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

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

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

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

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

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

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

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

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


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

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

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

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

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

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

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

The EFTA States shall participate fully in the Telematic Exchange of Information on Medicinal Products (IMP) programme.
The EFTA States shall be fully associated with the work of the Executive Steering Group on Shortages and Safety of Medicinal Products as set up by Article 3 of Regulation (EU) 2022/123 of the European Parliament and of the Council and shall have the same rights and obligations within it as the EU Member States, except for the right to vote.

The EFTA States shall be fully associated with the work of the Emergency Task Force as set up by Article 15 of Regulation (EU) 2022/123 of the European Parliament and of the Council and shall have the same rights and obligations within it as the EU Member States, except for the right to vote.

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

A centralised marketing authorisation granted for a medicinal product shall not be subject to any fees other than those referred to in Article 67(3) of Regulation (EC) No 726/2004 of the European Parliament and of the Council.

The Agency having legal personality shall enjoy in all the States of the Contracting Parties the most extensive legal capacity accorded to legal persons under their laws.

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


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

ACTS REFERRED TO


10.\(^{14}\) [ ]

11.\(^{14}\) [ ]


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.


12e. {19} **32019 R 1871:** Commission Regulation (EU) 2019/1871 of 7 November 2019 on reference points for action for non-allowed pharmacologically active substances present in food of animal origin and repealing Decision 2005/34/EC (OJ L 289, 8.11.2019, p. 41), as amended by:


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

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


-{28} **32011 R 0363:** Commission Regulation (EU) No 363/2011 of 13 April 2011 (OJ L 100, 14.4.2011, p. 28),

-{29} **32012 R 0084:** Commission Implementing Regulation (EU) No 84/2012 of 1 February 2012 (OJ L 30, 2.2.2012, p. 1),

-{30} **32012 R 0085:** Commission Implementing Regulation (EU) No 85/2012 of 1 February 2012 (OJ L 30, 2.2.2012, p. 4),

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\(\{46\}\) 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.

\(\{47\}\) Indent added by Decision No 190/2013 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.11.2013.


\(\{51\}\) Indent added by Decision No 81/2014 (OJ L 310, 30.10.2014, p. 34 and EEA Supplement No 63, 30.10.2014, p. 25), e.i.f. 17.5.2014.

\(\{52\}\) Indent added by Decision No 177/2014 (OJ L 202, 30.7.2015, p. 27 and EEA Supplement No 43, 30.7.2015, p. 27), e.i.f. 26.9.2014.

\(\{53\}\) Indent added by Decision No 177/2014 (OJ L 202, 30.7.2015, p. 27 and EEA Supplement No 43, 30.7.2015, p. 27), e.i.f. 26.9.2014.


(63) 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.


(68) 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.


- (76) 32015 R 1308: Commission Implementing Regulation (EU) 2015/1308 of 29 July 2015 (OJ L 200, 30.7.2015, p. 11),


- (81) 32016 R 0129: Commission Implementing Regulation (EU) 2016/129 of 1 February 2016 (OJ L 25, 2.2.2016, p. 44),

- (82) 32016 R 0305: Commission Implementing Regulation (EU) 2016/305 of 3 March 2016 (OJ L 58, 4.3.2016, p. 35),


- (85) 32016 R 0710: Commission Implementing Regulation (EU) 2016/710 of 12 May 2016 (OJ L 125, 13.5.2016, p. 6),


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(85) 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.


(96) Indent added by Decision No 262/2019 (OJ L 210, 27.7.2020, p. 29 and EEA Supplement No 44, 27.7.2020, p. 31), e.i.f. 30.3.2019.


(100) Indent added by Decision No 224/2019 (OJ L 4, 5.1.2023, p. 39 and EEA Supplement No 3, 5.1.2023, p. 39), e.i.f. 28.9.2019. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 10.12.2021


14. [ ]

15. [ ]


15b. [ ]

15c. [ ]

15d. [ ]

15e. [ ]

15f. [ ]
15h. [121]


[137] 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.


15i.\textsuperscript{138} \textbf{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. \textsuperscript{139} [ ]

15k. \textsuperscript{140} [ ]


\textsuperscript{143}The provisions of the Regulation shall, for the purposes of the present Agreement, be read with the following adaptation:

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.\textsuperscript{144} \textbf{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. \textsuperscript{146} [ ]


\[^{143}\] 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.

\[^{145}\] The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation:

Liechtenstein shall not be obliged to participate in the 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.


\[^{150}\] 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), c.i.f. 2.7.2011.


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.

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).

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).

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.

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).

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.


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.


Indent added by Decision No 371/2021 (OJ L, 2024/711, 14.3.2024 and EEA Supplement No 23, 14.3.2024, p. 76), e.i.f. pending.


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 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-ports 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.(172)  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:

-(173)  32013 D 0262: Commission Implementing Decision 2013/262/EU of 4 June 2013 (OJ L 152, 5.6.2013, p. 52),


-(175)  32013 D 0196: Commission Implementing Decision 2013/96/EU of 24 April 2013 (OJ L 113, 25.4.2013, p. 22),


-(177)  32019 D 0769: Commission Implementing Decision (EU) 2019/769 of 14 May 2019 (OJ L 126, 15.5.2019, p. 70),


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

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(173) 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. 9.10.2013.


(177) Indent added by Decision No 102/2020 (OJ L 172, 6.7.2023, p. 17 and EEA Supplement No 51, 6.7.2023, p. 17), e.i.f. 15.7.2020.


(179) Indent added by Decision No 59/2024 (OJ L, 2024/1598, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 30), e.i.f. 16.3.2024.


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 Inspektionen).


15r. [ ] \{187\}

15s. [ ] \{188\}

15t. [ ] \{189\}


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 authorizations within the decentralized procedure (DCP) and the mutual recognition procedure (MRP).


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 authorizations within the decentralized procedure (DCP) and the mutual recognition procedure (MRP).


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.


[201] Indent added by Decision No 151/2019 (OJ L 291, 10.11.2022, p. 28 and EEA Supplement No 74, 10.11.2022, p. 29), e.i.f. 15.6.2019.


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.


\textsuperscript{208} Point and adaptation text inserted by Decision No 49/2018 (OJ L 26, 30.1.2020, p. 28 and EEA Supplement No 6, 30.1.2020, p. 21), e.i.f. 24.3.2018.

\textsuperscript{209} Point and adaptation text inserted by Decision No 49/2018 (OJ L 26, 30.1.2020, p. 28 and EEA Supplement No 6, 30.1.2020, p. 21), e.i.f. 24.3.2018.


\textsuperscript{209} Indent and words “as amended by:” added by Decision No 212/2015 (OJ L 85, 30.3.2017, p. 29 and EEA Supplement No 19, 30.3.2017, p. 29), e.i.f. 26.9.2015.

\textsuperscript{211} Point inserted by Decision No 140/2007 (OJ L 100, 10.4.2008, p. 66 and EEA Supplement No 19, 10.4.2008, p. 68), e.i.f. 27.10.2007.

\textsuperscript{211} 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.

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


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\[^{21}\] Indent and words “; as amended by:” added by Decision No 126/2009 (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 126/2009 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).


\[^{23}\] 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.

\[^{24}\] 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.


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\[^{22}\] 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).
The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:

In Article 34(2), the words "or Article 53 of the EEA Agreement" shall be inserted after the words "Article 101 TFEU".

15zf. [ ] [229]


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


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\(\text{[225]}\) Three points and text inserted by Decision No 128/2009 (OJ L 62, 11.3.2010, p. 16 and EEA Supplement No 12, 11.3.2010, p. 15), e.i.f. 23.12.2009. For Liechtenstein, Decision 128/2009 entered into force 1.12.2010 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).

\(\text{[226]}\) Indent and words "", as amended by: added by Decision No 158/2013 (OJ L 58, 27.2.2014, p. 10 and EEA Supplement No 13, 27.2.2014, p. 11), e.i.f. 28.2.2014.


\(\text{[228]}\) Indent added by Decision No 335/2021 (OJ L 2024/641, 14.3.2024 and EEA Supplement No 23, 14.3.2024, p. 29), e.i.f. 11.12.2021.

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

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

15zk. [ ] [231]


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


{232} Indent added by Decision No 146/2012 (OJ L 309, 8.11.2012, p. 31 and EEA Supplement No 63, 8.11.2012, p. 36), e.i.f. 14.7.2012.

{233} Indent added by Decision No 146/2012 (OJ L 309, 8.11.2012, p. 31 and EEA Supplement No 63, 8.11.2012, p. 36), e.i.f. 14.7.2012.


{238} Indent added by Decision No 107/2018 (OJ L 368, 5.11.2020, p. 7 and EEA Supplement No 71, 5.11.2020, p. 8), e.i.f. 1.6.2018.


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)(h) and 23 shall only apply to Liechtenstein as far as necessary for transposing Articles 15 and 16 of the Directive.


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

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

15zo.\footnote{244} \textbf{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).


\footnote{241} Point inserted by Decision No 47/2012 (OJ L 207, 2.8.2012, p. 27 and EEA Supplement No 43, 2.8.2012, p. 31), e.i.f. 31.3.2012.

\footnote{242} Point inserted by Decision No 164/2013 (OJ L 58, 27.2.2014, p. 17 and EEA Supplement No 13, 27.2.2014, p. 19), e.i.f. 1.7.2014.

\footnote{243} Point inserted by Decision No 179/2014 (OJ L 202, 30.7.2015, p. 29 and EEA Supplement No 43, 30.7.2015, p. 29), e.i.f. 1.6.2015.


\footnote{246} Point inserted by Decision No 15/2015 (OJ L 93, 7.4.2016, p. 26 and EEA Supplement No 21, 7.4.2016, p. 23), e.i.f. 1.1.2016.


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.


\[249\] Indent added by Decision No 371/2021 (OJ L, 2024/711, 14.3.2024, p. 76), e.i.f. pending.


\[251\] Indent and words “, as amended by:” added by Decision No 87/2018 (OJ L 340, 15.10.2020, p. 18 and EEA Supplement No 66, 15.10.2020, p. 22), e.i.f. 28.4.2018.


\[253\] Indent added by Decision No 107/2021 (OJ L 249/1, 18.1.2024 and EEA Supplement No 5, 18.1.2024, p. 36), e.i.f. 20.3.2021.


\[257\] Point inserted by Decision No 213/2015 (OJ L 85, 30.3.2017, p. 31 and EEA Supplement No 19, 30.3.2017, p. 31), e.i.f. 1.8.2016. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 3.2.2017.


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

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


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

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

[289b] Point inserted by Decision No 28/2022 (OJ L 175, 30.6.2022, p. 45 and EEA Supplement No 42, 30.6.2022, p. 42), e.i.f. 5.2.2022.
[289c] Point inserted by Decision No 63/2022 (OJ L 182, 7.7.2022, p. 41 and EEA Supplement No 45, 7.7.2022, p. 27), e.i.f. 19.3.2022.
[289g] Point and adaptations text inserted by Decision No 371/2021 (OJ L 2024/711, 14.3.2024 and EEA Supplement No 23, 14.3.2024, p. 76), e.i.f. pending.
[289h] Indent and words “, as amended by:” added by Decision No 5/2022 (OJ L 175, 30.6.2022, p. 9 and EEA Supplement No 42, 30.6.2022, p. 8), e.i.f. pending.
[289i] Indent added by Decision No 179/2023 (OJ L 2024/800, 21.3.2024 and EEA Supplement No 26, 21.3.2024, p. 4), e.i.f. pending.
- Article 57
- Articles 103 to 115, included.

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

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


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

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


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

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

({271}) Indent and words “, as amended by:” added by Decision No 67/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

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

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

({274}) Point inserted by Decision No 119/2022 (OJ L 246, 22.9.2022, p. 60 and EEA Supplement No 61, 22.9.2022, p. 59), e.i.f. pending.

({275}) Point inserted by Decision No 119/2022 (OJ L 246, 22.9.2022, p. 60 and EEA Supplement No 61, 22.9.2022, p. 59), e.i.f. pending.

22h. {277} 32022 R 0209: Commission Implementing Regulation (EU) 2022/209 of 16 February 2022 establishing the format of the data to be collected and reported in order to determine the volume of sales and the use of antimicrobial medicinal products in animals in accordance with Regulation (EU) 2019/6 of the European Parliament and of the Council (OJ L 35, 17.2.2022, p. 7).


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

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


2.{282} 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**{284}

**ACTS REFERRED TO**


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


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

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

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

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

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

(c) In Article 52, as regards the EFTA States, the words “or the date of entry into force of Decision of the EEA Joint Committee No 77/2023 of 28 April 2023, whichever is the later” are inserted after the words “16 July 2022”.


[293] Point inserted by Decision No 60/2024 (OJ L, 2024/1612, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 31), c.i.f. 16.3.2024.


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

The Contracting Parties shall take note of the following acts:


**XV. DANGEROUS SUBSTANCES**

**ACTS REFERRED TO**

1. [ ] \{302\}

2. [ ] \{303\}
3. [ ] (385)

4. [ ] (386)

5. [ ] (387)

6. [ ] (388)

7. [ ] (389)


10. [ ] (392)

11. [ ] (393)

12. [ ] (394)

12a. [ ] (395)

12b. [ ] (396)

12c. [ ] (397)


12d. \[ \] \[137\]

12e. \[ \] \[138\]

12f. \[ \] \[139\]


12j. \[ \] \[143\]


[^228] 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.


[^315] 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.


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


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

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

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

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

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

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

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

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

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

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

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

“4. If the Commission amends a Union authorisation, the EFTA States shall amend the corresponding decision.”
(i) The following paragraph shall be added in Article 75:

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

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

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

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


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12np.{[32015 D 1751]} 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 to 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 256, 1.10.2015, p. 15).


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- [ ] [405]

The provisions of the Regulation shall, for the purposes of the present Agreement, be read with the following adaptations, in addition to the adaptations to the Directive 98/8/EC:

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

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

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


\[\begin{align*}
\end{align*}\]


\[446\] Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), e.i.f. 27.10.2018.


\[448\] Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), e.i.f. 27.10.2018.


\[450\] Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), e.i.f. 27.10.2018.


\[452\] Indent added by Decision No 210/2018 (OJ L 105, 25.3.2021, p. 4 and EEA Supplement No 21, 25.3.2021, p. 4), e.i.f. 27.10.2018.


Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), e.i.f. 15.6.2019.

Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), e.i.f. 15.6.2019.

Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), e.i.f. 15.6.2019.

Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), e.i.f. 15.6.2019.

Indent added by Decision No 153/2019 (OJ L 291, 10.11.2022, p. 30 and EEA Supplement No 74, 10.11.2022, p. 31), e.i.f. 15.6.2019.


Indent added by Decision No 205/2021 (OJ L 2024/311, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 35), e.i.f. 10.7.2021.


Indent added by Decision No 121/2022 (OJ L 246, 22.9.2022, p. 63 and EEA Supplement No 61, 22.9.2022, p. 62), e.i.f. 30.4.2022.

Indent added by Decision No 122/2022 (OJ L 246, 22.9.2022, p. 64 and EEA Supplement No 61, 22.9.2022, p. 63), e.i.f. 30.4.2022.


\(^{473}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{474}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{475}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{476}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{477}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{478}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{479}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{480}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{481}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{482}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{483}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{484}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{485}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{486}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{487}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{488}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{489}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{490}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{491}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{492}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{493}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{494}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{495}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{496}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{497}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{498}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{499}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.

\(^{500}\) Indent added by Decision No 169/2022 (OJ L 267, 13.10.2022, p. 8), e.i.f. 11.6.2022.


\textsuperscript{[487]} Indent added by Decision No 263/2023 (OJ L, 2024/1153, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 31), e.i.f. 28.10.2023.

\textsuperscript{[488]} Indent added by Decision No 265/2023 (OJ L, 2024/1149, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 33), e.i.f. 28.10.2023.

\textsuperscript{[489]} Indent added by Decision No 61/2024 (OJ L, 2024/1599, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 33), e.i.f. 16.3.2024.


\textsuperscript{[496]} Indent 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.

\textsuperscript{[497]} 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.


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

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

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

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

(ii) Article 12; and,

(iii) Article 13(3).

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


{505} Indent added by Decision No 264/2023 (OJ L., 2024/1151, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 32), e.i.f. 28.10.2023.

{506} Indent added by Decision No 267/2023 (OJ L., 2024/1136, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 38), e.i.f. 28.10.2023.


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.


\(^{[519]}\) 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.

\(^{[520]}\) 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.


\(^{[526]}\) 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.


[^{43}] 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.

[^{44}] 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.

[^{45}] Indent added by Decision No 83/2014 (OJ L 310, 30.10.2014, p. 36 and EEA Supplement No 63, 30.10.2014, p. 27), e.i.f. 17.5.2014.


[^{47}] 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.


[^{49}] Indent added by Decision No 219/2014 (OJ L 230, 3.9.2015, p. 18 and EEA Supplement No 52, 3.9.2015, p. 18), e.i.f. 1.11.2014.


[566] Indent added by Decision No 65/2019 (OJ L 210, 2.7.2020, p. 31 and EEA Supplement No 44, 2.7.2020, p. 35), e.i.f. 30.3.2019.


32022 R 0586: Commission Regulation (EU) 2022/586 of 8 April 2022 (OJ L 112, 11.4.2022, p. 6),


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:

Indent added by Decision No 14/2024 (OJ L, 2024/1554, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 28), e.i.f. 3.2.2024.

Indent added by Decision No 66/2024 (OJ L, 2024/1580, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 42), e.i.f. 16.3.2024.

Indent added by Decision No 91/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

Indent added by Decision No 92/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

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

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

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

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

“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. [ ]


12zt. [ ] [609]


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


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Footnotes:


(331) 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.


(335) 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.

(336) 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.


Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 10.12.2021


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

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

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

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

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

(ii) Article 51, in conjunction with Articles 4 and 46(1), with respect to the requirements for the classification, labelling and/or specific concentration limits for the substances or group of substances 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:
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.

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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chlorine released from chlorine as an existing active substance for use in biocidal products of product-types 2

acid generated from tetraacetyldiamine and sodium percarbonate as an existing active substance for

12zzzz.[32017 R 1277] Commission Implementing Regulation (EU) 2017/1277 of 14 July 2017 approving 2-octyl-
isothiazol-3(2H)-one as an active substance for use in biocidal products of product-type 8 (OJ L 184,
15.7.2017, p. 27).

12zzzz.[32017 R 1278] Commission Implementing Regulation (EU) 2017/1278 of 14 July 2017 approving 2-
methylisothiazol-3(2H)-one as an existing active substance for use in biocidal products of product-type 11

12zzzz.[32017 D 1282] Commission Implementing Decision (EU) 2017/1282 of 14 July 2017 not approving 2-
methyl-1,2-benzisothiazol-3(2H)-one as an active substance for use in biocidal products of product-type 13

approval of warfarin as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of chlorophacinone as an active substance for use in biocidal products of product-type 14 (OJ L

approval of coumatetralyl as an active substance for use in biocidal products of product-type 14 (OJ L

approval of difenacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of bromadiolone as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of brodifacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of difethialone as an active substance for use in biocidal products of product-type 14 (OJ L 194,

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chlorine released from chlorine as an existing active substance for use in biocidal products of product-types 2

acid generated from tetraacetyldiamine and sodium percarbonate as an existing active substance for

[32017 R 1277] Commission Implementing Regulation (EU) 2017/1277 of 14 July 2017 approving 2-octyl-
isothiazol-3(2H)-one as an active substance for use in biocidal products of product-type 8 (OJ L 184,
15.7.2017, p. 27).

[32017 R 1278] Commission Implementing Regulation (EU) 2017/1278 of 14 July 2017 approving 2-
methylisothiazol-3(2H)-one as an existing active substance for use in biocidal products of product-type 11

[32017 D 1282] Commission Implementing Decision (EU) 2017/1282 of 14 July 2017 not approving 2-
methyl-1,2-benzisothiazol-3(2H)-one as an active substance for use in biocidal products of product-type 13

approval of warfarin as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of chlorophacinone as an active substance for use in biocidal products of product-type 14 (OJ L

approval of coumatetralyl as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of difenacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of bromadiolone as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of brodifacoum as an active substance for use in biocidal products of product-type 14 (OJ L 194,

approval of difethialone as an active substance for use in biocidal products of product-type 14 (OJ L 194,


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12zzzzzzb.[85] 2019 D 1331: Commission Implementing Decision (EU) 2019/1331 of 5 August 2019 on the terms and conditions of the authorisation of a biocidal product containing peppermint oil and citronellal referred by

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\[\text{\textsuperscript{[76]}}\] Point inserted by Decision No 11/2020 (OJ L 49, 16.2.2023, p. 27 and EEA Supplement No 13, 16.2.2023, p. 28), e.i.f. 8.2.2020.

\[\text{\textsuperscript{[77]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[78]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[79]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[80]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[81]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[82]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[83]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[84]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[85]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[86]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[87]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[88]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.

\[\text{\textsuperscript{[89]}}\] Point inserted by Decision No 60/2020 (OJ L 72, 9.3.2023, p. 24 and EEA Supplement No 19, 9.3.2023, p. 25), e.i.f. 1.5.2020.


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\(^{(82)}\) Point inserted by Decision No 206/2021 (OJ L, 2024/305, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 37), e.i.f. 10.7.2021.

\(^{(83)}\) Point inserted by Decision No 206/2021 (OJ L, 2024/305, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 37), e.i.f. 10.7.2021.


Commission Implementing Decision (EU) 2021/1287 of 2 August 2021 postponing the expiry date of approval of active chlorine generated from sodium chloride by electrolysis as an active substance for use in biocidal products of product-type 1 (OJ L 70, 1.3.2021, p. 6).


Commission Implementing Decision (EU) 2021/1285 of 2 August 2021 postponing the expiry date of approval of active chlorine released from hypochlorous acid as an active substance for use in biocidal products of product-type 1 (OJ L 70, 1.3.2021, p. 9).


Commission Implementing Decision (EU) 2021/1283 of 2 August 2021 postponing the expiry date of approval of active chlorine generated from sodium chloride by electrolysis as an active substance for use in biocidal products of product-type 1 (OJ L 70, 1.3.2021, p. 6).


Commission Implementing Decision (EU) 2021/2149 of 3 December 2021 on unresolved objections regarding the terms and conditions of the provisional authorisation of a biocidal product containing 5-Chloro-2-methyl-2H-isothiazol-3-one (C(M)IT) referred by France in accordance with Article 36(1) of Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 434, 6.12.2021, p. 5).


Commission Implementing Decision (EU) 2022/0146 of 1 February 2022 determining whether a product containing Alkyl (C12-16) dimethylbenzyl ammonium chloride is a biocidal product,
pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 24, 3.2.2022, p. 133).


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


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

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

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


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[^50]: Point inserted by Decision No 266/2023 (OJ L, 2024/1148, 16.5.2024 and EEA Supplement No 32, 16.5.2024, p. 35), e.i.f. 28.10.2023.


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\[\text{ee}^3\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^4\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^5\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^6\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^7\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^8\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^9\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^{10}\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^{11}\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 34), e.i.f. 16.3.2024.

\[\text{ee}^{12}\]\ Point inserted by Decision No 62/2024 (OJ L, 2024/1597, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 37), e.i.f. 16.3.2024.

\[\text{ee}^{13}\]\ Point inserted by Decision No 91/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.


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**(87)** Point inserted by Decision No 92/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87a)** Point inserted by Decision No 92/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87b)** Point inserted by Decision No 93/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87c)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87d)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87e)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87f)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87g)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87h)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87i)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(87j)** Point inserted by Decision No 94/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.

**(88)** Point inserted by Decision No 117/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2024.

**(88a)** Point inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.i.f. 1.6.2015.

**(88b)** Indent and words “, as amended by:" added by Decision No 18/2015 (OJ L 93, 7.4.2016, p. 30 and EEA Supplement No 21, 7.4.2016, p. 26), e.i.f. pending. It shall apply provisionally pending the entry into force of the 2014 EEA Enlargement Agreement, from the day of entry into force of Decision of the Joint Committee No 203/2014 which e.i.f. 1.6.2015.

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

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

“Iceland, Norway”

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

“Liechtenstein”


Adaptations (i) to (j) are renumbered as adaptations (j) to (k) and adaptation text (i) inserted by Decision 77/2023 (OJ L, 2023/02233, 9.11.2023 and EEA Supplement No 81, 9.11.2023, p. 3), e.i.f. pending


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


(192) 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.

(193) 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.

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

(195) 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.

(196) 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.

(197) 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.

(198) 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.

(199) 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.

(200) 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.

(201) 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.

(202) 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.

(203) 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.

(204) 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.

(205) 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.

(206) 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.

(207) 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.

(208) 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.

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


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(**) 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.

(***) 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.

(****) 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.

(***** 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.

(******) 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.

(******* 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.

(******** 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.

********* 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.

********** 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.


(^[#16]^) 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.

(^[#17]^) 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.

(^[#18]^) 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.

(^[#19]^) 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.

(^[#20]^) 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.

(^[#21]^) 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.


Footnotes:

(97) 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.

(98) 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.

(99) 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.

(100) 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.

(101) 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.

(102) 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.

(103) 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.

(104) 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.

(105) 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.

(106) 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.

(107) 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.

(108) 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.


(110) Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 11.12.2015.

(111) 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.


Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (OJ L 202, 30.7.2015, p. 92 and EEA Supplement No 43, 30.7.2015, p. 91), e.i.f. 1.6.2015.

Indent added by Decision No 220/2014 (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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\[1006\] Indent added by Decision No 221/2014 (OJ L 230, 3.9.2015, p. 22 and EEA Supplement No 52, 3.9.2015, p. 22), e.i.f. 1.6.2015 and subsequently deleted by Decision No 17/2024 (OJ L 2024/1557, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 34), e.i.f. 3.2.2024.

\[1007\] Indent added by Decision No 221/2014 (OJ L 230, 3.9.2015, p. 22 and EEA Supplement No 52, 3.9.2015, p. 22), e.i.f. 1.6.2015.

\[1008\] Indent added by Decision No 221/2014 (OJ L 230, 3.9.2015, p. 22 and EEA Supplement No 52, 3.9.2015, p. 22), e.i.f. 1.6.2015.

\[1009\] Indent added by Decision No 221/2014 (OJ L 230, 3.9.2015, p. 22 and EEA Supplement No 52, 3.9.2015, p. 22), e.i.f. 1.6.2015.

\[1010\] 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.

\[1011\] 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.

\[1012\] 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.

\[1013\] 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.

\[1014\] 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.

\[1015\] 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.

\[1016\] 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.

\[1017\] 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.

\[1018\] 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.


\[^{1020}\] Indent added by Decision No 94/2015 (OJ L 211, 4.8.2016, p. 32 and EEA Supplement No 42, 4.8.2016, p. 31), e.i.f. 1.6.2015.

\[^{1021}\] 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.

\[^{1022}\] 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. Corrigendum to the EU act subsequently taken note of by the EEA Joint Committee on 10.6.2022.

\[^{1023}\] 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.

\[^{1024}\] 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.

\[^{1025}\] 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.


\[^{1031}\] 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.

\[^{1032}\] 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.


\[\text{Indent added by Decision No 182/2016} (OJ L 80, 22.3.2018, p. 26 and EEA Supplement No 19, 22.3.2018, p. 34), \text{e.i.f. 24.9.2016.}\]

\[\text{Indent added by Decision No 183/2016} (OJ L 80, 22.3.2018, p. 27 and EEA Supplement No 19, 22.3.2018, p. 36), \text{e.i.f. 24.9.2016.}\]

\[\text{Indent added by Decision No 183/2016} (OJ L 80, 22.3.2018, p. 27 and EEA Supplement No 19, 22.3.2018, p. 36), \text{e.i.f. 24.9.2016.}\]


\[\text{Indent added by Decision No 30/2017} (OJ L 297, 22.11.2018, p. 38 and EEA Supplement No 78, 22.11.2018, p. 44), \text{e.i.f. 4.2.2017.}\]

\[\text{Indent added by Decision No 30/2017} (OJ L 297, 22.11.2018, p. 38 and EEA Supplement No 78, 22.11.2018, p. 44), \text{e.i.f. 4.2.2017.}\]

\[\text{Indent added by Decision No 30/2017} (OJ L 297, 22.11.2018, p. 38 and EEA Supplement No 78, 22.11.2018, p. 44), \text{e.i.f. 4.2.2017.}\]

\[\text{Indent added by Decision No 30/2017} (OJ L 297, 22.11.2018, p. 38 and EEA Supplement No 78, 22.11.2018, p. 44), \text{e.i.f. 4.2.2017.}\]


- [1139] [ ] (1139)


\footnote{1155} 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.
\footnote{1156} 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.
\footnote{1157} 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.
\footnote{1158} 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.
\footnote{1159} 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.
\footnote{1160} 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.
\footnote{1161} Indent added by Decision No 119/2019 (OJ L 279, 27.10.2022, p. 16 and EEA Supplement No 69, 27.10.2022, p. 17), e.i.f. 1.6.2019.
\footnote{1162} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1163} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1164} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1165} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1166} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1167} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1168} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.
\footnote{1169} Indent added by Decision No 120/2019 (OJ L 279, 27.10.2022, p. 19 and EEA Supplement No 69, 27.10.2022, p. 19), e.i.f. 1.6.2019.


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


- (1214) 32020 R 0617: Commission Implementing Regulation (EU) 2020/617 of 5 May 2020 (OJ L 143, 6.5.2020, p. 6),

\[\text{(126)}\] 32020 R 0646: Commission Implementing Regulation (EU) 2020/646 of 13 May 2020 (OJ L 151, 14.5.2020, p. 3),


\[\text{(130)}\] 32020 R 1004: Commission Implementing Regulation (EU) 2020/1004 of 9 July 2020 (OJ L 221, 10.7.2020, p. 133),


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-\{1246\} 32021 R 0574: Commission Implementing Regulation (EU) 2021/574 of 30 March 2021 (OJ L 120, 8.4.2021, p. 9),


-\{1255\} Indent added by Decision No 67/2022 (OJ L 182, 7.7.2022, p. 46 and EEA Supplement No 45, 7.7.2022, p. 34), e.i.f. 19.3.2022.

-\{1256\} Indent added by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), e.i.f. 30.4.2022.

-\{1257\} Indent added by Decision No 127/2022 (OJ L 246, 22.9.2022, p. 73 and EEA Supplement No 61, 22.9.2022, p. 72), e.i.f. 30.4.2022.

-\{1258\} Indent added by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.


- (1247) 32022 R 0004: Commission Implementing Regulation (EU) 2022/4 of 4 January 2022 (OJ L 1, 5.1.2022, p. 5),

- (1248) 32022 R 0019: Commission Implementing Regulation (EU) 2022/19 of 7 January 2022 (OJ L 5, 10.1.2022, p. 9),


- (1250) 32022 R 0159: Commission Implementing Regulation (EU) 2022/159 of 4 February 2022 (OJ L 26, 7.2.2022, p. 7),


- (1252) 32022 R 0708: Commission Implementing Regulation (EU) 2022/708 of 5 May 2022 (OJ L 133, 10.5.2022, p. 1),

- (1253) 32022 R 0698: Commission Implementing Regulation (EU) 2022/698 of 3 May 2022 (OJ L 130, 4.5.2022, p. 3),

- (1254) 32022 R 0800: Commission Implementing Regulation (EU) 2022/800 of 20 May 2022 (OJ L 143, 23.5.2022, p. 4),

[1246] Indent added by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.


[1242] Indent added by Decision No 129/2022 (OJ L 246, 22.9.2022, p. 77 and EEA Supplement No 61, 22.9.2022, p. 76), e.i.f. 30.4.2022.


- [1313] Indent added by Decision No 16/2024 (OJ L, 2024/1553, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 32), e.i.f. 3.2.2024.


(1305) Indent added by Decision No 17/2024 (OJ L, 2024/1557, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 34), e.i.f. 3.2.2024.

(1306) Indent added by Decision No 17/2024 (OJ L, 2024/1557, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 34), e.i.f. 3.2.2024.

(1307) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1308) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1309) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1310) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1311) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1312) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1313) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1314) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1315) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1316) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1317) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1318) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

(1319) Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.
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.

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

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


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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 ljósnæmingu.

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

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

“IS: Efnið brennir húð og augi 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 yfirborðsvatni/Þoma skal í veg fyrir að mengun verði með afretni frá højarhlóðum og vegum.)

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

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

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

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

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

“IS: Þvoið allan hlífðarfatnað að lokinni notkun.

NO: Vask alt 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: Forbist innöndun reyks eftir að kveikt hefur verið í efninu og yfirgefið þegar í stað svæðið sem er til meðhöndlunar.


{1345} Point and adaptation text inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.i.f. 1.6.2015.
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 ílátíð 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: Loftræsta skal úðuð svæði/þurr í tilgreindan tíma/þar til úðinn hefur þornað) áður en farið er þangað inn aftur.

NO: De behandlede områder/vektathers ventileres (grundig/eller angivelse av tid/inntil produktet har tørket) 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/járðvegslífverur skal ekki nota þetta eða annað efnir som inniheldur (tilgreinið virkt efnir eða flokk virkra efnir eftir því sem við á) lengur eða oftur en (tilgreinið hversu lengi eða oft má nota efnin).

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/vatnallífverur skal ekki nota þetta efnir (á tilgreinda járðvegsgerð eða við tilgreindar aðstæður).

NO: For å beskytte (grunnvannet/vannlevende organismer) må dette produktet ikke brukes (på beskrevet jortype 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 vatnallífverur/plöntur utan markhóps/ljöður utan markhóps/skordýr má ekki nota efnir nær örekutsuð land/þurfsvatnini en (tilgreinu breidd svæðissem er óheimilt að úða).

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

(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 vatnallífverur/plöntur utan markhóps má ekki nota efnir á malbikað, steinsteypt, hellulagð eða malarbordið yfirborð eða vegi (jarnbrautar spor) eða önnur svæðir þar sem hætt er við afrensla út í umhverfið.

NO: For å beskytte (vannlevende organismer/viltlevende planter) må dette produktet ikke brukes på harde overflater som asfalterte, 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 fuglar/víllit spendýr verður að gæta þess vandlega að efnin sé algerlega hulið járðvegi; getið þess séstaklega að efnin sé hulið í endum raða.

NO: For å beskytte (fugler/ville pattedyr) skal produktet innblandes 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: Hreinsíð upp allt efni, sem hefur farið til spillis, til að vernda fugla/villt 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 efnið á varptíma fugla.

NO: Må ikke brukes i fuglenes hekketid.”

(q) 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 frævandi skordýr er óheimilt að nota efnið á blómstrandi nytjaplöntur/Óheimilt er að nota efnið þar sem blómstrug eru í fæðuleiti/Fjarlægið býkúpur meðan meðhöndlun með efniðu fer fram eða hyljið þær á meðan og í (tilgreiníð tíma) að lokinni meðhöndlun/Óheimilt er að nota efnið ef blómstrandi illgresi er til staðar/Eyða skal illgresi aður en það blómst og/Óheimilt er að nota efnið fyrir (tilgreiníð 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) etter behandlingen./Må ikke brukes i nærheten av blomstrende ugress./Fjern ugresset før det blomstrer./Må ikke brukes før (tidspunkt).”

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

“IS: Til að koma í veg fyrir þolmyndun skal ekki nota þetta eða annan varnarefni sem inniheldur (tilgreiníð virkt efni eða flokk virksom efna eftir því sem við á) oftar eða lengur en (tilgreiníð hversu oft eða lengi má nota efnið).

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

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

“IS: Beitu skal komið fyrir þannig að ekki sé hætta á að önnur dýr komist í hans. Festa skal beituna trygglæga þannig að nagdýr geti ekki dregið 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 velk av de gnagere som skal bekjempes.”

(t) 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 voldum storkuvarex 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.”

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

“IS: Hræ nagdýra skulu fjarlægi daglega af meðhöndlada svæðinu meðan meðhöndlun stendur yfir. Ekki skal setja hræin i opin sorpilät.

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

13f.{1346} 32020 R 1740: Commission Implementing Regulation (EU) 2020/1740 of 20 November 2020 setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in

{1346} Point inserted by Decision No 203/2014 (OJ L 202, 30.7.2015, p. 57 and EEA Supplement No 43, 30.7.2015, p. 57), e.i.f. 1.6.2015 and subsequently replaced by Decision No 341/2021 (OJ L, 2024/658, 14.3.2024 and EEA Supplement No 23, 14.3.2024, p. 58), e.i.f. pending.


13x.\{1366\} 32011 R 0942: Commission Implementing Regulation (EU) No 942/2011 of 22 September 2011 concerning the nonApproval of the active substance flufenoxuron, in accordance with Regulation (EC) No 1107/2009 of


\{1365\} Indent and words “as amended by:” added by Decision No 342/2015 (OJ L [to be published] and EEA Supplement No [to be published]), c.f. 11.12.2021.


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


(1378) 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.


(1380) 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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[137] 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.


[139] 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.

[139] 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.

[139] 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.

[152] Insert and words “as amended by:” added by Decision No 34/2020 (OJ L 57, 23.2.2020, p. 10) and EEA Supplement No 16, 23.2.2020, p. 10), e.i.f. 21.3.2020.

[139] 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.

[139] 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.

[139] 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.


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13zzze.\textsuperscript{1426}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:


\textsuperscript{-1427}32015 R 0052: Commission Implementing Regulation (EU) 2015/52 of 14 January 2015 (OJ L 9, 15.1.2015, p. 27),


\textsuperscript{-1435}32023 R 0543: Commission Implementing Regulation (EU) 2023/543 of 9 March 2023 (OJ L 73, 10.3.2023, p. 1),


13zzzf.\textsuperscript{1437}Commission Implementing Decision 2013/38/EU of 18 January 2013 allowing Member States to extend provisional authorisations granted for the new active substances enamectin and maltodextrin (OJ L 18, 22.1.2013, p. 17).


\textsuperscript{1425}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{1426}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{1427}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{1428}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{1429}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{1430}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{1431}Indent added by Decision No 229/2019 (OJ L 4, 5.1.2023, p. 46 and EEA Supplement No 3, 5.1.2023, p. 46), e.i.f. 28.9.2019.


\textsuperscript{1435}Indent added by Decision No 19/2024 (OJ L, 2024/1531, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 41), e.i.f. 3.2.2024.

\textsuperscript{1436}Indent added by Decision No 65/2024 (OJ L, 2024/1625, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 40), e.i.f. 16.3.2024.

\textsuperscript{1437}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{1438}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.
13zzh.\[^{1439}\] 32013 D 0205: Commission Implementing Decision 2013/205/EU of 25 April 2013 allowing Member States to extend provisional authorisations granted for the new active substances acequinocyl, aminopyralid, ascorbic acid, flubendiamide, gamma-cyhalothrin, ipconazole, metaflumizone, orthosulfamuron, \textit{Pseudomonas} sp. strain DSMZ 13134, pyridalil, pyroxasulam, spiroxifenes, thiencarbazone and topramezone (OJ L 117, 27.4.2013, p. 20).

13zzi.\[^{1440}\] 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:


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\(^{[1480]}\) 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.
\(^{[1481]}\) 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.
\(^{[1482]}\) 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.
\(^{[1483]}\) 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.
\(^{[1484]}\) 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.
\(^{[1485]}\) 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.
\(^{[1486]}\) 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.


13zzzrd. [ ] {[415]}


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

{[415]} Point inserted by Decision No 221/2014 (OJ L 230, 3.9.2015, p. 22 and EEA Supplement No 52, 3.9.2015, p. 22), e.i.f. 1.6.2015 and subsequently deleted by Decision No 17/2024 (OJ L 284/1557, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 34), e.i.f. 3.2.2024.


{[419]} Point inserted 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.


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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), e.i.f. 1.6.2015.


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


\begin{itemize}
  \item \textbf{32020 R 1295}: Commission Implementing Regulation (EU) 2020/1295 of 16 September 2020 (OJ L 303, 17.9.2020, p. 18),
  \item \textbf{32022 R 0043}: Commission Implementing Regulation (EU) 2022/43 of 13 January 2022 (OJ L 9, 14.1.2022, p. 7),
  \item \textbf{32022 R 1252}: Commission Implementing Regulation (EU) 2022/1252 of 19 July 2022 (OJ L 191, 20.7.2022, p. 41),
\end{itemize}


\footnote{1479} 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), e.i.f. 11.7.2015.
\footnote{1480} 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), e.i.f. 11.7.2015.
\footnote{1482} Indent and words “, as amended by” added by Decision No 69/2018 (OJ L 340, 15.10.2020, p. 20 and EEA Supplement No 66, 15.10.2020, p. 25), e.i.f. 28.4.2018.
\footnote{1483} Indent added by Decision No 33/2020 (OJ L 57, 23.2.2023, p. 7 and EEA Supplement No 16, 23.2.2023, p. 7), e.i.f. 21.3.2020.
\footnote{1484} Indent added by Decision No 33/2021 (OJ L 85, 30.3.2021 and EEA Supplement No 3, 11.1.2024, p. 68), e.i.f. 6.2.2021.
\footnote{1485} Indent added by Decision No 67/2022 (OJ L 182, 7.7.2022, p. 46 and EEA Supplement No 45, 7.7.2022, p. 34), e.i.f. 19.3.2022.
\footnote{1488} Indent added by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.
\footnote{1489} 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.

Point inserted e.i.f. Indent and words “, as amended by:” added e.i.f.


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1. **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.**
2. **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.**
3. **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.**
4. **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.**
5. **Point inserted by Decision No 265/2015 (OJ L 161, 22.6.2017, p. 48 and EEA Supplement No 38, 22.6.2017, p. 50), e.i.f. 1.11.2015.**
7. **Indent and words “, as amended by:” added by Decision No 229/2019 (OJ L 4, 5.1.2023, p. 46 and EEA Supplement No 3, 5.1.2023, p. 46), e.i.f. 28.9.2019.**
8. **Point inserted by Decision No 265/2015 (OJ L 161, 22.6.2017, p. 48 and EEA Supplement No 38, 22.6.2017, p. 50), e.i.f. 1.11.2015.**


13zzzzzm.**[1509]** 32015 R 2082: Commission Implementing Regulation (EU) 2015/2082 of 18 November 2015 concerning the non-approval of *Arctium lappa* L. (aerial parts) as a basic substance in accordance with Regulation

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3zzzzzzw. 32017 R 0240: Commission Implementing Regulation (EU) 2017/240 of 10 February 2017 concerning the non-approval of Satureja montana L. essential oil as a basic substance in accordance with Regulation

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Point inserted by Decision No 14/2019 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 9.2.2019.


13zzzzzzw.\textsuperscript{[175]}\textsuperscript{32019 R 0481}: Commission Implementing Regulation (EU) 2019/481 of 22 March 2019 approving the active substance flutianil, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and


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13zzzzzzzzzo.\textsuperscript{[1670]} Commission Implementing Regulation (EU) 2021/134 of 4 February 2021 renewing the approval of the low-risk active substance Akanthomyces muscardus strain Ve6 (formerly Lecanicillium muscarium strain Ve6) in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and


\textsuperscript{[1663]} Point inserted by Decision No 208/2021 (OJ L, 2024/303, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 40), e.i.f. 10.7.2021.

\textsuperscript{[1664]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1665]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1666]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1667]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1668]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1669]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.

\textsuperscript{[1670]} Point inserted by Decision No 239/2021 (OJ L, 2024/502, 22.2.2024 and EEA Supplement No 17, 22.2.2024, p. 21), e.i.f. 25.9.2021.


13zzzzzzzzzg.\[32021 R 2049\]: Commission Implementing Regulation (EU) 2021/2049 of 24 November 2021 renewing the approval of the active substance cypermethrin as a candidate for substitution in accordance


\[32021 R 1191\] Point inserted by Decision No 67/2022 (OJ L 182, 7.7.2022, p. 46 and EEA Supplement No 45, 7.7.2022, p. 34), e.i.f. 19.3.2022.

\[32021 R 1448\] Point inserted by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), e.i.f. 30.4.2022.

\[32021 R 1379\] Point inserted by Decision No 126/2022 (OJ L 246, 22.9.2022, p. 71 and EEA Supplement No 61, 22.9.2022, p. 70), e.i.f. 30.4.2022.

\[32021 R 1452\] Point inserted by Decision No 18/2024 (OJ L 204/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

\[32021 R 1455\] Point inserted by Decision No 127/2022 (OJ L 246, 22.9.2022, p. 73 and EEA Supplement No 61, 22.9.2022, p. 72), e.i.f. 30.4.2022.

\[32021 R 1451\] Point inserted by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

\[32021 R 2049\] Point inserted by Decision No 128/2022 (OJ L 246, 22.9.2022, p. 75 and EEA Supplement No 61, 22.9.2022, p. 74), e.i.f. 30.4.2022.

\[32021 R 0428\] Point inserted by Decision No 129/2022 (OJ L 246, 22.9.2022, p. 77 and EEA Supplement No 61, 22.9.2022, p. 76), e.i.f. 30.4.2022.


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Commission Implementing Regulation (EU) 2022/456 of 21 March 2022


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{1719} Point inserted by Decision No 16/2024 (OJ L, 2024/1553, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 32), e.i.f. 3.2.2024.

{1720} Point inserted by Decision No 17/2024 (OJ L, 2024/1557, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 34), e.i.f. 3.2.2024.

{1721} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

{1722} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

{1723} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

{1724} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

{1725} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.

{1726} Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), e.i.f. 3.2.2024.


Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), c.l.f. 3.2.2024.

Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), c.l.f. 3.2.2024.

Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), c.l.f. 3.2.2024.

Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), c.l.f. 3.2.2024.

Point inserted by Decision No 18/2024 (OJ L, 2024/1551, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 36), c.l.f. 3.2.2024.

Point inserted by Decision No 46/2024 (OJ L, 2024/1575, 4.7.2024 and EEA Supplement No 52, 4.7.2024, p. 38), c.l.f. 16.3.2024.


Point inserted by Decision No 118/2024 (OJ L [to be published] and EEA Supplement No [to be published]), c.l.f. 13.6.2024.

Point inserted by Decision No 118/2024 (OJ L [to be published] and EEA Supplement No [to be published]), c.l.f. 13.6.2024.


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(174) Point inserted by Decision No 118/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 13.6.2024.
(1738) Point inserted by Decision No 119/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 13.6.2024.
(1739) Point inserted by Decision No 160/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 6.7.2024.
(1740) Point inserted by Decision No 161/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 6.7.2024.
(1741) Point inserted by Decision No 162/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 6.7.2024.
(1742) Point inserted by Decision No 161/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.l.f. 6.7.2024.
(1745) Point inserted by Decision No 54/2015 (OJ L 129, 19.5.2015, p. 36 and EEA Supplement No 29, 19.5.2015, p. 37), e.l.f. 21.3.2015.
The Contracting Parties take note of the content of the following acts:

1. [ ]


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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. {1769} 32006 H 0283: Commission Recommendation of 11 April 2006 on risk reduction measures for the substances: Dibutylphthalate; 3,4-Dichloroaniline; Diisodecyl phthalate; 1,2-Benzenedicarboxylic acid, di-C9-11-branched alkyl esters, C10-rich; Di-isononyl phthalate; 1,2-Benzenedicarboxylic acid, di-C8-10-branched alkyl esters, C9-rich; Ethylenediaminetetraacetate; Methyl acetate; Monochloroacetic acid; n-Pentane; Tetrasodium ethylenediaminetetraacetate (OJ L 104, 13.4.2006, p. 45).

XVI. COSMETICS

ACTS REFERRED TO

1. [ ] (1779)


{1772} 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.i.f. 9.11.2013.

{1773} Indent added by Decision No 196/2013 (OJ L [to be published] and EEA Supplement No [to be published]), c.i.f. 9.11.2013.


{1777} Indent added by Decision No 19/2015 (OJ L 93, 7.4.2016, p. 32 and EEA Supplement No 21, 7.4.2016, p. 28), c.i.f. 26.2.2015.

{1778} Indent added by Decision No 20/2015 (OJ L 93, 7.4.2016, p. 33 and EEA Supplement No 21, 7.4.2016, p. 29), c.i.f. 26.2.2015.


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\{1812\} Indent added by Decision No 68/2022 (OJ L 182, 7.7.2022, p. 48 and EEA Supplement No 45, 7.7.2022, p. 36), e.i.f. 19.3.2022.
\{1813\} Indent added by Decision No 130/2022 (OJ L 246, 22.9.2022, p. 79 and EEA Supplement No 61, 22.9.2022, p. 78), e.i.f. 30.4.2022.
\{1816\} Indent added by Decision No 45/2023 (OJ L 2023/0248, 9.11.2023 and EEA Supplement No 81, 9.11.2023, p. 35), e.i.f. 4.9.2022.
\{1818\} Indent added by Decision No 269/2023 (OJ L 2023/0434, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 43), e.i.f. 28.10.2023.
\{1819\} Indent added by Decision No 270/2023 (OJ L 2023/0434, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 44), e.i.f. 28.10.2023.

9. [ ] [1825]

10. [ ] [1826]

11. [1827] [ ] [1828]

12. [1829] [ ] [1830]

13. [1831] [ ] [1832]


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE**[1834]

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


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XVII. ENVIRONMENT PROTECTION

ACTS REFERRED TO

1. [ ] \footnote{1836} 
2. [ ] \footnote{1837} 
3. [ ] \footnote{1838} 
4. [ ] \footnote{1839} 
5. [ ] \footnote{1840} 
6. [ ] \footnote{1841} 


\footnote{1846} 32011 L 0063: Commission Directive 2011/63/EU of 1 June 2011 (OJ L 147, 2.6.2011, p. 15),

\footnote{1846} Indent added by Decision No 74/2013 (OJ L 291, 31.10.2013, p. 31 and EEA Supplement No 61, 31.10.2013, p. 35), e.i.f. 1.11.2015.
\footnote{1847} Indent added by Decision No 55/2015 (OJ L 129, 19.5.2016, p. 38 and EEA Supplement No 29, 19.5.2016, p. 39), e.i.f. 1.11.2015.
The provisions of the Directive shall, for the purposes for 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. [ ]

6ab. [ ]

6ac. [ ]

6ad. [ ]


6af. [ ]

6ag. [ ]


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**1815** Point inserted by Decision No 156/2019 (OJ L 291, 10.11.2022, p. 36 and EEA Supplement No 74, 10.11.2022, p. 37), e.i.f. 15.6.2019.

**1816** Point inserted by Decision No 157/2019 (OJ L 291, 10.11.2022, p. 37 and EEA Supplement No 74, 10.11.2022, p. 38), e.i.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.


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”.


{(1905) 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 “or, and for Iceland for a period of 36 months following the latter date in Annex II, to finish the Icelandic stocks” shall be inserted after the word “force”.

9a. [ ]

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 Decision of the EEA Joint Committee No 160/2019 of 14 June 2019”.

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

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

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


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

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

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

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


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

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


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

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


{1925} 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.

{1926} 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.

{1927} 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.

9da. (\textsuperscript{32021 D 0958}) 32021 D 0958: Commission Implementing Decision (EU) 2021/958 of 31 May 2021 laying down the format for reporting data and information on fishing gear placed on the market and waste fishing gear collected in Member States and the format for the quality check report in accordance with Articles 13(1)(d) and 13(2) of Directive (EU) 2019/904 of the European Parliament and of the Council (OJ L 211, 15.6.2021, p. 51), as corrected by OJ L 34, 16.2.2022, p. 52.


9dc. (\textsuperscript{32022 D 0162}) 32022 D 0162: Commission Implementing Decision (EU) 2022/162 of 4 February 2022 laying down rules for the application of Directive (EU) 2019/904 of the European Parliament and of the Council as regards the calculation, verification and reporting on the reduction in the consumption of certain single-use plastic products and the measures taken by Member States to achieve such reduction (OJ L 26, 7.2.2022, p. 19), as amended by:


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


The provisions of the Directive shall, for the purposes of this Agreement, be read with the following adaptation: Articles 7, 11 and 16 shall not apply to Liechtenstein.

The provisions of the Regulation shall, for the purposes of this Agreement, be read with the following adaptation:
This Regulation shall not apply to Liechtenstein.


**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE** {1938}

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


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

**ACTS REFERRED TO**

1. [ ]{1939}

2. [ ]{1940}


4. [ ]{1941}

4a. [ ]{1942} 4b. [ ]{1943}

4b. [ ]{1944} 4b. [ ]{1945}

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4c. [1946] [ ]
4d. [1947] [ ]
4e. [1948] [ ]
4f. [1949] [ ]
4g. [1950] [ ]
4h. [1951] [ ]
4i. [ ] [1952]
4j. [ ] [1953]
4k. [ ] [1954]
4l. [ ] [1955]
4m. [1956] [ ] [1957]
4n. [ ] [1958]


4o. [1959]


4u. [ ] [1965]

4v. [ ] [1966]


4x. [ ] [1968]


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4ze. [ ]{\[1975\]}

4zf. [ ]{\[1976\]}

4zg. [ ]{\[1977\]}


4zu {1992} 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).


4xz {1990} 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).


4zhb {1999} 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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(1) Listed here for purposes of information only. For application, see Annex XIV on competition.

4zzq.\textsuperscript{[2012]} 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 (GMSS) (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)”.

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**


\[2017\] Indent and words “, as amended by:” added by Decision No 234/2021 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.


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

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

“The term ‘technical specification’ also covers production methods and processes used in respect of products intended for human and animal consumption, and in medicinal products as defined in Article 1 of Directive 2001/83/EC (as incorporated into point 15 q of Chapter XIII of Annex II to the Agreement by Decision of the EEA Joint Committee No 82/2002 of 25 June 2002\(^{[2023]}\)), 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.\(^{[2025]}\)

3a.\(^{[2026]}\)


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.


- 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),


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

(a) In Annex I, point 1(a) the following shall be added to the list of written indications concerning "upper":
   IS Efri hluti
   N Overdel
   \[2036\]

(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 bindssóli
   N För og bindsåle
   \[2037\]

(c) in Annex I, point 1(c) the following shall be added to the list of written indications concerning "outer sole":
   IS Slitsóli
   N Yttersåle
   \[2038\]

(d) In Annex I, point 2(a)(i) the following shall be added to the list of written indications concerning "leather":
   IS Leður
   N Lær
   \[2039\]

(e) In Annex I, point 2(a)(ii) the following shall be added to the list of written indications concerning "coated leather":
   IS Húðað leður
   N Belagt lær
   \[2040\]

(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
   \[2041\]

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

\[2035\] 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.
IS Ölj önnur efni
N Andre materialer
[ ]**(343)**,

3f. [ ]**(344)**


-**(345)** 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),

-**(346)** 1 94 N: Act concerning the conditions of accession of the Republic of Austria, the Republic of Finland and the Kingdom of Sweden and the adjustments of 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.**(352)** 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:


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


\[2053\] 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. 141), e.i.f. 11.11.2010.

\[2054\] 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. 141), e.i.f. 11.11.2010.

\[2055\] 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. 141), e.i.f. 11.11.2010.

\[2056\] 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. 141), e.i.f. 11.11.2010.


\[2058\] Indent added by Decision No 275/2014 (OJ L 311, 26.11.2015, p. 28 and EEA Supplement No 71, 26.11.2015, p. 27), e.i.f. 13.12.2014.

\[2059\] Indent added by Decision No 97/2015 (OJ L 211, 4.8.2016, p. 37 and EEA Supplement No 42, 4.8.2016, p. 36), e.i.f. 1.5.2015.


\[2063\] 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.

\[2064\] Indent added by Decision No 196/2012 (OJ L 21, 24.1.2013, p. 44 and EEA Supplement No 6, 24.1.2013, p. 10), e.i.f. 1.11.2012.

\[2065\] Point inserted by Decision No 196/2012 (OJ L 21, 24.1.2013, p. 44 and EEA Supplement No 6, 24.1.2013, p. 10), e.i.f. 1.11.2012.


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.

3r.\textsuperscript{[277]} 32011 D 0477: Commission Decision 2011/477/EU of 27 July 2011 on the safety requirements to be met by European standards to address certain risks posed to children by internal blinds, corded window coverings and related products within the Community (OJ L 170, 6.7.2010, p. 39), as amended by:

- 32011 D 0677: Commission Decision 2011/677/EU of 31 July 2011 on the safety requirements to be met by European standards to address certain risks posed to children by internal blinds, corded window coverings and related products within the Community (OJ L 311, 25.11.2011, p. 1),
- \textsuperscript{[266]} Point inserted by Decision No 196/2012 (OJ L 21, 24.1.2013, p. 44 and EEA Supplement No 6, 24.1.2013, p. 10), e.i.f. 1.11.2012.
- \textsuperscript{[269]} 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.
- \textsuperscript{[274]} Indent added by Decision No 159/2019 (OJ L 291, 10.11.2022, p. 39 and EEA Supplement No 74, 10.11.2022, p. 40), e.i.f. 15.6.2019.
- \textsuperscript{[275]} Indent added by Decision No 46/2023 (OJ L 203, 23.3.2023 and EEA Supplement No 77, 26.10.2023, p. 36), e.i.f. 18.3.2023.
- \textsuperscript{[276]} Indent added by Decision No 20/2024 (OJ L 204/1545, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 42), e.i.f. 3.2.2024.


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

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

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

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

(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.


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

(a) Unless otherwise specified, references to Union law shall be understood as referring to the EEA Agreement.

(b) Article 3 shall be amended as follows:

(i) in paragraph 24, the words “or the customs administrations of the EFTA States responsible for applying the customs legislation and any other authorities of the EFTA States empowered under
national law to apply certain customs legislation” shall be added after the reference to Regulation (EU) No 952/2013;

(ii) in paragraph 25, the words “or, as regards the EFTA States, the corresponding procedures in accordance with their respective national customs legislation” shall be added after the reference to Regulation (EU) No 952/2013;

(iii) in paragraph 26, the words “or within the customs territories of the EFTA States” shall be added after the words “customs territory of the Union”.

(c) In paragraph 2 of Article 14, the words “, including the principles of the Charter of Fundamental Rights of the European Union” shall not apply to the EFTA States.

(d) In paragraphs 3 and 4 of Article 25 and in the second subparagraph of Article 28(4), references to Regulation (EU) No 952/2013 of the European Parliament and of the Council shall, as regards the EFTA States, be understood to refer to corresponding provisions of national customs law.

(e) As regards Liechtenstein, obligations on authorities designated under Article 25(1) shall be governed by national law.

(f) Products exported from Liechtenstein to the other Contracting Parties may be subjected to controls in accordance with Articles 25 to 28 when entering the EEA.

(g) Articles 25(2), (4) and (6) and 34(6) shall not apply to Liechtenstein.

(h) Article 26(4) shall not apply to the EFTA States.

(i) In Article 28, as regards the EFTA States, the words “the customs data-processing system” shall read “any notification issued to the affected parties in accordance with national procedures”.

(j) The EFTA States shall participate fully, without the right to vote, in the Union Product Compliance Network in accordance with Articles 29-31. The EFTA Surveillance Authority shall participate as observer.

3va. [2021] R 1121: Commission Implementing Regulation (EU) 2021/1121 of 8 July 2021 specifying the details of the statistical data to be submitted by the Member States as regards controls on products entering the Union market with regard to product safety and compliance (OJ L 243, 9.7.2021, p. 37).

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

(a) In paragraphs 1(c) and 4 of Article 1, references to Commission Delegated Regulation (EU) 2015/2446 shall, as regards the EFTA States, be understood to refer to corresponding provisions of national customs law.

(b) In paragraph 1(c)(ix) of Article 1, the words “Union legislation” shall be replaced by the words “provisions of the EEA Agreement”.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

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


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

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


XX. FREE MOVEMENT OF GOODS – GENERAL

ACTS REFERRED TO[2006]


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

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

2. [2007] 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. [2105] [ ]


**XXI. CONSTRUCTION PRODUCTS**

**ACTS REFERRED TO**


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{2102} Point inserted by Decision No 49/2010 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 1.5.2010.


{2104} Indent and words “, as amended by:” added by Decision No 21/2015 (OJ L 93, 7.4.2016, p. 35 and EEA Supplement No 21, 7.4.2016, p. 30), e.i.f. 26.2.2015.

{2105} Indent added by Decision No 22/2015 (OJ L 93, 7.4.2016, p. 36 and EEA Supplement No 21, 7.4.2016, p. 31), e.i.f. 26.2.2015.


1n. [ ] [2120]


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1zr. [ ] [188]


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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 then the appropriate information is sent to the EEA Joint Committee with the approval of the President of EOTA and the Technical Board votes on the application. If the Technical Board votes in favour of the application then the President of EOTA issues an authority to issue ETAs. If the Technical Board votes against the application 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 EEA Joint Committee then the appropriate information is sent with the approval of the President of EOTA to the EEA Agreement on Common Technical Regulations and EEA Supplement No [to be published]’.
Board it will be passed to the Executive Commission for decision, as to whether it should be submitted to the EEA Joint Committee.’

(b) In the third paragraph of chapter 3.2.2. of the Annex the term 'Commission of the EC' shall be replaced by the term 'EEA Joint Committee'.

(c) In the last sentence of chapter 3.2.5. of the Annex the words 'shall be referred to the Standing Committee on Construction (the Directive, Article 9.2) via the Commission of the EC.' shall be replaced by 'shall be referred to the EEA Joint Committee.'


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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. [ ] {2218}


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-\[(227)\]

-\[(228)\]

-\[(229)\]

-\[(230)\]

-\[(231)\]

-\[(232)\]

-\[(233)\]

-\[(234)\]

-\[(235)\]

-\[(236)\]

**ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE[^2]**

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

2. **C/87/33/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**


[^232]: Indent added by Decision No 209/2021 (OJ L 2024/304, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 42), e.i.f. 10.7.2021.

[^233]: Indent added by Decision No 209/2021 (OJ L 2024/304, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 42), e.i.f. 10.7.2021.

[^234]: Indent added by Decision No 210/2021 (OJ L 2024/295, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 44), e.i.f. 10.7.2021.

[^235]: Indent added by Decision No 210/2021 (OJ L 2024/295, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 44), e.i.f. 10.7.2021.


ACTS REFERRED TO

1. [ ]\(^{2239}\)

1a. [ ]\(^{2240}\)

1b. [ ]\(^{2241}\)


\(^{2243}\) 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.


\(^{2246}\) 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.


\(^{2249}\) Indent added by Decision No 38/2020 (OJ L 57, 23.2.2023, p. 16 and EEA Supplement No 16, 23.2.2023, p. 16), e.i.f. 1.8.2020.

\(^{2250}\) Indent added by Decision No 70/2022 (OJ L 182, 7.7.2022, p. 50 and EEA Supplement No 45, 7.7.2022, p. 38), e.i.f. 19.3.2022.


ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

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


**XXV. TOBACCO**

**ACTS REFERRED TO**

1. \textit{C/2261} [\textit{2261}]

2. \textit{C/2262} [\textit{2262}]

3. \textit{C/2263} [\textit{2263}]

4. \textit{C/2264} [\textit{2264}]


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

(a) In the second subparagraph of Article 5(1), as regards the EFTA States, the words “by 20 November 2016” shall read “at the latest six months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

(b) The following subparagraph shall be added to Article 6(4) and Article 7(13):

“In cases concerning manufacturers and importers in the EFTA States, the EFTA Surveillance Authority shall collect any fees charged by the Commission.”.

(c) With regard to Norway, the following subparagraph shall be added to Article 12(1):

“Taking into account the specific national circumstances supported by statistics regarding the health risks related to the use and use patterns of tobacco for oral use, tobacco for oral use placed on the market in Norway may carry the following alternative health warning:

“This tobacco product increases the risk of harm to the foetus and stillbirth”.”


\textit{[2265]} Indent added by Decision No 239/2023 (OJ L 2024/993, 25.4.2024 and EEA Supplement No 35, 25.4.2024, p. 85), e.i.f. pending.
In Article 15(13), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

In Article 16(3), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

The prohibition in Article 17 shall not apply to the placing on the market in Norway of the product defined in Article 2, point (8). Norway shall ban export of the product defined in Article 2, point (8), to all Contracting Parties to this Agreement, with the exception of Sweden.

In Article 30, as regards the EFTA States, the words “20 May 2017” shall read “one year after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

In Article 30, points (a) and (c), as regards the EFTA States, the words “20 May 2016” shall read “the date of entry into force Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.

In Article 30, point (b), as regards the EFTA States, the words “20 November 2016” shall read “six months after the date of entry into force of Decision of the EEA Joint Committee No 6/2022 of 4 February 2022”.


3f. 32016 D 0786: Commission Implementing Decision (EU) 2016/786 of 18 May 2016 laying down the procedure for the establishment and operation of an independent advisory panel assisting Member States and the Commission in determining whether tobacco products have a characterising flavour (OJ L 131, 20.5.2016, p. 79).

3g. 32016 D 0787: Commission Implementing Decision (EU) 2016/787 of 18 May 2016 laying down a priority list of additives contained in cigarettes and roll-your-own tobacco subject to enhanced reporting obligations (OJ L 131, 20.5.2016, p. 88).

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

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3a Inserted by Decision No 90/2016 (OJ L 300, 16.11.2017, p. 39 and EEA Supplement No 73, 16.11.2017, p. 43), e.i.f. pending.

3b Inserted by Decision No 90/2016 (OJ L 300, 16.11.2017, p. 39 and EEA Supplement No 73, 16.11.2017, p. 43), e.i.f. pending.

3c Inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

3d Inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

3e Inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

3f Inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.

3g Inserted by Decision No 7/2022 (OJ L 175, 30.6.2022, p. 15 and EEA Supplement No 42, 30.6.2022, p. 13), e.i.f. pending.
In Article 2, as regards the EFTA States, the words “1 January 2017” shall read “the date of entry into force of Decision of the EEA Joint Committee No 7/2022 of 4 February 2022”.


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

(a) In Article 7(6)(a), as regards the EFTA States, the words “20 May 2020” shall read “sixteen months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(b) In Article 7(6)(b), as regards the EFTA States, the words “20 May 2021” shall read “two years after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(c) In Article 37(1), as regards the EFTA States, the words “20 May 2019” shall read “sixteen months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(d) In Article 37(2), as regards the EFTA States, the words “20 May 2024” shall read “five years after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

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

(a) In Article 3(4), as regards the EFTA States, the words “20 September 2018” shall read “eight months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(b) In Article 4(3), as regards the EFTA States, the words “20 September 2018” shall read “eight months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.


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

(a) In Article 3(4), as regards the EFTA States, the words “20 September 2018” shall read “eight months after the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(b) Indent and words “as amended by:” added by Decision No 21/2024 (OJ L 2024/1548, 27.6.2024 and EEA Supplement No 51, 27.6.2024, p. 43), e.i.f. 3.2.2024.
(c) In Article 9(1), the words “20 May 2019” shall, as regards the EFTA States, read “sixteen months after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”,

In Article 9(1), the words “20 May 2020” shall, as regards the EFTA States, read “two years and four months after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.

(d) In Article 9(2), the words “20 May 2024” shall, as regards the EFTA States, read “five years after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”,

In Article 9(2), the words “20 May 2026” shall, as regards the EFTA States, read “seven years after the date of the entry into force of Decision of the EEA Joint Committee No 9/2022 of 4 February 2022”.


XXVI. ENERGY

ACTS REFERRED TO

1.  \[^{2280}\] \[^{2281}\]

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.

\[^{2281}\] 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.  \[^{2282}\] \[^{2283}\]
2. \[ \] \[(228)\]

3. \[ \] \[(228)\]


9.\[(228)\] 32019 R 0787: Regulation (EU) 2019/787 of the European Parliament and of the Council of 17 April 2019 on the definition, description, presentation and labelling of spirit drinks, the use of the names of spirit drinks in the presentation and labelling of other foodstuffs, the protection of geographical indications for spirit drinks, the


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

(a) The provisions of the Regulation shall not prejudice the right of the EFTA States to prohibit, on a non-discriminatory basis, the placing on their national market of spirit drinks for direct human consumption which exceed an alcoholic strength of 60%.

(b) The EFTA States shall be invited to send observers to the meetings of the Committee for Spirit Drinks, as referred to in Article 47, dealing with matters which fall within acts referred to in the Agreement. The representatives of the EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.

(c) Paragraph 4(d) of Protocol 1 to the Agreement shall not apply to Chapter III of the Regulation.

9a. [ ] \[229\]


9af. (32020 R 0179) Commission Implementing Regulation (EU) 2020/179 of 3 February 2020 approving amendments to the specification for a spirit drink whose name is registered as a geographical indication (Berliner Kümme1) (OJ L 37, 10.2.2020, p. 4).


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

In Article 3(1), as regards the EFTA States, the words “25 August 2021” shall read “three months after the date of entry into force of the Decision of the EEA Joint Committee No 241/2022 of 23 September 2022”.

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Note: The numbers (32020 R) refer to the page numbers in the original document. The specific regulations and their dates are referenced throughout the text.


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

The following shall be added in Article 13:

“Notwithstanding the provisions of Protocol 1 to the Agreement, the communications made by the competent authorities of the EFTA States to the Commission pursuant to Article 13(1) shall follow the procedure set out in point (b). Point 4 of Protocol 1 shall not apply to Article 13.”


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The EFTA States concerned shall be invited to send observers to the meetings of the Committee on aromatised wine products and the listing of the existing geographical designations in that register (OJ L 42, 14.2.2020, p. 8).

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.

The establishment of the register of geographical indications protected in the sector of aromatised wine products shall not have the right to vote.

The EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.

The EFTA States shall participate fully in the work of the Committee, but shall not have the right to vote.

XXVIII. CULTURAL GOODS

ACTS REFERRED TO


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

ACTS REFERRED TO


{2346} ACTS OF WHICH THE CONTRACTING PARTIES SHALL TAKE NOTE

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


XXX. MEDICAL DEVICES {2347}

ACTS REFERRED TO

1. {2348}


\(^{2351}\) 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.


\[-[\(^{2353}\)] [ ]


5. [ ] [\(^{2359}\)]


\(^{2360}\) 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.

7. [ ]


9. [ ]


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

(a) The EFTA States shall participate fully in the Medical Device Coordination Group (‘MDCG’) established under Article 103, but shall not have the right to vote.

(b) The EFTA States shall participate in the European database on medical devices (Eudamed) set up by the Commission as referred to in Article 33.


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

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


\[\text{indent added by Decision No } 242/2022 (OJ L 246, 22.9.2022, p. 82), \text{c.i.f. } 30.4.2022.\]


\[\text{indent inserted by Decision No } 48/2023 (OJ L 233, 26.10.2023), \text{c.i.f. } 18.3.2023.\]


\[\text{indent and words "", as amended by:": added by Decision No } 276/2023 (OJ L 244/2024, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 51), \text{c.i.f. } 28.10.2023.\]

\[\text{indent inserted by Decision No } 48/2023 (OJ L 233, 26.10.2023 and EEA Supplement No 77, 26.10.2023, p. 38), \text{c.i.f. } 18.3.2023.\]

\[\text{indent inserted by Decision No } 301/2019 (OJ L 68, 5.3.2020, p. 44 and EEA Supplement No 14, 5.3.2020, p. 50), \text{c.i.f. } 12.6.2020.\]

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

\[\text{indent and words "", as amended by:": added by Decision No } 177/2022 (OJ L 267, 13.10.2022, p. 21 and EEA Supplement No 66, 13.10.2022, p. 19), \text{c.i.f. } 11.6.2022.\]

\[\text{indent added by Decision No } 153/2023 (OJ L 250, 30.11.2023 and EEA Supplement No 87, 30.11.2023, p. 16), \text{c.i.f. } \text{pending.}\]

\[\text{indent added by Decision No } 275/2023 (OJ L 212, 16.5.2024 and EEA Supplement No 42, 16.5.2024, p. 50), \text{c.i.f. } \text{pending.}\]


Modalities for the association of the EFTA States in accordance with Article 101 of this Agreement:

The EFTA States shall be fully associated with the work of the Executive Steering Group on Shortages of Medical Devices and shall have the same rights and obligations within it as the EU Member States, except for the right to vote.

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

In Article 34(2), the words “or Article 53 of the EEA Agreement” shall be inserted after the words “Article 101 TFUE”.

\textit{Acts of which the Contracting Parties shall take note}\textsuperscript{(290)}


\textsuperscript{(285)} Point inserted by Decision No 122/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 13.6.2024.


\textsuperscript{(288)} Point inserted by Decision No 211/2021 (OJ L, 2024/284, 8.2.2024 and EEA Supplement No 13, 8.2.2024, p. 46), e.i.f. 10.7.2021.

\textsuperscript{(289)} Point and adaptation text inserted by Decision No 167/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. pending.

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


XXXI. RECREATIONAL CRAFT{[2395]}

ACTS REFERRED TO

1. [ ] {[2389]}


XXXII. MARINE EQUIPMENT{[2396]}

ACTS REFERRED TO

1. [ ] {[2399]}


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[2403] Point inserted by Decision No 162/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 6.7.2024.
APPENDIX 1\textsuperscript{(2404)}

ENERGY LABELS

SECTION 1\textsuperscript{(2405)} [ ]


SECTION 2\footnote{2406} [ ]

SECTION 3[2407] [ ]

SECTION 4\(^{(2488)}\)

Commission Directive 96/60/EC

(household combined washer-driers)

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### Energi

#### Merke

<table>
<thead>
<tr>
<th>Modell</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
<th>F</th>
<th>G</th>
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#### Lavt forbruk

<table>
<thead>
<tr>
<th>Energiforbruk (Ved da G vaskning og tørring med full kapasitet) kWh</th>
<th>X.YZ</th>
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</thead>
<tbody>
<tr>
<td>Vask &amp; centrifugering kWh</td>
<td>X.YZ</td>
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#### Høyt forbruk

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<tr>
<th>Ytelsevne</th>
<th>A: høy</th>
<th>B: middels</th>
<th>C: lav</th>
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<tbody>
<tr>
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<td>y.z</td>
<td>y.z</td>
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<tr>
<td>Tørring</td>
<td>y.z</td>
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#### Vannforbruk (totalt) l

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<th>Lydnivå</th>
<th>dB(A) (Støy)</th>
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<tbody>
<tr>
<td>Vaskning</td>
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<tr>
<td>Sentrifugering</td>
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<tr>
<td>Tørring</td>
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SECTION 5[\ref{2409}] [ ]

<table>
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<th><strong>Energi</strong></th>
<th><strong>Oppvaskmaskin</strong></th>
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<tr>
<td><strong>Merke</strong></td>
<td>Logo</td>
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<td><strong>Modell</strong></td>
<td>ABC 123</td>
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<td>B</td>
</tr>
<tr>
<td><strong>Høyt forbruk</strong></td>
<td>X.YZ</td>
</tr>
</tbody>
</table>

**Energiforbruk kWh/oppvask**
(på grunnlag av testresultater for normal/program ved kraftvernstilsnitt)

Den faktiske energibruken avhenger av hvordan maskinen brukes.

**Rengjøringsøvne**
A: høy  G: lav

**Tørkeøvne**
A: høy  G: lav

**Standardkuverter**

**Vannforbruk l/oppvask**

**Lydnivå**
DB(A) dB, (Støy)

Produktbrosjyrene inneholder ytterligere opplysninger

**Europeisk standard EN 50542**

**Direktiv 97/17/EF om energimerkning av oppvaskmaskiner**
SECTION 6{2410}

Commission Directive 2002/40/EC

(household electric ovens)

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<th>Stór</th>
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<table>
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<th>Notkunarrými (lítrar)</th>
<th>XYZ</th>
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<tbody>
<tr>
<td>Orkunotkun (kWh)</td>
<td></td>
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</table>

Hitun: Hefðbundinn Blástursofn

(mðað við staðalálag)

Nánari upplýsingar er að finna í bæklingum sem fylgja vörunum

Norm EN 50304
Rafmagnsþekkingar
Tilskipun 2002/40/EB um orkumerkingar
<table>
<thead>
<tr>
<th>Orka</th>
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### Góð nýtni

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<th>D</th>
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### Slæm nýtni

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<tr>
<td>Blástursofn</td>
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### Þróð (míðað vóð staðalálag)

<table>
<thead>
<tr>
<th>Notkunarrými (litrar)</th>
<th>XYZ</th>
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</table>

### Stærð:

<table>
<thead>
<tr>
<th>Lítill</th>
<th>Medal</th>
<th>Stór</th>
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### Hávaði (dB(A) re 1 pW)

Nánari upplýsingar er að finna í bæklingum sem fylgja vörunum

**Norm EN 50304**

Rafmagnsbökunarofnar

**Tilskipun 2002/40/EB um orkumerkingar**
**Energi**

**Elektrisk stekeøvn**

<table>
<thead>
<tr>
<th>Merke</th>
<th>Logo</th>
<th>ABC 1 2 3</th>
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</table>

**Høyt energiforbruk**

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

**Lavt energiforbruk**

- **X.YZ**
- **X.YZ**

**Energiforbruk (kWh)**

- Oppvarmingsfunksjon:
  - Tradisjonell oppvarming
  - Varmluft

**(basert på standardbelastning)**

| Nettovolum (liter) | XYZ |

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<tr>
<td>Liten</td>
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<tr>
<td>Middels stor</td>
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<tr>
<td>Stor</td>
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| Lydnivå (støy) (dB(A) re 1 pW) |

Produktbrosjyrene inneholder ytterligere opplysninger

Standard EN 50304
Elektriske stekeovner
Direktiv 2002/40/EF om energimerking
**Energi**

**Merk**

**Modell**

**Høy energiforbruk**

A

B

C

D

E

F

G

**Lav energiforbruk**

Energiforbruk (kWh)

Oppvarmingsfunksjon:

- Tradisjonell oppvarming
- Varmluft (basert på standardbelastning)

Nettovolum (liter)

Type:

- Liten
- Middels stor
- Stor

Lydnivå (støy) (dB(A) re 1 pW)

Produktbrosjyrene inneholder ytterligere opplysninger

Standard EN 50304
Elektriske stekeovner
Direktiv 2002/40/EF om energimerking
SECTION 7{2411} [ ]

APPENDIX 2\(^{[2412]}\)

ENERGY TABLES

SECTION I\(^{[2413]}\) [ ]


SECTION 2[\textsuperscript{2414}] []

\textsuperscript{2414} Section 2 (Commission Directive 95/12/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 tables correspond to the tables previously contained in former adaptation b) to the second indent of point 4 of Chapter IV, subsequently reinserted as point 4b 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 3{2415} []

**SECTION 4**

**Commission Directive 96/60/EC**

*(household combined washer-driers)*

<table>
<thead>
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<th>IS</th>
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<tbody>
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<td>Mail order</td>
<td></td>
</tr>
<tr>
<td>Annex I</td>
<td>Annex II</td>
<td>Annex III</td>
<td></td>
</tr>
</tbody>
</table>

### 1. **Energy**

- **EN**: Energi
- **NO**: Orka
- **IS**: Þvottavél – þurrkari

### 2. **Washer-drier**

- **EN**: Kombinert vaske- og tørkemaskin
- **NO**: Framleiðandi
- **IS**: Fræðiþeimönði

### 3. **Manufacturer**

- **EN**: Merke
- **NO**: Gerð
- **IS**: Gerð

### 4. **Model**

- **EN**: Modell
- **NO**: Gerð
- **IS**: Gerð

### 5. **Energy**

#### More efficient
- **EN**: Lavt forbruk
- **NO**: Góð nýtni
- **IS**: Látt samþykkja

#### Less efficient
- **EN**: Høyt forbruk
- **NO**: Slæm nýntni
- **IS**: Þakk taksin

### 6. **Energy efficiency class**

- **EN**: Relativ energibruk
- **NO**: Orkunýtniflokkur
- **IS**: Ólgerð

### 7. **Energy consumption**

- **EN**: Forbruk
- **NO**: Energiforbruk
- **IS**: Orkunotkun

<table>
<thead>
<tr>
<th>Energy consumption for washing, spinning and drying</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>EN</strong>: Energibruk til vasking, sentrifugering og tørking</td>
</tr>
<tr>
<td><strong>NO</strong>: Orkunotkun til ðvott, þeytivindingu og þurrkun</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Energy consumption for washing and spinning only</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>EN</strong>: Energibruk pr vask og sentrifugering alene</td>
</tr>
<tr>
<td><strong>NO</strong>: Orkunotkun við þvott og þeytivindigu eingöngu</td>
</tr>
</tbody>
</table>

### 8. **Washing performance**

- **EN**: Vaskeevne
- **NO**: Þvottahæfni
- **IS**: Þorrkari

### 9. **Washing performance class**

| Washing performance class | **EN**: Vaskeevne
|---------------------------|----------------|
| A (higher)                | **NO**: Þvottahæfni
| G (lower)                 | **IS**: Þorrkari

### 10. **Energy consumption**

<table>
<thead>
<tr>
<th>Energy consumption for washing and spinning only</th>
</tr>
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<tr>
<td><strong>EN</strong>: Energibruk pr vask og sentrifugering alene</td>
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<tr>
<td><strong>NO</strong>: Orkunotkun við þvott og þeytivindigu eingöngu</td>
</tr>
</tbody>
</table>

### 11. **Actual consumption**

- **EN**: Den faktiske energibruk
- **NO**: Raunnotkun
- **IS**: Stykkur

- **EN**: Braking capacity used
- **NO**: Ósamtaksanunnin
- **IS**: Ósamtaksanunnin

### 12. **Energy consumption for washing and spinning only**

<table>
<thead>
<tr>
<th>Energy consumption for washing and spinning only</th>
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</thead>
<tbody>
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<td><strong>EN</strong>: Energibruk pr vask og sentrifugering alene</td>
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<td><strong>NO</strong>: Orkunotkun við þvott og þeytivindigu eingöngu</td>
</tr>
</tbody>
</table>

### 13. **Washing performance**

- **EN**: Vaskeevne
- **NO**: Þvottahæfni
- **IS**: Þorrkari

### 14. **Washing performance class**

| Washing performance class | **EN**: Vaskeevne
|---------------------------|----------------|
| A (higher)                | **NO**: Þvottahæfni
| G (lower)                 | **IS**: Þorrkari

### Footnote

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<td>Annex II</td>
<td>Annex II</td>
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<td>8</td>
<td>5</td>
<td>Water remaining after spin ...% (as a proportion of dry weight of wash)</td>
<td>Restvanninnhold etter sentrifugering ....% (i forhold til vekten av tørt tøy)</td>
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<td>VIII</td>
<td>9</td>
<td>6</td>
<td>Spin speed (rpm)</td>
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<td>IX/X</td>
<td>10/11</td>
<td>7/8</td>
<td>Capacity (cotton) kg</td>
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<td>X</td>
<td>10</td>
<td>7</td>
<td>Washing</td>
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<td>IX</td>
<td>11</td>
<td>8</td>
<td>Drying</td>
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<tr>
<td>XI</td>
<td>Water consumption (total)</td>
<td>Vannforbruk (totalt)</td>
<td>Vatnsnotkun (alls)</td>
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<td>9</td>
<td>Water consumption washing, spinning and drying</td>
<td>Vannforbruk vasking, sentrifugering og tøring</td>
</tr>
<tr>
<td>13</td>
<td>10</td>
<td>Water consumption for washing and spinning only</td>
<td>Vannforbruk til vask-og sentrifugering alene</td>
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<tr>
<td>14</td>
<td>Washing and drying time</td>
<td>Vaske- og tørketid</td>
<td>Þvotta- og þurkftími</td>
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<td>16</td>
<td>11</td>
<td>Estimated annual consumption for a 4-person household, always using the drier (200 cycles)</td>
<td>Anslått årlig forbruk for en husstand på fire personer som alltid tørker tøyet i maskinen (200 ganger)</td>
</tr>
<tr>
<td>17</td>
<td>12</td>
<td>Estimated annual consumption for a 4-person household, never using the drier (200 cycles)</td>
<td>Anslått årlig forbruk for en husstand på fire personer som aldri tørker tøyet i maskinen (200 ganger)</td>
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<td>XII</td>
<td>18</td>
<td>13</td>
<td>Noise (dB(A) re 1 pW)</td>
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<td>Washing</td>
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<td>13</td>
<td>Spinning</td>
<td>Sentrifugering</td>
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<tr>
<td>18</td>
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<td>Drying</td>
<td>Törking</td>
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<td>Europeisk standard EN 50229</td>
<td>Staðall EN 50229</td>
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<td>Washer-drier Label Directive 96/60/EC</td>
<td>Direktiv 96/60/EF om energimerking av kombinerte vaske- og tørkemaskiner</td>
<td>Tilskipun 96/60/EB um merkingar þvottavéla-þurrkara</td>
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SECTION 5[^247] [ ]

### SECTION 6\(^{24/18}\)

*Commission Directive 2002/40/EC*

*(household electric ovens)*

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<td>Electric oven</td>
<td>Rafmagnsbökunarofn</td>
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<td>Framleiðandi</td>
<td>Merke</td>
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<td>Góð nýtni</td>
<td>Læt energiforbruk</td>
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<td>Less efficient</td>
<td>Slæm nýtni</td>
<td>Háyt energiforbruk</td>
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<td>Orkunýtniflokkur …á kvarðanum A (góð nýtni) til G (slæm nýtni)</td>
<td>Klassifisering av energieeffektivitet etter en skala fra A (lavt energiforbruk) til G (høyt nergiforbruk)</td>
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<td>Stekeoverflate</td>
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\(^{24/18}\) Section, including heading and tables, added by Decision No 141/2003 (OJ L 41, 12.02.2004, p. 11 and EEA Supplement No 7, 12.02.2004, p. 9), e.i.f. 11.8.2003.
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<td>kWh</td>
<td>kWh</td>
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<td>Large</td>
<td>Stór</td>
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<td>6</td>
<td>Time to cook standard load</td>
<td>Bökunartími við staðalálag</td>
<td>Koketid ved standardbelastning</td>
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<td>Noise (dB(A) re 1 pW)</td>
<td>Hávaði (dB(A) re 1 pW)</td>
<td>Lydnivå (støy) (dB(A) re 1 pW)</td>
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<td>Produktbrosjyrene inneholder ytterligere opplysninger</td>
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<td></td>
<td>11</td>
<td></td>
<td>The area of the largest baking sheet</td>
<td>Stærð stærstu bökunarplötu</td>
<td>Arealet til den største stekeplaten</td>
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SECTION 7\(^{[2419]}\) [ ]
HAZARD AND PRECAUTIONARY STATEMENTS IN ICELANDIC

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

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<tbody>
<tr>
<td>H200</td>
<td>Óstöðugt, sprengifím efní.</td>
</tr>
<tr>
<td>H201</td>
<td>Sprengifím efní, hætta á alsprengingu.</td>
</tr>
<tr>
<td>H202</td>
<td>Sprengifím efní, mikil hætta á sprengibroti.</td>
</tr>
<tr>
<td>H203</td>
<td>Sprengifím efní, hætta á bruna, höggbylgju eða sprengibrotum.</td>
</tr>
<tr>
<td>H204</td>
<td>Hætta á bruna eða sprengibrotum.</td>
</tr>
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<td>H205</td>
<td>Hætta á alsprengingu í bruna.</td>
</tr>
<tr>
<td>H220</td>
<td>Afar eldfím lofttegund.</td>
</tr>
<tr>
<td>H221</td>
<td>Eldfím lofttegund.</td>
</tr>
<tr>
<td>H222</td>
<td>Úðabrúsi með afar eldfínum efnínum.</td>
</tr>
<tr>
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<td>Úðabrúsi með eldfínum efnínum.</td>
</tr>
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<td>Afar eldfímur vöki og gufa.</td>
</tr>
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<td>Mjög eldfímur vöki og gufa.</td>
</tr>
<tr>
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<td>Eldfímur vöki og gufa.</td>
</tr>
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<td>Eldfím, fast efní.</td>
</tr>
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<td>Sprengifím við hitun.</td>
</tr>
<tr>
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<td>Eldfím eða sprengifím við hitun.</td>
</tr>
<tr>
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<td>Eldfím við hitun.</td>
</tr>
<tr>
<td>H250</td>
<td>Kvíknjar í sjálfkrafa við snertingu við loft.</td>
</tr>
<tr>
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<td>Sjálfhitandi, hætta á sjálfsíkviknun.</td>
</tr>
<tr>
<td>H252</td>
<td>Sjálfhitandi í miklu efnísmagni, hætta á sjálfsíkviknun.</td>
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<td>Í snertingu við vatn myndast eldfímar lofttegundir sem er hætt við sjálfsíkviknun.</td>
</tr>
<tr>
<td>H261</td>
<td>Eldfímar lofttegundir myndast við snertingu við vatn</td>
</tr>
<tr>
<td>H270</td>
<td>Getur valdið eða aukið bruna, eldmyndandi (oxandi).</td>
</tr>
<tr>
<td>H271</td>
<td>Getur valdið bruna eða sprengingu, mjög eldmyndandi (oxandi).</td>
</tr>
<tr>
<td>H272</td>
<td>Getur aukið bruna, eldmyndandi (oxandi).</td>
</tr>
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(2422) 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 and subsequently amended by Decision No 89/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.
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<td>Inniheldur kælda lofttegund, getur valdið kalsárum.</td>
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<td>Getur verið ætandi fyrir málma.</td>
</tr>
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<td>Banvænt við innöðun.</td>
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</tr>
<tr>
<td>H302</td>
<td>Hættulegt við innöðun.</td>
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<tr>
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Eitrað í snertingu við augu.

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Hættulegt ósonlaginu.

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### EUH 203
Innheldur sexgilt króm. Getur framkallað ofnæmisviðbrógð.

### EUH 204
Innheldur ísósýanöt. Getur framkallað ofnæmisviðbrógð.

### EUH 205
Innheldur epoxýefnisþætti. Getur framkallað ofnæmisviðbrógð.

### EUH 206
Varð! Notist ekki með öðrum vörum. Getur gefið frá sér hættulegar lofttegundir (klór).

### EUH 207

### EUH 208
Innheldur (heiti næmandi efnis). Getur framkallað ofnæmisviðbrógð.

### EUH 209/ 209A
Getur orðið mjög elfdint við notkun. Getur orðið elfdint við notkun.

### EUH 210
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### EUH 380
Getur valdið ínknirlatruflunum hjá mönnum

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Fylgð notkunarleiðbeiningum til að varast hættu fyrir heilbrigði manna og umhverfið.

### EUH 430
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### EUH 440
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### EUH 441
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### EUH 450
Getur valdið langvarandi og dreifðri mengum vatnsauðlinda

### EUH 451
Getur valdið mjög langvarandi og dreifðri mengun vatnsauðlinda

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The following shall be added to Part 2 of Annex IV of Regulation (EC) No 1272/2008:

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**APPENDIX 6{2423)**

HAZARD AND PRECAUTIONARY STATEMENTS IN NORWEGIAN

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

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{2423} 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 and subsequently amended by Decision No 89/2024 (OJ L [to be published] and EEA Supplement No [to be published]), e.i.f. 27.4.2024.
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<tr>
<td>P101</td>
<td>Dersom det er nødvendig med legehjelp, ha produktets beholder eller etikett for hånden.</td>
</tr>
<tr>
<td>P102</td>
<td>Oppbevares utilgjengelig for barn.</td>
</tr>
<tr>
<td>P103</td>
<td>Les etiketten før bruk.</td>
</tr>
<tr>
<td>P201</td>
<td>Innhent særskilt instruks før bruk.</td>
</tr>
<tr>
<td>P202</td>
<td>Skal ikke håndteres før alle advarsler er lest og oppfattet.</td>
</tr>
<tr>
<td>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/opbevares i nærheten av tøy /.../ brennbare materialer.</td>
</tr>
<tr>
<td>P221</td>
<td>Må ikke blandes med brennbare stoffer.</td>
</tr>
<tr>
<td>P222</td>
<td>Unngå kontakt med luft.</td>
</tr>
<tr>
<td>P223</td>
<td>Unngå all kontakt med vann, på grunn av fare for voldsom reaksjon og eksplosjonsaktig brann.</td>
</tr>
<tr>
<td>P230</td>
<td>Holdes fuktet med …</td>
</tr>
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<td>P231</td>
<td>Håndteres under inertgass.</td>
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<td>P232</td>
<td>Beskyttes mot fuktighet.</td>
</tr>
<tr>
<td>P233</td>
<td>Hold beholderen tett lukket.</td>
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<tr>
<td>P234</td>
<td>Oppbevares bare i originalbeholder.</td>
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<tr>
<td>P235</td>
<td>Oppbevares kjølig.</td>
</tr>
<tr>
<td>P240</td>
<td>Beholder og mottaksutstyr jordes/potensialutlignes.</td>
</tr>
<tr>
<td>P241</td>
<td>Bruk elektrisk materiell /ventilasjonsmateriell/belysningsmateriell som er eksplosjonssikkert.</td>
</tr>
<tr>
<td>P242</td>
<td>Bruk bare verktøy som ikke avgir gnister.</td>
</tr>
<tr>
<td>P243</td>
<td>Treff tiltak mot statisk elektrisitet.</td>
</tr>
<tr>
<td>P244</td>
<td>Reduksjonsventiler skal holdes fri for fett og olje.</td>
</tr>
<tr>
<td>P250</td>
<td>Må ikke utsettes for sliping/støt/…/friksjon.</td>
</tr>
<tr>
<td>P251</td>
<td>Beholder under trykk: Må ikke punkteres eller brennes, selv ikke etter bruk.</td>
</tr>
<tr>
<td>P260</td>
<td>Ikke innånd støv/røyk/gass/tåke/damp/aerosoler.</td>
</tr>
<tr>
<td>P261</td>
<td>Unngå innånding av støv/røyk/gass/tåke/damp/aerosoler.</td>
</tr>
<tr>
<td>P262</td>
<td>Må ikke komme i kontakt med øyne, huden eller klær.</td>
</tr>
<tr>
<td>P263</td>
<td>Unngå kontakt under graviditet/amming.</td>
</tr>
<tr>
<td>P264</td>
<td>Vask … grundig etter bruk.</td>
</tr>
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<tr>
<td>P270</td>
<td>Ikke spis, drikk eller røyk ved bruk av produktet.</td>
</tr>
<tr>
<td>P271</td>
<td>Brukes bare utendørs eller i et godt ventileret område.</td>
</tr>
<tr>
<td>P272</td>
<td>Tilsølte arbeidsklær må ikke fjernes fra arbeidsplassen.</td>
</tr>
<tr>
<td>P273</td>
<td>Unngå utslipp til miljøet.</td>
</tr>
<tr>
<td>P280</td>
<td>Benytt vernehansker /verneklær/vernebriller/ansiktsskjerm.</td>
</tr>
<tr>
<td>P281</td>
<td>Bruk påkrevd personlig verneutstyr.</td>
</tr>
<tr>
<td>P282</td>
<td>Bruk kuldeisolerende hansker /visir/øyevern.</td>
</tr>
<tr>
<td>P283</td>
<td>Benytt brannbestandige/flammehemmende klær.</td>
</tr>
<tr>
<td>P284</td>
<td>Bruk åndedrettsvern.</td>
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<td>P285</td>
<td>Ved utilstrekkelig ventilasjon skal åndedrettsvern benyttes.</td>
</tr>
<tr>
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<td>VED SVELGING:</td>
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<td>VED KONTAKT MED KŁÆR:</td>
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<tr>
<td>P310</td>
<td>Kontakt umiddelbart et GIFTINFORMASJONSSENTER eller lege.</td>
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<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>
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<td>P314</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>
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<td>P321</td>
<td>Særlig behandling (se … på etiketten).</td>
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<tr>
<td>P322</td>
<td>Særlige tiltak (se … på etiketten).</td>
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<td>P330</td>
<td>Skyll munnen.</td>
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<tr>
<td>P331</td>
<td>IKKE framkall brekning.</td>
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<td>P332</td>
<td>Ved hudirritasjon:</td>
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<td>P333</td>
<td>Ved hudirritasjon eller utslett:</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>
<tr>
<td>P338</td>
<td>Fjern eventuelle kontaktlinser dersom dette enkelt lar seg gjøre. Fortsett skyllingen.</td>
</tr>
<tr>
<td>P340</td>
<td>Flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P341</td>
<td>Ved pustevansker, flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P342</td>
<td>Ved symptomer i luftveiene:</td>
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<td>P350</td>
<td>Vask forsiktig med mye såpe og vann.</td>
</tr>
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<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>
</tr>
<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>
</tr>
<tr>
<td>P371</td>
<td>Ved større brann og store mengder:</td>
</tr>
<tr>
<td>P372</td>
<td>Eksplosjonsfare ved brann.</td>
</tr>
<tr>
<td>P373</td>
<td>IKKE bekjemp brannen når den når eksplosive varer.</td>
</tr>
<tr>
<td>P374</td>
<td>Bekjemp brannen med normal forsiktighet på behørig avstand.</td>
</tr>
<tr>
<td>P375</td>
<td>Bekjemp brannen på avstand på grunn av eksplosjonsfare.</td>
</tr>
<tr>
<td>P376</td>
<td>Stopp lekkasje dersom dette kan gjøres på en sikker måte.</td>
</tr>
<tr>
<td>P377</td>
<td>Brann ved gasslekkasje: Ikke slukk med mindre lekkasjen kan stanses på en sikker måte.</td>
</tr>
<tr>
<td>P378</td>
<td>Slukk med:.....</td>
</tr>
<tr>
<td>P380</td>
<td>Evakuier området.</td>
</tr>
<tr>
<td>P381</td>
<td>Fjern alle tennkilder dersom dette kan gjøres på en sikker måte.</td>
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<tr>
<td>P390</td>
<td>Absorber spill for å hindre materiell skade.</td>
</tr>
<tr>
<td>P391</td>
<td>Samle opp spill.</td>
</tr>
<tr>
<td>P301 + P310</td>
<td><strong>VED SVELGING:</strong> Kontakt umiddelbart et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P301 + P312</td>
<td><strong>VED SVELGING:</strong> Kontakt et GIFTINFORMASJONSSENTER eller lege ved ubehag.</td>
</tr>
<tr>
<td>P301 + P330 + P331</td>
<td><strong>VED SVELGING:</strong> Skyl Munnen. IKKE framkall brekning.</td>
</tr>
<tr>
<td>P302 + P334</td>
<td><strong>VED HUDKONTAKT:</strong> Skyll i kaldt vann / anvend våt kompress.</td>
</tr>
<tr>
<td>P302 + P350</td>
<td><strong>VED HUDKONTAKT:</strong> Vask forsiktig med mye såpe og vann.</td>
</tr>
<tr>
<td>P302 + P352</td>
<td><strong>VED HUDKONTAKT:</strong> Vask med mye såpe og vann.</td>
</tr>
<tr>
<td>P303 + P361 + P353</td>
<td><strong>VED HUDKONTAKT</strong> (eller håret): Tilsølte klær må fjernes straks. Skyll/dusj huden med vann.</td>
</tr>
<tr>
<td>P304 + P340</td>
<td><strong>VED INNÅNDING:</strong> Flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P304 + P341</td>
<td><strong>VED INNÅNDING:</strong> Ved pustevansker, flytt personen til frisk luft og sørg for at vedkommende hviler i en stilling som letter åndedrettet.</td>
</tr>
<tr>
<td>P305 + P351 + P338</td>
<td><strong>VED KONTAKT MED ØYNENE:</strong> Skyll forsiktig med vann i flere minutter. Fjern eventuelle kontaktlinser dersom dette enkelt lar seg gjøre. Fortsett skyllingen.</td>
</tr>
<tr>
<td>P306 + P360</td>
<td><strong>VED KONTAKT MED KLÆR:</strong> Skyll umiddelbart tilsølte klær og hud med mye vann før klærne fjernes.</td>
</tr>
<tr>
<td>P308 + P313</td>
<td>Ved eksponering eller mistanke om eksponering: Søk legehjelp.</td>
</tr>
<tr>
<td>P309 + P311</td>
<td>Ved eksponering eller ubehag: Kontakt et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P332 + P313</td>
<td>Ved hudirritasjon: Søk legehjelp.</td>
</tr>
<tr>
<td>P335 + P334</td>
<td>Børst bort løse partikler fra huden. Skyll i kaldt vann / anvend våt kompress.</td>
</tr>
<tr>
<td>No.</td>
<td>Norwegian</td>
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<tr>
<td>P337 + P313</td>
<td>Ved vedvarende øyeirritasjon: Søk legehjelp.</td>
</tr>
<tr>
<td>P342 + P311</td>
<td>Ved symptomer i luftveiene: Kontakt et GIFTINFORMASJONSSENTER eller lege.</td>
</tr>
<tr>
<td>P370 + P376</td>
<td>Ved brann: Stopp lekkasje dersom dette kan gjøres på en sikker måte.</td>
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<tr>
<td>P370 + P378</td>
<td>Ved brann: Sluk med …</td>
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<td>P401</td>
<td>Oppbevares …</td>
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<tr>
<td>P402</td>
<td>Oppbevares tørt.</td>
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<tr>
<td>P403</td>
<td>Oppbevares på et godt ventilert sted.</td>
</tr>
<tr>
<td>P404</td>
<td>Oppbevares i lukket beholder.</td>
</tr>
<tr>
<td>P405</td>
<td>Oppbevares innelåst.</td>
</tr>
<tr>
<td>P406</td>
<td>Oppbevares i korrosjonsbestandig/… beholder med korrosjonsbestandig indre belegg.</td>
</tr>
<tr>
<td>P407</td>
<td>Se til at det er luft mellom stabler/paller.</td>
</tr>
<tr>
<td>P410</td>
<td>Beskyttes mot sollys.</td>
</tr>
<tr>
<td>P411</td>
<td>Oppbevares ved en temperatur som ikke er høyere enn …°C /… °F.</td>
</tr>
<tr>
<td>P412</td>
<td>Må ikke utsettes for temperaturer høyere enn 50 °C /122 °F.</td>
</tr>
<tr>
<td>P413</td>
<td>Bulkmengder på over …kg/…lbs oppbevares ved en temperatur som ikke er høyere enn …°C /… °F.</td>
</tr>
<tr>
<td>P420</td>
<td>Må oppbevares adskilt fra andre materialer.</td>
</tr>
<tr>
<td>P422</td>
<td>Oppbevar innholdet under …</td>
</tr>
<tr>
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<td>Oppbevares tørt. Oppbevares i lukket beholder.</td>
</tr>
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<td>Oppbevares på et godt ventilert sted. Hold beholderen tett lukket.</td>
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<td>Oppbevares på et godt ventilert sted. Oppbevares kjølig.</td>
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<td>P410 + P403</td>
<td>Beskyttes mot sollys. Oppbevares på et godt ventilert sted.</td>
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<tr>
<td>P410 +</td>
<td>Beskyttes mot sollys. Må ikke utsettes for temperaturer høyere enn 50 °C /122 °F.</td>
</tr>
<tr>
<td>P412</td>
<td></td>
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<tr>
<td>P411 +</td>
<td>Oppbevares ved en temperatur som ikke er høyere enn ...°C /... °F. Oppbevares kjølig.</td>
</tr>
<tr>
<td>P235</td>
<td></td>
</tr>
<tr>
<td>P501</td>
<td>Innhold/beholder leveres til …</td>
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</tbody>
</table>