\textsuperscript{1757} Indent added by Decision No 198/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.}


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

[1787] 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.
[1794] Indent added by Decision No 91/2018 (OJ L [to be published]) and EEA Supplement No [to be published], e.i.f. 28.4.2018.
(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 25, 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 4d) of Protocol 1 to the Agreement shall not apply to Chapter III of the Regulation.

(d) The following shall be added in Annex III:

<table>
<thead>
<tr>
<th>Product category</th>
<th>Geographical indication</th>
<th>Country of origin</th>
</tr>
</thead>
<tbody>
<tr>
<td>15. Vodka</td>
<td>Íslenskt Vodka/Icelandic Vodka</td>
<td>Iceland</td>
</tr>
<tr>
<td></td>
<td>Norsk Vodka/Norwegian Vodka</td>
<td>Norway</td>
</tr>
<tr>
<td></td>
<td>Norsk akevitt/Norsk Aquavit/Norsk Akvavit/Norwegian Aquavit</td>
<td>Norway</td>
</tr>
</tbody>
</table>
| Other spirit drinks | The geographical indications mentioned under this point concern products which are not defined in the Regulation. Therefore, they must be completed with the sales description “spirit drink”.

The EFTA States producing these spirit drinks shall inform the other Contracting Parties of the national definitions of these products. |


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.

\footnote{1771} Point inserted by Decision No 223/2014 (OJ L 230, 3.9.2015, p. 25 and EEA Supplement No 52, 3.9.2015, p. 25), e.i.f. 1.11.2014.


10. [ ] \{(1773)\}


**XXVIII. CULTURAL GOODS \{(1775)\}**

**ACTS REFERRED TO**

1. [ ] \{(1776)\}


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 \{(1778)\}**

**ACTS REFERRED TO**


3. [ ]


[Acts of which the Contracting Parties shall take note]


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.


Indent and words “as amended by” added by Decision No 109/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2018.

Indent added by Decision No 109/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2018.

Indent added by Decision No 109/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.6.2018.
The Contracting Parties take note of the content of the following acts:


XXX. MEDICAL DEVICES {[1793]}

**ACTS REFERRED TO**

1. [ ] (1794)


   ([1797]) 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.


   - ([1799])


5. [ ] (1803)


7. [ ] [1856]


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.

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[1854] 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.


[1860] Indent and words “as amended by:” added by Decision No 89/2020 (OJ L [to be published]) and EEA Supplement No [to be published], e.i.f. 18.6.2020.


[1862] Indent and words “as amended by:” added by Decision No 90/2020 (OJ L [to be published]) and EEA Supplement No [to be published], e.i.f. 18.6.2020.

**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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XXXI. RECREATIONAL CRAFT

ACTS REFERRED TO

1. [ ]


XXXII. MARINE EQUIPMENT

ACTS REFERRED TO

1. [ ]


---

1818 Point inserted by Decision No 224/2014 (OJ L 230, 3.9.2015, p. 27 and EEA Supplement No 52, 3.9.2015, p. 27), e.i.f. 1.11.2014. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2015.
1824 Point inserted by Decision No 147/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 7.7.2018
1825 Point inserted by Decision No 244/2018 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.12.2018
APPENDIX I{\textsuperscript{1826}}

ENERGY LABELS

SECTION I{\textsuperscript{1827}} [ ]


\small{\textsuperscript{1827}} Section I (Commission Directive 94/2/EC) and heading added by Decision No 22/98 (OJ L 342, 17.12.1998, p. 32 and EEA Supplement No 52, 17.12.1998, p. 1), e.i.f. 1.4.1998. The following labels correspond to the labels previously contained in former adaptation a) to the first indent of point 4 of Chapter IV, subsequently reinserted as point 4a by Decision No 22/98 subsequently deleted by Decision No 218/2012 (OJ L 81, 21.3.2013, p. 18 and EEA Supplement No 18, 21.3.2013, p. 21), e.i.f. 1.6.2013.
SECTION 2\[^{1828}\]
SECTION 3\(^{(1829)}\) [ ]

SECTION 4\textsuperscript{(1830)}

Commission Directive 96/60/EC

\textit{(household combined washer-driers)}

<table>
<thead>
<tr>
<th>Energi</th>
<th>Kombinert vaske- og tørrmaskin</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Merke</strong></td>
<td><strong>Modell</strong></td>
</tr>
<tr>
<td><strong>Lavt forbruk</strong></td>
<td><strong>LOGO</strong></td>
</tr>
<tr>
<td></td>
<td>A B C 1 2 3</td>
</tr>
<tr>
<td><strong>Høyt forbruk</strong></td>
<td><strong>X.YZ</strong></td>
</tr>
<tr>
<td><strong>Energiforbruk</strong></td>
<td>kWh</td>
</tr>
<tr>
<td>(Ved av C vaskning og tørring med full kapasitet)</td>
<td>X.YZ</td>
</tr>
<tr>
<td>Vask &amp; sentrifugering</td>
<td>kWh</td>
</tr>
<tr>
<td>Den faktiske energibruk avhenger av</td>
<td>X.YZ</td>
</tr>
<tr>
<td>hvordan vaske- og tørrmaskinen brukes</td>
<td></td>
</tr>
<tr>
<td><strong>Vaskeevne</strong></td>
<td>A B C D E F G</td>
</tr>
<tr>
<td>A: høy</td>
<td>G: lav</td>
</tr>
<tr>
<td>sentrifugeringshastighet (omr/min.)</td>
<td>1100</td>
</tr>
<tr>
<td><strong>Kapasitet</strong></td>
<td><strong>Vasking</strong></td>
</tr>
<tr>
<td>(bamull) kg</td>
<td>Vasking</td>
</tr>
<tr>
<td><strong>Vannforbruk</strong> (totalt)</td>
<td><strong>yx</strong></td>
</tr>
<tr>
<td><strong>Lydninå</strong></td>
<td><strong>Vasking</strong></td>
</tr>
</tbody>
</table>

Produktbelysning inneholder ytterligere opplysninger

European standard EN 50287
Unika levert oppen av kompetente vaske og tørrmaskiner
SECTION 5[\textsuperscript{1831}] [ ]

Orka
Framleiðandi
Gerð

Góð nýtni

A
B
C
D
E
F
G

Slæm nýtni

Orkunotkun kWh/lotu
(Lýgur á prófúnnarfræðinum staketafrottlotu þar sem notað er kafl vatn)

Raunnotkun fer etir því hvemig tekið er notað

Pvottahæðni
A: meiri G; minni

Purrkunarhæðni
A: meiri G; minni

Staðalborðbúnaður YZ

Valnsnotkun l/lotu YX

 Hávaði
(dB(A) re 1 pW)

Nánari uppfyllingar er að finna í beæklingum sem tylgja vörnum

Uppþvottavél

Logo
ABC
123

X.YZ

Stofn EN 50042
Tilhefni 97/13/EB um mækinar uppþvottavél
## Energi

<table>
<thead>
<tr>
<th>Merke</th>
<th>Oppvaskmaskin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Modell</td>
<td>Logo ABC 123</td>
</tr>
</tbody>
</table>

### Lavt forbruk

- A
- B
- C
- D
- E
- F
- G

### Høyt forbruk

Energiforbruk kWh/oppvask  
(på grunnlag av testresultater for normalprogram ved kraftvernomsetning)

Den faktiske energibruken avhenger av hvordan maskinen brukes.

<table>
<thead>
<tr>
<th>Rengjøringsvevne</th>
<th>A: høy</th>
<th>G: lav</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tørkeevne</td>
<td>A: høy</td>
<td>G: lav</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Standardkviderer</th>
<th>YZ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vanntilkjøring</td>
<td>YX</td>
</tr>
</tbody>
</table>

Lydtilstand

EB(A), (Støy)

Produktbrosjyrene inneholder ytterligere opplysninger
SECTION 6\(^{1832}\)

Commission Directive 2002/40/EC

*(household electric ovens)*

### Orka

<table>
<thead>
<tr>
<th>Framleiðandi</th>
<th>Gerð</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### Góð nýtti

<table>
<thead>
<tr>
<th>Grade</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Good</td>
</tr>
<tr>
<td>B</td>
<td>Good</td>
</tr>
<tr>
<td>C</td>
<td>Good</td>
</tr>
<tr>
<td>D</td>
<td>Good</td>
</tr>
<tr>
<td>E</td>
<td>Good</td>
</tr>
<tr>
<td>F</td>
<td>Good</td>
</tr>
<tr>
<td>G</td>
<td>Poor</td>
</tr>
</tbody>
</table>

#### Slæm nýtti

<table>
<thead>
<tr>
<th>Orkunotkun (kWh)</th>
<th>Hefðbundinn</th>
<th>Blástursofn</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Notkunarrými (lítar)</th>
<th>X.YZ</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Stærð:</th>
<th>Lítill</th>
<th>Meðal</th>
<th>Stór</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Hávaði (dB(A) re 1 pW)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
</tbody>
</table>

Nánari upplýsingar er að finna í bæklingum sem fylgja vörurnum

Norm EN 50304
Rafmagnsþekkingar
Tilskipun 2002/40/EB um orkumerkingar
<table>
<thead>
<tr>
<th>Energi</th>
<th>Elektrisk stekeovn</th>
</tr>
</thead>
<tbody>
<tr>
<td>Merke</td>
<td>Logo ABC 123</td>
</tr>
<tr>
<td>Modell</td>
<td></td>
</tr>
</tbody>
</table>

### Høyt energiforbruk

<table>
<thead>
<tr>
<th>Grade</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td></td>
</tr>
<tr>
<td>B</td>
<td></td>
</tr>
<tr>
<td>C</td>
<td></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>

### Lavt energiforbruk

- **Energiforbruk** (kWh)
- **Oppvarmingsfunksjon:**
  - Tradisjonell oppvarming
  - Varmluft
- **(basert på standardbelasting)**
- **Nettvolum** (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
## Energieredning

<table>
<thead>
<tr>
<th>Merke</th>
<th>Modell</th>
<th>Høyt energiforbruk</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>A</td>
</tr>
<tr>
<td></td>
<td></td>
<td>B</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
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<td></td>
<td></td>
<td>D</td>
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<tr>
<td></td>
<td></td>
<td>E</td>
</tr>
<tr>
<td></td>
<td></td>
<td>F</td>
</tr>
<tr>
<td></td>
<td></td>
<td>G</td>
</tr>
</tbody>
</table>

## Elektriske stekeovner

### Energiforbruk (kWh)

<table>
<thead>
<tr>
<th>Oppvarmingsfunksjon</th>
<th>X.YZ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tradisjonell oppvarming</td>
<td></td>
</tr>
<tr>
<td>Varmluft</td>
<td></td>
</tr>
</tbody>
</table>

### Nettovolum (liter)

<table>
<thead>
<tr>
<th>Type</th>
<th>X.YZ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Liten</td>
<td></td>
</tr>
<tr>
<td>Middels stor</td>
<td></td>
</tr>
<tr>
<td>Stor</td>
<td></td>
</tr>
</tbody>
</table>

### Lydnivå (støy)

<table>
<thead>
<tr>
<th>dB(A) re 1 pW</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
</tbody>
</table>

Produktbrosjyrene inneholder ytterligere opplysninger.

Standard EN 50304
Elektriske stekeovner
Direktiv 2002/40/EF om energimerking
SECTION 7[1833] [ ]

APPENDIX 2{\textsuperscript{1834}}

ENERGY TABLES

SECTION I{\textsuperscript{1835}} [ ]
SECTION 2[1836] [ ]

SECTION 3 (\{1837\} [ ]
**SECTION 4**

*Commission Directive 96/60/EC*

*(household combined washer-driers)*

<table>
<thead>
<tr>
<th>Note</th>
<th>EN</th>
<th>NO</th>
<th>IS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Label</td>
<td>Fiche</td>
<td>Mail order</td>
</tr>
<tr>
<td></td>
<td>Annex I</td>
<td>Annex II</td>
<td>Annex III</td>
</tr>
<tr>
<td></td>
<td>Energy</td>
<td>Energi</td>
<td>Orka</td>
</tr>
<tr>
<td></td>
<td>Washer-drier</td>
<td>Kombinert vaske- og tørkemaskin</td>
<td>Þvottavél – þurrkari</td>
</tr>
<tr>
<td>I 1</td>
<td>Manufacturer</td>
<td>Merke</td>
<td>Framleiðandi</td>
</tr>
<tr>
<td>II 2</td>
<td>Model</td>
<td>Modell</td>
<td>Gerð</td>
</tr>
<tr>
<td></td>
<td>More efficient</td>
<td>Lavt forbruk</td>
<td>Góð nýtni</td>
</tr>
<tr>
<td></td>
<td>Less efficient</td>
<td>Høyt forbruk</td>
<td>Slæm nýtni</td>
</tr>
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Commission Directive 2002/40/EC

(*household electric ovens*)

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SECTION 7[1841] [ ]

HAZARD AND PRECAUTIONARY STATEMENTS IN ICELANDIC

The following shall be added to Annex III to Regulation (EC) No 1272/2008:

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<tr>
<td>H373</td>
<td>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 för með sér).</td>
</tr>
<tr>
<td>H400</td>
<td>Mjög eitrað lífi í vatni.</td>
</tr>
<tr>
<td>H410</td>
<td>Mjög eitrað lífi í vatni, hefur langvinn áhrif.</td>
</tr>
<tr>
<td>H411</td>
<td>Eitrað lífi í vatni, hefur langvinn áhrif.</td>
</tr>
<tr>
<td>H412</td>
<td>Skaðlegt lífi í vatni, hefur langvinn áhrif.</td>
</tr>
<tr>
<td>H413</td>
<td>Getur valdið langvinnum, skaðlegum áhrifum á líf í vatni.</td>
</tr>
<tr>
<td>H350i</td>
<td>Getur valdið krabbameini við innöndun.</td>
</tr>
<tr>
<td>H360F</td>
<td>Getur haft skaðleg áhrif á frjósemi.</td>
</tr>
<tr>
<td>H360D</td>
<td>Getur haft skaðleg áhrif á börn í móðurkviði.</td>
</tr>
<tr>
<td>H361f</td>
<td>Grunað um að hafa skaðleg áhrif á frjósemi.</td>
</tr>
<tr>
<td>H361d</td>
<td>Grunað um að hafa skaðleg áhrif á börn í móðurkviði.</td>
</tr>
<tr>
<td>H360FD</td>
<td>Getur haft skaðleg áhrif á frjósemi. Getur haft skaðleg áhrif á börn í móðurkviði.</td>
</tr>
<tr>
<td>H361fd</td>
<td>Grunað um að hafa skaðleg áhrif á frjósemi. Grunað um að hafa skaðleg áhrif á börn í móðurkviði.</td>
</tr>
<tr>
<td>H360Fd</td>
<td>Getur haft skaðleg áhrif á frjósemi. Grunað um að hafa skaðleg áhrif á börn í móðurkviði.</td>
</tr>
<tr>
<td>H360Df</td>
<td>Getur haft skaðleg áhrif á börn í móðurkviði. Grunað um að hafa skaðleg áhrif á frjósemi.</td>
</tr>
<tr>
<td>EUH 001</td>
<td>Sprengifímt sem þurrefni.</td>
</tr>
<tr>
<td>EUH 006</td>
<td>Sprengifímt með og án andrúmslofts.</td>
</tr>
<tr>
<td>EUH 014</td>
<td>Hvarfast kröftuglega við vatn</td>
</tr>
<tr>
<td>EUH 018</td>
<td>Getur myndað eldfimar eða sprengifimar blöndur af efnagufu og andrúmslofti við notkun.</td>
</tr>
<tr>
<td>EUH 019</td>
<td>Getur myndað sprengifíms efnasambönd (peroxíð).</td>
</tr>
<tr>
<td>EUH 044</td>
<td>Sprengifímt við hitun í fokuðu rými.</td>
</tr>
<tr>
<td>EUH 029</td>
<td>Myndar eitraða lofttegund í snertingu við vatn.</td>
</tr>
<tr>
<td>EUH 031</td>
<td>Myndar eitraða lofttegund í snertingu við sýru.</td>
</tr>
<tr>
<td>EUH 032</td>
<td>Myndar mjög eitraða lofttegund í snertingu við sýru.</td>
</tr>
<tr>
<td>EUH 066</td>
<td>Endurtekin snerting getur valdið þurri eða sprunginni húð.</td>
</tr>
</tbody>
</table>
The following shall be added to Part 2 of Annex IV of Regulation (EC) No 1272/2008:

<table>
<thead>
<tr>
<th>No.</th>
<th>Icelandic</th>
</tr>
</thead>
<tbody>
<tr>
<td>P101</td>
<td>Ef leita þarf læknis skal hafa flát eða merkimiða tiltæk.</td>
</tr>
<tr>
<td>P102</td>
<td>Gyemist þar sem börn ná ekki til.</td>
</tr>
<tr>
<td>P103</td>
<td>Lesið merkimiðann fyrir notkun.</td>
</tr>
<tr>
<td>P201</td>
<td>Affið sérstakra leiðbeiningar fyrir notkun.</td>
</tr>
<tr>
<td>P202</td>
<td>Nauðsýnlegt er að lesa og skilja allar viðvaranir áður en efnið er notað.</td>
</tr>
<tr>
<td>P210</td>
<td>Haldið frá hitagjöfum, neistagjöfum, opnum eldi og heitum flötum. — Reykningar bannaðar.</td>
</tr>
<tr>
<td>P211</td>
<td>Má ekki úða á opinn eld eða annan íkveikjuvald.</td>
</tr>
<tr>
<td>P220</td>
<td>Má ekki nota eða geyma í námunda við fatnað/.../brennanleg efni.</td>
</tr>
<tr>
<td>P221</td>
<td>Gætið þess að blanda efninu ekki saman við brennanleg efni/...</td>
</tr>
<tr>
<td>P222</td>
<td>Má ekki komast í snertingu við andrúmsloft.</td>
</tr>
<tr>
<td>P223</td>
<td>Má alls ekki komast í snertingu við vatn vegna hættu á kröftugu hvarfi og leiftureldi.</td>
</tr>
<tr>
<td>P230</td>
<td>Haldið röku með....</td>
</tr>
<tr>
<td>P231</td>
<td>Meðhöndlið undir Óhvarfgjarni lofttegund.</td>
</tr>
</tbody>
</table>
Icelandic

<table>
<thead>
<tr>
<th>No.</th>
<th>Icelandic</th>
</tr>
</thead>
<tbody>
<tr>
<td>P232</td>
<td>Verjið gegn raka.</td>
</tr>
<tr>
<td>P233</td>
<td>Ílát skal vera vel lukt.</td>
</tr>
<tr>
<td>P234</td>
<td>Má aðeins geyma í upprunalegu ílátí.</td>
</tr>
<tr>
<td>P235</td>
<td>Geymist á köldum stað.</td>
</tr>
<tr>
<td>P240</td>
<td>Jarðtengið/spennujafnið ílát og viðökubúnað.</td>
</tr>
<tr>
<td>P241</td>
<td>Notið sprengiheld rafföng/lofræstibúnað/lysinu/...</td>
</tr>
<tr>
<td>P242</td>
<td>Notið ekki verkfæri sem mynda neista.</td>
</tr>
<tr>
<td>P243</td>
<td>Gerið varúðarráðstafanir gegn stöðfurafmagni</td>
</tr>
<tr>
<td>P244</td>
<td>Gætið þess að ekki sé feiti og olía á þrystingslokum.</td>
</tr>
<tr>
<td>P250</td>
<td>Má ekki verða fyrir hnjski/höggi/.../núningi</td>
</tr>
<tr>
<td>P251</td>
<td>Þrystihylki: Ekki mæ gata eða brenda hylki jafnvel þott þau sér tóm.</td>
</tr>
<tr>
<td>P260</td>
<td>Andið ekki að ykkur rykri/reykk/lofttegund/úða/gufu/ýringi.</td>
</tr>
<tr>
<td>P261</td>
<td>Gætið þess að anda ekki inn rykri/reykk/lofttegund/úða/gufu/ýringi.</td>
</tr>
<tr>
<td>P262</td>
<td>Má ekki koma í augu eða á húð eða fót.</td>
</tr>
<tr>
<td>P263</td>
<td>Forðist alla snertingu við efninð meðan á meðgöngu og brjóstagjöf stendur.</td>
</tr>
<tr>
<td>P264</td>
<td>Þvoið...vandlega eftir meðhöndlun.</td>
</tr>
<tr>
<td>P270</td>
<td>Neytið ekki matar, drykkjar eða tóbaks við notkun þessarar vörur.</td>
</tr>
<tr>
<td>P271</td>
<td>Notið eingöngu utandyra eða í vel loftfræstu rými.</td>
</tr>
<tr>
<td>P272</td>
<td>Ekki skal farið með vinnufót af vinnustað hafi þau óheinkast af efninu.</td>
</tr>
<tr>
<td>P273</td>
<td>Forðist losun út í umhverfið.</td>
</tr>
<tr>
<td>P280</td>
<td>Notið hliðarfahska/hlíðarfatnað/augnhlífar/andlitshlífar.</td>
</tr>
<tr>
<td>P281</td>
<td>Notið tilskildar persónuhlífar.</td>
</tr>
<tr>
<td>P282</td>
<td>Klæðist kuldaeinangrandi hönskum/andlitshlífar/augnhlífum.</td>
</tr>
<tr>
<td>P283</td>
<td>Klæðist brunarþolnum/eldþolnum/eldtefjandi fatnæði.</td>
</tr>
<tr>
<td>P284</td>
<td>Notið öndunarhlífar.</td>
</tr>
<tr>
<td>P285</td>
<td>Notið öndunarhlífar ef loftfræsting er öftunnægjandi.</td>
</tr>
<tr>
<td>P301</td>
<td>EFTIR INNTÖKU:</td>
</tr>
<tr>
<td>P302</td>
<td>BERIST EFNÍÐ Á HÚÐ:</td>
</tr>
<tr>
<td>P303</td>
<td>BERIST EFNÍÐ Á HÚÐ (eða í hár):</td>
</tr>
<tr>
<td>P304</td>
<td>EFTIR INNÖNDUN:</td>
</tr>
<tr>
<td>No.</td>
<td>Icelandic</td>
</tr>
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<td>-----</td>
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</tr>
<tr>
<td>P305</td>
<td>BERIST EFNÍD Í AUGU:</td>
</tr>
<tr>
<td>P306</td>
<td>EF EFNÍD FER Á FÓT:</td>
</tr>
<tr>
<td>P307</td>
<td>EF um váhrif er að ræða:</td>
</tr>
<tr>
<td>P308</td>
<td>EF um váhrif eða huganleg váhrif er að ræða:</td>
</tr>
<tr>
<td>P309</td>
<td>EF um váhrif er að ræða eða ef lasleika verður vart:</td>
</tr>
<tr>
<td>P310</td>
<td>Hringið umsvifalaust í EITRUNARMÍ DSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P311</td>
<td>Hringið í EITRUNARMÍ DSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P312</td>
<td>Hringið í EITRUNARMÍ DSTÖÐ eða lækni ef lasleika verður vart.</td>
</tr>
<tr>
<td>P313</td>
<td>Leitið læknis.</td>
</tr>
<tr>
<td>P314</td>
<td>Leitið læknis ef lasleika verður vart.</td>
</tr>
<tr>
<td>P315</td>
<td>Leitið umsvifalaust læknis.</td>
</tr>
<tr>
<td>P320</td>
<td>Brynt er að fá sérstaka meðferð (sjá .... á þessum merkimiða).</td>
</tr>
<tr>
<td>P321</td>
<td>Sérstök meðferð (sjá .... á þessum merkimiða).</td>
</tr>
<tr>
<td>P322</td>
<td>Sérstakar råðstafanir (sjá .... á þessum merkimiða).</td>
</tr>
<tr>
<td>P330</td>
<td>Skolið munninn.</td>
</tr>
<tr>
<td>P331</td>
<td>EKKI framkalla uppköst.</td>
</tr>
<tr>
<td>P332</td>
<td>Ef efnið ertir húð.</td>
</tr>
<tr>
<td>P333</td>
<td>Ef efnið ertir húð eða útbrot koma fram:</td>
</tr>
<tr>
<td>P334</td>
<td>Sökkvið í kalt vatn/vefið með blautu sárabindi.</td>
</tr>
<tr>
<td>P335</td>
<td>Dustið lausar agnir af húðinni.</td>
</tr>
<tr>
<td>P337</td>
<td>Ef augnerting er viðvarandi:</td>
</tr>
<tr>
<td>P338</td>
<td>Fjarlægði snertilinsur ef það er auðvelt. Skolið áfram.</td>
</tr>
<tr>
<td>P340</td>
<td>Flytjið viðkomandi í ferskt loft og látið hann hvilast í stellingu sem léttir öndun.</td>
</tr>
<tr>
<td>P341</td>
<td>Ef viðkomandi á erfitt með öndun skal flytja hann í ferskt loft og látu hann hvilast í stellingu sem léttir öndun.</td>
</tr>
<tr>
<td>P342</td>
<td>Ef vart verður einkenna frá öndunarvegi:</td>
</tr>
<tr>
<td>P350</td>
<td>Þvoið varlega með mikilli sápu og vatni.</td>
</tr>
<tr>
<td>P351</td>
<td>Þvoið varlega með mikilli sápu og vatni.</td>
</tr>
<tr>
<td>P352</td>
<td>Þvoið með mikilli sápu og vatni.</td>
</tr>
<tr>
<td>P353</td>
<td>Skolið húðina með vatni/Farið í sturtu.</td>
</tr>
<tr>
<td>P360</td>
<td>Föt og húð, sem öhrinkast af efninu, skal skola strax með miklu vatni abur en farið er úr féttunum.</td>
</tr>
<tr>
<td>P361</td>
<td>Farið strax úr féttum sem öhrinkast af efninu.</td>
</tr>
<tr>
<td>No.</td>
<td>Icelandic</td>
</tr>
<tr>
<td>-----</td>
<td>-----------</td>
</tr>
<tr>
<td>P362</td>
<td>Farið úr fötum, sem öhrinkast af efninu, og þvoið fyrir næstu notkun.</td>
</tr>
<tr>
<td>P363</td>
<td>Þvoið fót, sem öhrinkast af efninu, fyrir næstu notkun.</td>
</tr>
<tr>
<td>P370</td>
<td>Ef eldur kemur upp:</td>
</tr>
<tr>
<td>P371</td>
<td>Þegar um mikinn eld og mikilð efnismagn er að ræda:</td>
</tr>
<tr>
<td>P372</td>
<td>Sprengihætta ef eldur kemur upp.</td>
</tr>
<tr>
<td>P373</td>
<td>EKKI reyna að slökkva eld ef hann kemst að sprengifimum efnun.</td>
</tr>
<tr>
<td>P374</td>
<td>Beitið eðlilegu varðbæðistöfunum við slökkkvistörf og verið í hæfilegt fjrarlagð frá eldnum.</td>
</tr>
<tr>
<td>P375</td>
<td>Verið í fjarlægð frá eldnum við slökkkvistörf vegna sprengihættu.</td>
</tr>
<tr>
<td>P376</td>
<td>Stöðvið leka ef það er öhætt.</td>
</tr>
<tr>
<td>P377</td>
<td>Eldur í lekandi gæsi: Reynið ekk að slökkva eldinn nema hægt sé að stöðva lekann á öruggan máta.</td>
</tr>
<tr>
<td>P378</td>
<td>Notið ... til að slökkva eldinn.</td>
</tr>
<tr>
<td>P380</td>
<td>Rýmið svæðið.</td>
</tr>
<tr>
<td>P381</td>
<td>Fjarlagið alla Íkveikjuvalda ef það er öhætt.</td>
</tr>
<tr>
<td>P390</td>
<td>Sogið upp allt sem hellist niður til að afstýra eignatjóni.</td>
</tr>
<tr>
<td>P391</td>
<td>Safnið upp því sem hellist niður.</td>
</tr>
<tr>
<td>P301 + P310</td>
<td>EFTIR INNTÖKU: Hringið umsvifalaust í EITRUNARMÍDSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P301 + P312</td>
<td>EFTIR INNTÖKU: Hringið í EITRUNARMÍDSTÖÐ eða lækni ef lasleika verður vart.</td>
</tr>
<tr>
<td>P301 + P330 + P331</td>
<td>EFTIR INNTÖKU: Skolið munninn. EKKI framkalla uppköst.</td>
</tr>
<tr>
<td>P302 + P334</td>
<td>BERIST EFNIÐ Á HÚÐ: Sökkvið í kalt vatn/verfjöð með blautu sárabindi.</td>
</tr>
<tr>
<td>P302 + P350</td>
<td>BERIST EFNIÐ Á HÚÐ: Þvoið varlega með mikili sápu og vatni.</td>
</tr>
<tr>
<td>P302 + P352</td>
<td>BERIST EFNIÐ Á HÚÐ: Þvoið með mikili sápu og vatni.</td>
</tr>
<tr>
<td>P304 + P340</td>
<td>EFTIR INNÖNDUND: Flytjöð viðkomandi í ferskt loft og látið hann hvílast í stellingu sem létir öndun.</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>P306 + P360</td>
<td>EF EFINÍD FER Á FÓT: Föt og húð, 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 váhrif er að ræða: Hringið í EITRUNARMÍÐSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P308 + P313</td>
<td>EF um váhrif eða hugsanleg váhrif er að ræða: Leitið læknis.</td>
</tr>
<tr>
<td>P309 + P311</td>
<td>EF um váhrif er að ræða eða ef lasleika verður vart: Hringið í EITRUNARMÍÐSTÖÐ eða lækni.</td>
</tr>
<tr>
<td>P332 + P313</td>
<td>Ef efníð ertir húð: Leitið læknis.</td>
</tr>
<tr>
<td>P333 + P313</td>
<td>Ef efníð ertir húð eða útbrot koma fram: Leitið læknis.</td>
</tr>
<tr>
<td>P335 + P334</td>
<td>Dustið lausar agnir af húðinni. Sökkvið í kalt vatni/vefjið með blautu sárabindi.</td>
</tr>
<tr>
<td>P337 + P313</td>
<td>Ef augnering 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 efnismagn 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 þurrun stað.</td>
</tr>
<tr>
<td>P403</td>
<td>Geymist á vel loftfræstum stað.</td>
</tr>
<tr>
<td>P404</td>
<td>Geymist í lokuðu ílái.</td>
</tr>
</tbody>
</table>
APPENDIX 6\(^{(1845)}\)

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>

\(^{(1845)}\) Appendix and heading added 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.
<table>
<thead>
<tr>
<th>No.</th>
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<tbody>
<tr>
<td>H204</td>
<td>Fare for brann eller utkast av fragmenter.</td>
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<tr>
<td>H205</td>
<td>Fare for masseeksplosjon ved brann.</td>
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<tr>
<td>H220</td>
<td>Ekstremt brannfarlig gass.</td>
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<tr>
<td>H221</td>
<td>Brannfarlig gass.</td>
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<tr>
<td>H222</td>
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<td>H223</td>
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<td>H251</td>
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<tr>
<td>H252</td>
<td>Selvopphetende i store mengder; kan selvantenne.</td>
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<tr>
<td>H260</td>
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</tr>
<tr>
<td>H261</td>
<td>Ved kontakt med vann utvikles brannfarlige gasser.</td>
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<tr>
<td>H270</td>
<td>Kan forårsake eller forsterke brann; oksiderende.</td>
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<tr>
<td>H271</td>
<td>Kan forårsake brann eller eksplosjon; sterkt oksiderende.</td>
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<tr>
<td>H272</td>
<td>Kan forsterke brann; oksiderende.</td>
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<td>H280</td>
<td>Inneholder gass under trykk; kan eksplodere ved oppvarming.</td>
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<td>H281</td>
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<tr>
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<td>Kan være etsende for metaller.</td>
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<td>H300</td>
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<tr>
<td>H302</td>
<td>Farlig ved svelging.</td>
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<tr>
<td>H304</td>
<td>Kan være dødelig ved svelging om det kommer ned i luftveiene.</td>
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<td>Dødelig ved hudkontakt.</td>
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<td>Irriterer huden.</td>
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<td>H317</td>
<td>Kan utløse en allergisk hudreaksjon.</td>
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<tr>
<td>H318</td>
<td>Gir alvorlig øyeskade.</td>
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<td>H319</td>
<td>Gir alvorlig øyeirritasjon.</td>
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<tr>
<td>H330</td>
<td>Dødelig ved innånding.</td>
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<td>H331</td>
<td>Giftig ved innånding.</td>
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<tr>
<td>H332</td>
<td>Farlig ved innånding.</td>
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<tr>
<td>H334</td>
<td>Kan gi allergi eller astmasymptomer eller pustevansker ved innånding.</td>
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<tr>
<td>H335</td>
<td>Kan forårsake irritasjon av luftveiene.</td>
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<td>H336</td>
<td>Kan forårsake døsighet eller svimmelhet.</td>
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<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>
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<tr>
<td>H410</td>
<td>Meget giftig, med langtidsvirkning, for liv i vann.</td>
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<tr>
<td>H411</td>
<td>Giftig, med langtidsvirkning, for liv i vann.</td>
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<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>
</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>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>
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<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 eksponering kan gi øttr 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>
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<tr>
<td>EUH 059</td>
<td>Farlig for ozonlaget.</td>
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<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>

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<tr>
<th>No.</th>
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<tbody>
<tr>
<td>P101</td>
<td>Dersom det er nødvendig med legehjelp, ha produktets beholder eller etikett for hånden.</td>
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<tr>
<td>P102</td>
<td>Oppbevares utilgjengelig for barn.</td>
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<td>P103</td>
<td>Les etiketten før bruk.</td>
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<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>
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<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>
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<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>
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<td>P231</td>
<td>Håndteres under inertgass.</td>
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<td>P234</td>
<td>Oppbevares bare i originalbeholder.</td>
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<tr>
<td>P235</td>
<td>Oppbevares kjølig.</td>
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<td>P240</td>
<td>Beholder og mottaksutstyr jordes/potensialutlignes.</td>
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<td>P241</td>
<td>Bruk elektrisk materiell /ventilasjonsmateriell/belysningsmateriell som er eksplosjonssikkert.</td>
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<tr>
<td>P242</td>
<td>Bruk bare verktøy som ikke avger gnister.</td>
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<tr>
<td>P243</td>
<td>Treff tiltak mot statisk elektrisitet.</td>
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<tr>
<td>P244</td>
<td>Reduksjonsventiler skal holdes fri for fett og olje.</td>
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<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>
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<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>
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<tr>
<td>P263</td>
<td>Unngå kontakt under graviditet/amming.</td>
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<tr>
<td>P264</td>
<td>Vask … grundig etter bruk.</td>
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<tr>
<td>P270</td>
<td>Ikke spis, drikk eller røyk ved bruk av produktet.</td>
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<tr>
<td>P271</td>
<td>Brukes bare utendørs eller i et godt ventilert område.</td>
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<tr>
<td>P272</td>
<td>Tilsølte arbeidshjelm må ikke fjernes fra arbeidsplassen.</td>
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<td>P273</td>
<td>Unngå utslipp til miljøet.</td>
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<tr>
<td>P280</td>
<td>Benytt vernehansker /verneklær/vernebriller/ansiktsskjerm.</td>
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<td>P281</td>
<td>Bruk påkrevd personlig verneutstyr.</td>
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<td>P282</td>
<td>Bruk kuldeisolerende hansker /visir/øyevern.</td>
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<tr>
<td>P283</td>
<td>Benytt brannbestandige/flammehemmende klær.</td>
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<tr>
<td>P284</td>
<td>Bruk åndedrettsvern.</td>
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<tr>
<td>P285</td>
<td>Ved utilstrekkelig ventilasjon skal åndedrettsvern benyttes.</td>
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<td>P301</td>
<td>VED SVELGING:</td>
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<tr>
<td>P302</td>
<td>VED HUDKONTAKT:</td>
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<tr>
<td>P303</td>
<td>VED HUDKONTAKT (eller håret):</td>
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<td>P304</td>
<td>VED INNÅNDING:</td>
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<tr>
<td>P305</td>
<td>VED KONTAKT MED ØYNENE</td>
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<tr>
<td>P306</td>
<td>VED KONTAKT MED KLÆR:</td>
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<td>P307</td>
<td>Ved eksponering:</td>
</tr>
<tr>
<td>P308</td>
<td>Ved eksponering eller misanke om eksponering:</td>
</tr>
<tr>
<td>P309</td>
<td>Ved eksponering eller ubehag:</td>
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<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>
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<td>P313</td>
<td>Søk legehjelp.</td>
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<td>P314</td>
<td>Søk legehjelp ved ubehag.</td>
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<td>P315</td>
<td>Søk legehjelp umiddelbart.</td>
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<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>
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<tr>
<td>P330</td>
<td>Skyll munnen.</td>
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<tr>
<td>P331</td>
<td>IKKE framkall brekning.</td>
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<tr>
<td>P332</td>
<td>Ved hudirritasjon:</td>
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<tr>
<td>P333</td>
<td>Ved hudirritasjon eller utsett:</td>
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<tr>
<td>P334</td>
<td>Skyll i kaldt vann / anvend våt kompress.</td>
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<tr>
<td>P335</td>
<td>Børst bort løse partikler fra huden.</td>
</tr>
<tr>
<td>P336</td>
<td>Varm opp frostskadede legemsdeler med lunkent vann. Ikke gni på det skadede området.</td>
</tr>
<tr>
<td>P337</td>
<td>Ved vedvarende øyeirritasjon:</td>
</tr>
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<td>Norwegian</td>
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<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>
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<td>P342</td>
<td>Ved symptomer i luftveiene:</td>
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<td>P350</td>
<td>Vask forsiktig med mye såpe og vann.</td>
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<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 klærne 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>
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<tr>
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