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Legislation

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Contents

EN

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

The titles of all other acts are printed in bold type and preceded by an asterisk.

DECISIONS

[1] Text with EEA relevance

II

(Non-legislative acts)

INTERNATIONAL AGREEMENTS

COUNCIL DECISION (EU) 2016/1995

of 11 November 2016

on the signing, on behalf of the European Union, of the Agreement in the form of an Exchange of Letters between the European Union and the Federative Republic of Brazil pursuant to Article XXIV:6 and Article XXVIII of the General Agreement on Tariffs and Trade (GATT) 1994 relating to the modification of concessions in the schedule of the Republic of Croatia in the course of its accession to the European Union

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on the Functioning of the European Union, and in particular the first subparagraph of Article 207(4), in conjunction with Article 218(5) thereof,

Having regard to the proposal from the European Commission,

Whereas:

(1) On 15 July 2013 the Council authorised the Commission to open negotiations with certain other Members of the World Trade Organization under Article XXIV:6 of the General Agreement on Tariffs and Trade (GATT) 1994, in the course of the accession to the Union of the Republic of Croatia.

(2) Negotiations were conducted by the Commission in accordance with the negotiating directives adopted by the Council.

(3) Those negotiations have been concluded and an Agreement in the form of an Exchange of Letters between the European Union and the Federative Republic of Brazil pursuant to Article XXIV:6 and Article XXVIII of the GATT 1994 relating to the modification of concessions in the schedule of the Republic of Croatia in the course of its accession to the European Union (the 'Agreement') was initialled on 12 July 2016.

(4) The Agreement should be signed,

HAS ADOPTED THIS DECISION:

Article 1

The signing on behalf of the Union of the Agreement in the form of an Exchange of Letters between the European Union and the Federative Republic of Brazil pursuant to Article XXIV:6 and Article XXVIII of GATT 1994 relating to the modification of concessions in the schedule of the Republic of Croatia in the course of its accession to the European Union is hereby authorised, subject to the conclusion of that Agreement ([1]).

Article 2

The President of the Council is hereby authorised to designate the person(s) empowered to sign the Agreement on behalf of the Union.

([1]) The text of the Agreement will be published together with the decision on its conclusion.

Article 3

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

Done at Brussels, 11 November 2016.

For the Council
The President
P. ŽIGA

REGULATIONS

COUNCIL IMPLEMENTING REGULATION (EU) 2016/1996

of 15 November 2016

implementing Regulation (EU) No 36/2012 concerning restrictive measures in view of the situation in Syria

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Regulation (EU) No 36/2012 of 18 January 2012 concerning restrictive measures in view of the situation in Syria and repealing Regulation (EU) No 442/2011 [1], and in particular Article 32(1) thereof,

Having regard to the proposal from the High Representative of the Union for Foreign Affairs and Security Policy,

Whereas:

(1) On 18 January 2012, the Council adopted Regulation (EU) No 36/2012, concerning restrictive measures in view of the situation in Syria.

(2) Two entities should be removed from the list of entities subject to restrictive measures, as set out in Section B of Annex II to Regulation (EU) No 36/2012.

(3) Annex II to Regulation (EU) No 36/2012 should therefore be amended accordingly,

HAS ADOPTED THIS REGULATION:

Article 1

Annex II to Regulation (EU) No 36/2012 is amended as set out in the Annex to this Regulation.

Article 2

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

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

Done at Brussels, 15 November 2016.

For the Council
The President
I. KORČOK

[1] OJ L 16, 19.1.2012, p. 1.

ANNEX

I. The following entities and the related entries are deleted from the list set out in Section B of Annex II to Regulation (EU) No 36/2012:

55. Tri-Ocean Trading

55a. Tri-Ocean Energy

COMMISSION IMPLEMENTING REGULATION (EU) 2016/1997

of 15 November 2016

amending Implementing Regulation (EU) No 808/2014 as regards the amendment of rural development programmes and monitoring of actions to support integration of third-country nationals, and correcting that Regulation

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) No 1305/2013 of the European Parliament and of the Council of 17 December 2013 on support for rural development by the European Agricultural Fund for Rural Development (EAFRD) and repealing Council Regulation (EC) No 1698/2005 ([1]), and in particular Article 12, Article 66(5), Article 67 and Article 75(5) thereof,

Whereas:

(1) Article 4(2) of Commission Implementing Regulation (EU) No 808/2014 ([2]) sets the maximum number of amendments to rural development programmes that Member States may submit to the Commission. Experience has shown that the maximum number of programme amendments should be increased to enable Member States to submit a limited number of additional amendments over the programming period. The cases in which the maximum number of programme amendments does not apply should be clarified and should include the amendments related to the adoption of certain emergency measures or to the new delimitation of areas facing significant natural constraints referred to in Article 32(5)(b) of Regulation (EU) No 1305/2013.

(2) The success of rural development programmes depends not only on good governance and their full implementation, but also on the readiness to adapt to new challenges and changing circumstances such as the migration crisis. In order to ensure good coordination of all existing intervention mechanisms, EAFRD support to actions targeted to integration of third-country nationals should be monitored at Union level.

(3) In point 1 of Part 2 of Annex III to Implementing Regulation (EU) No 808/2014, the Leader logo has erroneously not been inserted. This should be corrected. In point 1 of Annex IV, the reference to less favoured areas should be corrected. It should be replaced by a reference to areas facing natural or other specific constraints.

(4) Implementing Regulation (EU) No 808/2014 should therefore be amended accordingly.

(5) The measures provided for in this Regulation are in accordance with the opinion of the Rural Development Committee,

HAS ADOPTED THIS REGULATION:

Article 1

Implementing Regulation (EU) No 808/2014 is amended as follows:

(1) Article 4(2) is replaced by the following:

‘2. Programme amendments of the type referred to in Article 11(a)(i) of Regulation (EU) No 1305/2013 may be proposed no more than three times during the duration of the programming period.

([1]) OJ L 347, 20.12.2013, p. 487.
([2]) Commission Implementing Regulation (EU) No 808/2014 of 17 July 2014 laying down rules for the application of Regulation (EU) No 1305/2013 of the European Parliament and of the Council on support for rural development by the European Agricultural Fund for Rural Development (EAFRD) (OJ L 227, 31.7.2014, p. 18).

For all other types of amendments combined:

(a) a single amendment proposal may be submitted per calendar year and per programme, with the exception of the year 2023 in which year more than a single amendment proposal may be submitted for amendments concerning exclusively the adaptation of the financing plan, including any resulting changes to the indicator plan;

(b) three additional amendment proposals per programme may be submitted during the duration of the programming period.

The maximum number of amendments referred to in the first and second subparagraphs shall not apply:

(a) in case emergency measures due to natural disasters, catastrophic events or adverse climatic events formally recognized by the competent national public authority, or due to a significant and sudden change in the socio-economic conditions of the Member State or region, including significant and sudden demographic changes resulting from migration or reception of refugees, need to be taken;

(b) in case an amendment is necessary following a change to the Union legal framework;

(c) following the performance review referred to in Article 21 of Regulation (EU) No 1303/2013;

(d) in case of a change in the EAFRD contribution planned for each year referred to in Article 8(1)(h)(i) of Regulation (EU) No 1305/2013 resulting from developments relating to the annual breakdown by Member State referred to in Article 58(7) of that Regulation; the proposed amendments may include consequential changes in the description of measures;

(e) in case of changes related to the introduction of financial instruments referred to in Article 37 of Regulation (EU) No 1303/2013; or

(f) in case of changes related to the introduction of the new delimitation referred to in Article 32(5)(b) of Regulation (EU) No 1305/2013.';

(2) Article 5(4) is replaced by the following:

'4. Except in cases of emergency measures due to natural disasters, catastrophic events or adverse climatic events formally recognized by the competent national public authority, or due to a significant and sudden change in the socioeconomic conditions of the Member State or region, including significant and sudden demographic changes resulting from migration or reception of refugees, changes to the legal framework, or changes resulting from the performance review referred to in Article 21 of Regulation (EU) No 1303/2013, requests for amendment of the national framework referred to in paragraph 2 may be submitted only once per calendar year before 1 April. By way of derogation from the second subparagraph of Article 4(2), changes in programmes that result from such revision may be done in addition to the amendment proposals submitted in accordance with that subparagraph.';

(3) Article 14(4) is replaced by the following:

'4. For types of operations where a potential contribution to focus areas referred to in Article 5, first paragraph, point (2)(a), Article 5, first paragraph, points (5)(a) to (d), and Article 5, first paragraph, point(6)(a) of Regulation (EU) No 1305/2013 is identified, or for types of operations where a potential contribution to the integration of third-country nationals is identified, the electronic record of the operations referred to in Article 70 of Regulation (EU) No 1305/2013 shall include flag(s) to identify those cases where the operation has a component contributing to one or more of those focus areas or goal.';

(4) Annexes III, IV and VII are amended as set out in the Annex to this Regulation.

Article 2

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

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

Done at Brussels, 15 November 2016.

For the Commission
The President
Jean-Claude JUNCKER

ANNEX

Annexes III, IV and VII to Implementing Regulation (EU) No 808/2014 are amended as follows:

(1) in point 1 of Part 2 of Annex III, point (b) is replaced by the following:

'(b) for the actions and measures financed by LEADER, the LEADER logo:

';

(2) in point 1 of Annex IV, indicator C32 is replaced by the following:

'C32. Areas facing natural or other specific constraints';

(3) in point 1(b) of Annex VII, the entry 'Table C' is replaced by the following:

'— Table C: Breakdown for relevant outputs and measures by type of area, gender and/or age, and by operations contributing to the integration of third-country nationals.'

COMMISSION IMPLEMENTING REGULATION (EU) 2016/1998

of 15 November 2016

withdrawing the acceptance of the undertaking for five exporting producers under Implementing Decision 2013/707/EU confirming the acceptance of an undertaking offered in connection with the anti-dumping and anti-subsidy proceedings concerning imports of crystalline silicon photovoltaic modules and key components (i.e. cells) originating in or consigned from the People's Republic of China for the period of application of definitive measures

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) 2016/1036 of the European Parliament and of the Council of 8 June 2016 on protection against dumped imports from countries not members of the European Union [1] ('the basic anti-dumping Regulation'), and in particular Article 8 thereof,

Having regard to Regulation (EU) 2016/1037 of the European Parliament and of the Council of 8 June 2016 on protection against subsidised imports from countries not members of the European Union [2] ('the basic anti-subsidy Regulation'), and in particular Article 13 thereof,

Informing the Member States,

Whereas:

A. UNDERTAKING AND OTHER EXISTING MEASURES

(1) By Regulation (EU) No 513/2013 [3], the European Commission ('the Commission') imposed a provisional anti-dumping duty on imports into the European Union ('the Union') of crystalline silicon photovoltaic modules ('modules') and key components (i.e. cells and wafers) originating in or consigned from the People's Republic of China ('the PRC').

(2) A group of exporting producers gave a mandate to the China Chamber of Commerce for Import and Export of Machinery and Electronic Products ('CCCME') to submit a price undertaking on their behalf to the Commission, which they did. It is clear from the terms of that price undertaking that it constitutes a bundle of individual price undertakings for each exporting producer, which is, for reasons of practicality of administration, coordinated by the CCCME.

(3) By Decision 2013/423/EU [4], the Commission accepted that price undertaking with regard to the provisional anti-dumping duty. By Regulation (EU) No 748/2013 [5], the Commission amended Regulation (EU) No 513/2013 to introduce the technical changes necessary due to the acceptance of the undertaking with regard to the provisional anti-dumping duty.

(4) By Implementing Regulation (EU) No 1238/2013 [6], the Council imposed a definitive anti-dumping duty on imports into the Union of modules and cells originating in or consigned from the PRC ('the products concerned'). By Implementing Regulation (EU) No 1239/2013 [7], the Council also imposed a definitive countervailing duty on imports into the Union of the products concerned.

(5) Following the notification of an amended version of the price undertaking by a group of exporting producers ('the exporting producers') together with the CCCME, the Commission confirmed by Implementing Decision 2013/707/EU [8] the acceptance of the price undertaking as amended ('the undertaking') for the period of application of definitive measures. The Annex to this Decision lists the exporting producers for whom the undertaking was accepted, inter alia:

(a) Wuxi Suntech Power Co. Ltd, Suntech Power Co. Ltd, Wuxi Sunshine Power Co. Ltd, Luoyang Suntech Power Co. Ltd, Zhenjiang Rietech New Energy Science Technology Co. Ltd and Zhenjiang Ren De New Energy Science Technology Co. Ltd together with their related companies in the Union, jointly covered by the TARIC additional code: B796 ('Wuxi Suntech');

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[1] OJ L 176, 30.6.2016, p. 21.
[2] OJ L 176, 30.6.2016, p. 55.
[3] OJ L 152, 5.6.2013, p. 5.
[4] OJ L 209, 3.8.2013, p. 26.
[5] OJ L 209, 3.8.2013, p. 1.
[6] OJ L 325, 5.12.2013, p. 1.
[7] OJ L 325, 5.12.2013, p. 66.
[8] OJ L 325, 5.12.2013, p. 214.

(b) Jinko Solar Co. Ltd, Jinko Solar Import and Export Co. Ltd, ZHEJIANG JINKO SOLAR CO. LTD and ZHEJIANG JINKO SOLAR TRADING CO. LTD together with their related companies in the Union, jointly covered by the TARIC additional code: B845 ('Jinko Solar');

(c) Risen Energy Co., Ltd, together with its related company in the Union, jointly covered by the TARIC additional code: B868 ('Risen Energy');

(d) JingAo Solar Co. Ltd, Shanghai JA Solar Technology Co. Ltd, JA Solar Technology Yangzhou Co. Ltd, Hefei JA Solar Technology Co. Ltd and Shanghai JA Solar PV Technology Co. Ltd, together with their related company in the Union, jointly covered by TARIC additional code: B794 ('JA Solar'); and

(e) Sumec Hardware & Tools Co. Ltd and Phono Solar Technology Co. Ltd, together with their related companies in the Union, jointly covered by TARIC additional code: B866 ('Sumec').

(6) By Implementing Decision 2014/657/EU ([1]) the Commission accepted a proposal by the exporting producers together with the CCCME for clarifications concerning the implementation of the undertaking for the products concerned covered by the undertaking, that is modules and cells originating in or consigned from the PRC, currently falling within CN codes ex 8541 40 90 (TARIC codes 8541 40 90 21, 8541 40 90 29, 8541 40 90 31 and 8541 40 90 39) produced by the exporting producers ('product covered'). The anti-dumping and countervailing duties referred to in recital 4 above, together with the undertaking, are jointly referred to as 'measures'.

(7) By Implementing Regulation (EU) 2015/866 ([2]) the Commission withdrew the acceptance of the undertaking for three exporting producers.

(8) By Implementing Regulation (EU) 2015/1403 ([3]) the Commission withdrew the acceptance of the undertaking for another exporting producer.

(9) By Implementing Regulation (EU) 2015/2018 ([4]) the Commission withdrew the acceptance of the undertaking for two exporting producers.

(10) The Commission initiated an expiry review investigation of the anti-dumping measures by a Notice of Initiation published in the *Official Journal of the European Union* ([5]) on 5 December 2015.

(11) The Commission initiated an expiry review investigation of the countervailing measures by a Notice of Initiation published in the *Official Journal of the European Union* ([6]) on 5 December 2015.

(12) The Commission also initiated a partial interim review of the anti-dumping and countervailing measures by a Notice of Initiation published in the *Official Journal of the European Union* ([7]) on 5 December 2015.

(13) By Implementing Regulation (EU) 2016/115 ([8]) the Commission withdrew the acceptance of the undertaking for another exporting producer.

(14) By Implementing Regulation (EU) 2016/185 ([9]), the Commission extended the definitive anti-dumping duty imposed by Implementing Regulation (EU) No 1238/2013 on imports of the products concerned originating in or consigned from the People's Republic of China to imports of the product concerned consigned from Malaysia and Taiwan, whether declared as originating in Malaysia and in Taiwan or not.

(15) By Implementing Regulation (EU) 2016/184 ([10]), the Commission extended the definitive countervailing duty imposed by Implementing Regulation (EU) No 1239/2013 on imports of the products concerned originating in or consigned from the People's Republic of China to imports of the product concerned consigned from Malaysia and Taiwan, whether declared as originating in Malaysia and in Taiwan or not.

([1]) OJ L 270, 11.9.2014, p. 6.
([2]) OJ L 139, 5.6.2015, p. 30.
([3]) OJ L 218, 19.8.2015, p. 1.
([4]) OJ L 295, 12.11.2015, p. 23.
([5]) OJ C 405, 5.12.2015, p. 8.
([6]) OJ C 405, 5.12.2015, p. 20.
([7]) OJ C 405, 5.12.2015, p. 33.
([8]) OJ L 23, 29.1.2016, p. 47.
([9]) OJ L 37, 12.2.2016, p. 76.
([10]) OJ L 37, 12.2.2016, p. 56.

(16) By Implementing Regulation (EU) 2016/1045 ([1]) the Commission withdrew the acceptance of the undertaking for another exporting producer.

(17) By Implementing Regulation (EU) 2016/1382 ([2]) the Commission withdrew the acceptance of the undertaking for another five exporting producers.

(18) By Implementing Regulation (EU) 2016/1402 ([3]) the Commission withdrew the acceptance of the undertaking for another three exporting producers.

B. TERMS OF THE UNDERTAKING AND VOLUNTARY WITHDRAWALS

(19) As per the undertaking, any exporting producer may voluntarily withdraw its undertaking at any time during its application.

(20) Wuxi Suntech notified the Commission in August 2016 that it wished to withdraw from the undertaking.

(21) Jinko Solar, Risen Energy, JA Solar and Sumec notified the Commission in September 2016 that they also wished to withdraw from the undertaking.

C. WITHDRAWAL OF THE ACCEPTANCE OF THE UNDERTAKING AND IMPOSITIONS OF DEFINITIVE DUTIES

(22) Therefore, in accordance with Article 8(9) of the basic anti-dumping Regulation, Article 13(9) of the basic anti-subsidy Regulation and also in accordance with the terms of the undertaking, the Commission has concluded that the acceptance of the undertaking for Wuxi Suntech, Jinko Solar, Risen Energy, JA Solar and Sumec shall be withdrawn.

(23) Accordingly, pursuant to Article 8(9) of the basic anti-dumping Regulation and Article 13(9) of the basic anti-subsidy Regulation, the definitive anti-dumping duty imposed by Article 1 of Implementing Regulation (EU) No 1238/2013 and the definitive countervailing duty imposed by Article 1 of Implementing Regulation (EU) No 1239/2013 automatically apply to imports originating in or consigned from the PRC of the product concerned and produced by Wuxi Suntech (TARIC additional code: B796), Jinko Solar (TARIC additional code: B845), Risen Energy (TARIC additional code: B868), JA Solar (TARIC additional code: B794) and Sumec (TARIC additional code: B866) as of the day of entry into force of this Regulation.

(24) For information purposes the table in Annex to this Regulation lists the exporting producers for whom the acceptance of the undertaking by Implementing Decision 2013/707/EU is not affected,

HAS ADOPTED THIS REGULATION:

Article 1

Acceptance of the undertaking in relation to:

(a) Wuxi Suntech Power Co. Ltd, Suntech Power Co. Ltd, Wuxi Sunshine Power Co. Ltd, Luoyang Suntech Power Co. Ltd, Zhenjiang Rietech New Energy Science Technology Co. Ltd and Zhenjiang Ren De New Energy Science Technology Co. Ltd together with their related companies in the Union, jointly covered by the TARIC additional code: B796 ('Wuxi Suntech');

(b) Jinko Solar Co. Ltd, Jinko Solar Import and Export Co. Ltd, ZHEJIANG JINKO SOLAR CO. LTD and ZHEJIANG JINKO SOLAR TRADING CO. LTD together with their related companies in the PRC and in the Union, jointly covered by the TARIC additional code: B845 ('Jinko Solar');

(c) Risen Energy Co., Ltd, together with its related company in the Union, jointly covered by the TARIC additional code: B868 ('Risen Energy');

([1]) OJ L 170, 29.6.2016, p. 5.
([2]) OJ L 222, 17.8.2016, p. 10.
([3]) OJ L 228, 23.8.2016, p. 16.

(d) JingAo Solar Co. Ltd, Shanghai JA Solar Technology Co. Ltd, JA Solar Technology Yangzhou Co. Ltd, Hefei JA Solar Technology Co. Ltd and Shanghai JA Solar PV Technology Co. Ltd, together with their related company in the Union, jointly covered by TARIC additional code: B794 ('JA Solar'); and

(e) Sumec Hardware & Tools Co. Ltd and Phono Solar Technology Co. Ltd, together with their related companies in the Union, jointly covered by TARIC additional code: B866 ('Sumec');

is hereby withdrawn.

Article 2

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

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

Done at Brussels, 15 November 2016.

For the Commission
The President
Jean-Claude JUNCKER

ANNEX

List of companies

Name of the company	TARIC additional code
Jiangsu Aide Solar Energy Technology Co. Ltd	B798
Alternative Energy (AE) Solar Co. Ltd	B799
Anhui Chaoqun Power Co. Ltd	B800
Anji DaSol Solar Energy Science & Technology Co. Ltd	B802
Anhui Schutten Solar Energy Co. Ltd Quanjiao Jingkun Trade Co. Ltd	B801
Anhui Titan PV Co. Ltd	B803
Xi'an SunOasis (Prime) Company Limited TBEA SOLAR CO. LTD XINJIANG SANG'O SOLAR EQUIPMENT	B804
Changzhou NESL Solartech Co. Ltd	B806
Changzhou Shangyou Lianyi Electronic Co. Ltd	B807
CHINALAND SOLAR ENERGY CO. LTD	B808
ChangZhou EGing Photovoltaic Technology Co. Ltd	B811
CIXI CITY RIXING ELECTRONICS CO. LTD ANHUI RINENG ZHONGTIAN SEMICONDUCTOR DEVELOPMENT CO. LTD HUOSHAN KEBO ENERGY & TECHNOLOGY CO. LTD	B812
CSG PVtech Co. Ltd	B814
China Sunergy (Nanjing) Co. Ltd CEEG Nanjing Renewable Energy Co. Ltd CEEG (Shanghai) Solar Science Technology Co. Ltd China Sunergy (Yangzhou) Co. Ltd China Sunergy (Shanghai) Co. Ltd	B809
Dongfang Electric (Yixing) MAGI Solar Power Technology Co. Ltd	B816
EOPLLY New Energy Technology Co. Ltd SHANGHAI EBEST SOLAR ENERGY TECHNOLOGY CO. LTD JIANGSU EOPLLY IMPORT & EXPORT CO. LTD	B817
Zhejiang Era Solar Technology Co. Ltd	B818

Name of the company	TARIC additional code
GD Solar Co. Ltd	B820
Greenway Solar-Tech (Shanghai) Co. Ltd Greenway Solar-Tech (Huaian) Co. Ltd	B821
Konca Solar Cell Co. Ltd Suzhou GCL Photovoltaic Technology Co. Ltd Jiangsu GCL Silicon Material Technology Development Co. Ltd Jiangsu Zhongneng Polysilicon Technology Development Co. Ltd GCL-Poly (Suzhou) Energy Limited GCL-Poly Solar Power System Integration (Taicang) Co. Ltd GCL SOLAR POWER (SUZHOU) LIMITED GCL Solar System (Shuzhou) Limited GCL System Integration Technology Co. Ltd	B850
Guodian Jintech Solar Energy Co. Ltd	B822
Hangzhou Bluesun New Material Co. Ltd	B824
Hanwha SolarOne (Qidong) Co. Ltd	B826
Hengdian Group DMEGC Magnetics Co. Ltd	B827
HENGJI PV-TECH ENERGY CO. LTD	B828
Himin Clean Energy Holdings Co. Ltd	B829
Jetion Solar (China) Co. Ltd Junfeng Solar (Jiangsu) Co. Ltd Jetion Solar (Jiangyin) Co. Ltd	B830
Jiangsu Green Power PV Co. Ltd	B831
Jiangsu Hosun Solar Power Co. Ltd	B832
Jiangsu Jiasheng Photovoltaic Technology Co. Ltd	B833
Jiangsu Runda PV Co. Ltd	B834
Jiangsu Sainty Photovoltaic Systems Co. Ltd Jiangsu Sainty Machinery Imp. And Exp. Corp. Ltd	B835
Jiangsu Seraphim Solar System Co. Ltd	B836
Jiangsu Shunfeng Photovoltaic Technology Co. Ltd Changzhou Shunfeng Photovoltaic Materials Co. Ltd Jiangsu Shunfeng Photovoltaic Electronic Power Co. Ltd	B837
Jiangsu Sinski PV Co. Ltd	B838

Name of the company	TARIC additional code
Jiangsu Sunlink PV Technology Co. Ltd	B839
Jiangsu Zhongchao Solar Technology Co. Ltd	B840
Jiangxi Risun Solar Energy Co. Ltd	B841
Jiangxi LDK Solar Hi-Tech Co. Ltd LDK Solar Hi-Tech (Nanchang) Co. Ltd LDK Solar Hi-Tech (Suzhou) Co. Ltd	B793
Jiangyin Hareon Power Co. Ltd Hareon Solar Technology Co. Ltd Taicang Hareon Solar Co. Ltd Hefei Hareon Solar Technology Co. Ltd Jiangyin Xinhui Solar Energy Co. Ltd Altusvia Energy (Taicang) Co. Ltd	B842
Jiangyin Shine Science and Technology Co. Ltd	B843
Jinzhou Yangguang Energy Co. Ltd Jinzhou Huachang Photovoltaic Technology Co. Ltd Jinzhou Jinmao Photovoltaic Technology Co. Ltd Jinzhou Rixin Silicon Materials Co. Ltd Jinzhou Youhua Silicon Materials Co. Ltd	B795
Juli New Energy Co. Ltd	B846
Jumao Photonic (Xiamen) Co. Ltd	B847
King-PV Technology Co. Ltd	B848
Kinve Solar Power Co. Ltd (Maanshan)	B849
Lightway Green New Energy Co. Ltd Lightway Green New Energy(Zhuozhou) Co. Ltd	B851
Nanjing Daqo New Energy Co. Ltd	B853
NICE SUN PV CO. LTD LEVO SOLAR TECHNOLOGY CO. LTD	B854
Ningbo Huashun Solar Energy Technology Co. Ltd	B856
Ningbo Jinshi Solar Electrical Science & Technology Co. Ltd	B857
Ningbo Komaes Solar Technology Co. Ltd	B858
Ningbo South New Energy Technology Co. Ltd	B861

Name of the company	TARIC additional code
Ningbo Sunbe Electric Ind Co. Ltd	B862
Ningbo Ulica Solar Science & Technology Co. Ltd	B863
Perfectenergy (Shanghai) Co. Ltd	B864
Perlight Solar Co. Ltd	B865
SHANGHAI ALEX SOLAR ENERGY SCIENCE & TECHNOLOGY CO. LTD SHANGHAI ALEX NEW ENERGY CO. LTD	B870
Shanghai BYD Co. Ltd BYD(Shangluo)Industrial Co. Ltd	B871
Shanghai Chaori Solar Energy Science & Technology Co. Ltd	B872
Propsolar (Zhejiang) New Energy Technology Co. Ltd Shanghai Propsolar New Energy Co. Ltd	B873
SHANGHAI SHANGHONG ENERGY TECHNOLOGY CO. LTD	B874
SHANGHAI SOLAR ENERGY S&T CO. LTD Shanghai Shenzhou New Energy Development Co. Ltd Lianyungang Shenzhou New Energy Co. Ltd	B875
Shanghai ST Solar Co. Ltd Jiangsu ST Solar Co. Ltd	B876
Shenzhen Sacred Industry Co. Ltd	B878
Shenzhen Topray Solar Co. Ltd Shanxi Topray Solar Co. Ltd Leshan Topray Cell Co. Ltd	B880
Sopray Energy Co. Ltd Shanghai Sopray New Energy Co. Ltd	B881
SUN EARTH SOLAR POWER CO. LTD NINGBO SUN EARTH SOLAR POWER CO. LTD Ningbo Sun Earth Solar Energy Co. Ltd	B882
SUZHOU SHENGLONG PV-TECH CO. LTD	B883
TDG Holding Co. Ltd	B884
Tianwei New Energy Holdings Co. Ltd Tianwei New Energy (Chengdu) PV Module Co. Ltd Tianwei New Energy (Yangzhou) Co. Ltd	B885

Name of the company	TARIC additional code
Wenzhou Jingri Electrical and Mechanical Co. Ltd	B886
Shanghai Topsolar Green Energy Co. Ltd	B877
Shenzhen Sungold Solar Co. Ltd	B879
Wuhu Zhongfu PV Co. Ltd	B889
Wuxi Saijing Solar Co. Ltd	B890
Wuxi Shangpin Solar Energy Science and Technology Co. Ltd	B891
Wuxi Solar Innova PV Co. Ltd	B892
Wuxi Taichang Electronic Co. Ltd China Machinery Engineering Wuxi Co. Ltd Wuxi Taichen Machinery & Equipment Co. Ltd	B893
Xi'an Huanghe Photovoltaic Technology Co. Ltd State-run Huanghe Machine-Building Factory Import and Export Corporation Shanghai Huanghe Fengjia Photovoltaic Technology Co. Ltd	B896
Yingli Energy (China) Co. Ltd Baoding Tianwei Yingli New Energy Resources Co. Ltd Hainan Yingli New Energy Resources Co. Ltd Hengshui Yingli New Energy Resources Co. Ltd Tianjin Yingli New Energy Resources Co. Ltd Lixian Yingli New Energy Resources Co. Ltd Baoding Jiasheng Photovoltaic Technology Co. Ltd Beijing Tianneng Yingli New Energy Resources Co. Ltd Yingli Energy (Beijing) Co. Ltd	B797
Yuhuan BLD Solar Technology Co. Ltd Zhejiang BLD Solar Technology Co. Ltd	B899
Yuhuan Sinosola Science & Technology Co. Ltd	B900
Zhangjiagang City SEG PV Co. Ltd	B902
Zhejiang Fengsheng Electrical Co. Ltd	B903
Zhejiang Global Photovoltaic Technology Co. Ltd	B904
Zhejiang Heda Solar Technology Co. Ltd	B905
Zhejiang Jiutai New Energy Co. Ltd Zhejiang Topoint Photovoltaic Co. Ltd	B906
Zhejiang Kingdom Solar Energy Technic Co. Ltd	B907

Name of the company	TARIC additional code
Zhejiang Koly Energy Co. Ltd	B908
Zhejiang Mega Solar Energy Co. Ltd Zhejiang Fortune Photovoltaic Co. Ltd	B910
Zhejiang Shuqimeng Photovoltaic Technology Co. Ltd	B911
Zhejiang Shinew Photoelectronic Technology Co. Ltd	B912
Zhejiang Sunflower Light Energy Science & Technology Limited Liability Company Zhejiang Yauchong Light Energy Science & Technology Co. Ltd	B914
Zhejiang Sunrupu New Energy Co. Ltd	B915
Zhejiang Tianming Solar Technology Co. Ltd	B916
Zhejiang Trunsun Solar Co. Ltd Zhejiang Beyondsun PV Co. Ltd	B917
Zhejiang Wanxiang Solar Co. Ltd WANXIANG IMPORT & EXPORT CO LTD	B918
ZHEJIANG YUANZHONG SOLAR CO. LTD	B920
Zhongli Talesun Solar Co. Ltd	B922

COMMISSION IMPLEMENTING REGULATION (EU) 2016/1999

of 15 November 2016

establishing the standard import values for determining the entry price of certain fruit and vegetables

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) No 1308/2013 of the European Parliament and of the Council of 17 December 2013 establishing a common organisation of the markets in agricultural products and repealing Council Regulations (EEC) No 922/72, (EEC) No 234/79, (EC) No 1037/2001 and (EC) No 1234/2007 ([1]),

Having regard to Commission Implementing Regulation (EU) No 543/2011 of 7 June 2011 laying down detailed rules for the application of Council Regulation (EC) No 1234/2007 in respect of the fruit and vegetables and processed fruit and vegetables sectors ([2]), and in particular Article 136(1) thereof,

Whereas:

(1) Implementing Regulation (EU) No 543/2011 lays down, pursuant to the outcome of the Uruguay Round multilateral trade negotiations, the criteria whereby the Commission fixes the standard values for imports from third countries, in respect of the products and periods stipulated in Annex XVI, Part A thereto.

(2) The standard import value is calculated each working day, in accordance with Article 136(1) of Implementing Regulation (EU) No 543/2011, taking into account variable daily data. Therefore this Regulation should enter into force on the day of its publication in the *Official Journal of the European Union*,

HAS ADOPTED THIS REGULATION:

Article 1

The standard import values referred to in Article 136 of Implementing Regulation (EU) No 543/2011 are fixed in the Annex to this Regulation.

Article 2

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

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

Done at Brussels, 15 November 2016.

For the Commission,
On behalf of the President,
Jerzy PLEWA
Director-General for Agriculture and Rural Development

([1]) OJ L 347, 20.12.2013, p. 671.
([2]) OJ L 157, 15.6.2011, p. 1.

ANNEX

Standard import values for determining the entry price of certain fruit and vegetables

(EUR/100 kg)

CN code	Third country code (1)	Standard import value
0702 00 00	MA	92,9
	ZZ	92,9
0707 00 05	TR	141,4
	ZZ	141,4
0709 93 10	MA	105,4
	TR	102,4
	ZZ	103,9
0805 20 10	MA	88,2
	ZZ	88,2
0805 20 30, 0805 20 50, 0805 20 70, 0805 20 90	PE	122,6
	TR	67,3
	ZZ	95,0
0805 50 10	AR	67,2
	CL	69,9
	TR	83,0
	ZZ	73,4
0806 10 10	BR	293,4
	IN	166,9
	PE	319,6
	TR	136,7
	US	353,3
	ZA	345,1
	ZZ	269,2
0808 10 80	CL	174,1
	NZ	139,2
	ZA	122,8
	ZZ	145,4
0808 30 90	CN	44,3
	TR	168,6
	ZZ	106,5

(1) Nomenclature of countries laid down by Commission Regulation (EU) No 1106/2012 of 27 November 2012 implementing Regulation (EC) No 471/2009 of the European Parliament and of the Council on Community statistics relating to external trade with non-member countries, as regards the update of the nomenclature of countries and territories (OJ L 328, 28.11.2012, p. 7). Code 'ZZ' stands for 'of other origin'.

DECISIONS

COUNCIL IMPLEMENTING DECISION (CFSP) 2016/2000

of 15 November 2016

implementing Decision 2013/255/CFSP concerning restrictive measures against Syria

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on European Union, and in particular Article 31(2) thereof,

Having regard to Council Deision 2013/255/CFSP of 31 May 2013 concerning restrictive measures against Syria ([1]), and in particular Article 30(1) thereof,

Having regard to the proposal from the High Representative of the Union for Foreign Affairs and Security Policy,

Whereas:

(1) On 31 May 2013, the Council adopted Decision 2013/255/CFSP, concerning restrictive measures against Syria.

(2) Two entities should be removed from the list of entities subject to restrictive measures, as set out in Section B of Annex I to Decision 2013/255/CFSP.

(3) Annex I to Decision 2013/255/CFSP should therefore be amended accordingly,

HAS ADOPTED THIS DECISION:

Article 1

Annex I to Decision 2013/255/CFSP is amended as set out in the Annex to this Decision.

Article 2

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

Done at Brussels, 15 November 2016.

For the Council
The President
I. KORČOK

([1]) OJ L 147, 1.6.2013, p. 14.

ANNEX

I. The following entities and the related entries are deleted from the list set out in Section B of Annex I to Decision 2013/255/CFSP:

55. Tri-Ocean Trading

55a. Tri-Ocean Energy

COUNCIL DECISION (CFSP) 2016/2001

of 15 November 2016

on a Union contribution to the establishment and the secure management of a Low Enriched Uranium (LEU) Bank under the control of the International Atomic Energy Agency (IAEA) in the framework of the EU Strategy against the Proliferation of Weapons of Mass Destruction

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on European Union, and in particular Article 28 thereof,

Having regard to the proposal of the High Representative of the Union for Foreign Affairs and Security Policy,

Whereas:

(1) On 12 December 2003, the European Council adopted the EU Strategy against the Proliferation of Weapons of Mass Destruction ('the Strategy'), Chapter III of which contains a list of measures to combat such proliferation and which need to be taken both within the Union and in third countries.

(2) The Union is actively implementing the Strategy and is giving effect to the measures listed in Chapter III thereof, in particular through releasing financial resources to support specific projects conducted by multilateral institutions, such as the International Atomic Energy Agency (IAEA).

(3) Article IV of the Treaty on the Non-Proliferation of Nuclear Weapons (NPT) stipulates the inalienable right of all the Parties to the NPT to develop research, production and use of nuclear energy for peaceful purposes without discrimination and in conformity with Articles I and II of the NPT. It also stipulates that all the Parties to the NPT undertake to 'cooperate in contributing alone or together with other States or international organizations to the further development of the applications of nuclear energy for peaceful purposes, especially in the territories of non-nuclear-weapon States Party to the Treaty, with due consideration for the needs of the developing areas of the world'.

(4) Multilateral approaches to the nuclear fuel cycle have the potential to provide countries which have decided to resort to nuclear energy for peaceful uses with an alternative to the development of national nuclear fuel cycles, while avoiding proliferation risks.

(5) Under Article III of its Statute, the IAEA is authorised to perform any operation, including acquiring nuclear fuel, services and equipment and establishing its own facilities and plants, in order to facilitate the practical application of nuclear energy for peaceful purposes.

(6) In September 2006, the Nuclear Threat Initiative (NTI), an independent non-governmental organisation based in the United States, offered a grant of USD 50 000 000 to the IAEA to help create a low enriched uranium stockpile owned and managed by the IAEA on the condition that the Agency should be able to collect an additional amount of USD 100 000 000, including grants from other IAEA Member States and donors, and set up a nuclear fuel reserve.

(7) On 8 December 2008, the Council adopted conclusions in support of the establishment and the secure management of a nuclear fuel bank placed under the control of the IAEA. It also stated that the Union was planning to contribute up to EUR 25 000 000 to that project, once the conditions and modalities for the Bank have been defined and approved by the IAEA Board of Governors. The European Commission has already provided EUR 20 000 000 for the acquisition of the LEU.

(8) On 3 December 2010, the IAEA Board of Governors adopted resolution GOV/2010/70 approving the establishment of an IAEA Low-Enriched Uranium (LEU) Bank and affirmed that the IAEA LEU Bank's operations would be funded exclusively through extra-budgetary contributions.

(9) Paragraph 15 of GOV/2010/67, entitled 'Assurance of Supply': Establishment of an IAEA Low Enriched Uranium (LEU) Bank for the Supply of LEU to Member States provides 'the Agency shall be the owner of the LEU in the IAEA LEU bank and the LEU shall be under its control and in its formal legal possession. The Agency shall be responsible for storing and protecting materials in its possession by ensuring, through any Host State Agreement, that the LEU is safeguarded against natural and other hazards, unauthorized removal or diversion, damage or

destruction, including sabotage, and forcible seizure. In addition, the Agency through any Host State Agreement shall ensure the application of IAEA safeguards to the LEU in the IAEA LEU bank, as well as the application of the safety standards and measures, and the physical protection measures by the Host State or States'. Paragraph 16 of GOV/2010/67 further provides that '[t]he Agency, with Board approval, shall conclude with any Host State a Host State Agreement, similar to the present IAEA Headquarters Agreement, that shall provide for the safety and security and appropriate liability coverage of the storage facility and shall afford those privileges and immunities to the Agency that are necessary for the independent operation of the IAEA LEU bank, including the right to transport LEU to and from the IAEA LEU bank as determined by the Agency in accordance with the Statute and the Host State(s) agreement. In addition, if necessary, guaranteed transit arrangements shall be concluded with States neighbouring the Host State'.

(10) The IAEA LEU Bank will be a stock of up to 60 Type 30B cylinders containing standard commercial low-enriched uranium hexafluoride. The IAEA LEU Bank will be located in the IAEA LEU storage facility, operated by Ulba Metallurgical Plant, and regulated by the Committee for Atomic and Energy Supervision and Control of the Republic of Kazakhstan.

(11) The basic legal framework between the IAEA and the host State Kazakhstan has been concluded. The Transit Agreement with the Russian Federation, approved by the IAEA Board of Governors (GOV/2015/36) has been signed. The design of a new IAEA LEU storage facility has been completed and the IAEA has concluded that it meets the applicable provisions of the IAEA safety standards and security guidance. A detailed cost estimate of the new IAEA LEU storage facility has been undertaken and has been independently validated. A Partnership Agreement between the IAEA and the facility operator that establishes the terms and conditions of cooperation for the construction of the IAEA LEU storage facility has been finalised. The IAEA is now planning activities in preparation for the acquisition of LEU.

(12) According to the Project and Financial Plan as described in the updated report by the IAEA Director-General (GOV/INF/2016/8) 'Assurance of Supply: Establishment of an IAEA Low Enriched Uranium (LEU) Bank for the Supply of LEU to Member States', the total cost of the LEU project is expected to be EUR 118 863 000,

HAS ADOPTED THIS DECISION:

Article 1

1. For the purposes of giving immediate and practical implementation to some elements of the EU Strategy against the Proliferation of Weapons of Mass Destruction, the Union shall contribute to the establishment and the secure management of a Low-Enriched Uranium (LEU) Bank placed under the control of the International Atomic Energy Agency ('IAEA', or 'the Agency') in order to reduce the growing proliferation risks caused by the spread of sensitive nuclear fuel cycle technologies. The Union shall undertake activities to support the IAEA LEU Bank, in the form of a LEU reserve with the following objectives:

(a) to enable countries to enjoy their rights under Article IV of the NPT while avoiding proliferation risks; and

(b) to serve as a mechanism of last resort to support the commercial market without distorting it, in the event that an IAEA member state's supply of LEU is disrupted and cannot be restored by commercial means and that such IAEA Member State fulfils the eligibility criteria.

2. In order to achieve the objectives referred in paragraph 1, the Union shall contribute to the establishment and the secure management of the LEU Bank, under the control of the IAEA, by financing security-related activities, including physical protection, transport, safe guarding, and contributions to the secure management of the LEU Bank. The project shall be carried out for the benefit of all countries which have decided to resort to nuclear energy for peaceful uses.

A detailed description of the project is set out in the Annex.

Article 2

1. The High Representative of the Union for Foreign Affairs and Security Policy (HR) shall be responsible for the implementation of this Decision.

2. The technical implementation of the project referred to in Article 1(2) shall be carried out by the IAEA. It shall perform this task under the control of the HR. For this purpose, the HR shall enter into the necessary arrangements with the IAEA.

Article 3

1. The financial reference amount for the implementation of the activities referred to in Article 1(2) shall be EUR 4 362 200.

2. The expenditure financed by the amount stipulated in paragraph 1 shall be managed in accordance with the procedures and rules applicable to the Union budget.

3. The Commission shall supervise the proper management of the expenditure referred to in paragraph 1. For this purpose, it shall conclude a financing agreement with the IAEA. The agreement shall stipulate that the IAEA is to ensure visibility of the Union contribution, appropriate to its size.

4. The Commission shall endeavour to conclude the financing agreement referred to in paragraph 3 as soon as possible after the entry into force of this Decision. It shall inform the Council of any difficulties in that process and of the date of conclusion of the financing agreement.

Article 4

1. The HR shall report to the Council on the implementation of this Decision on the basis of regular reports prepared by the IAEA. Those IAEA reports shall form the basis for the evaluation carried out by the Council.

2. The Commission shall provide information on the financial aspects of the implementation of the project referred to in Article 1(2).

Article 5

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

It shall expire 60 months after the date of the conclusion of the financing agreement referred to in Article 3(3). However, it shall expire 6 months after its entry into force if no financing agreement has been concluded by that time.

Article 6

This Decision shall be published in the *Official Journal of the European Union*.

Done at Brussels, 15 November 2016.

For the Council
The President
I. KORČOK

ANNEX

Union contribution to the establishment and the secure management of a Low Enriched Uranium (LEU) Bank under the control of the International Atomic Energy Agency (IAEA) in the framework of the EU Strategy against the Proliferation of Weapons of Mass Destruction

I. INTRODUCTION

Background

In December 2010, the IAEA Director-General received a mandate from the Board of Governors to launch the setting-up of a Low Enriched Uranium (LEU) Bank and has presented a detailed plan for its establishment and secure management.

On 20 December 2011, the IAEA confirmed to the Permanent Mission of Kazakhstan to the IAEA that, based on the information provided to the Agency by Kazakhstan in its 'Expression of Interest' and with reference to the requirements set out in document GOV/INF/2011/7, the Ulba Metallurgical Plant (UMP) was suitable as a host site for the IAEA LEU Bank.

Several missions were conducted by the IAEA to Kazakhstan between 2011 and 2016, aimed at assessing the UMP facility and the national regulatory framework, to ensure that the LEU Bank would be able to fulfil the applicable provisions of the IAEA safety standards and security guidance. The assessments were performed in the areas of facility safety, seismic safety, emergency preparedness and response, transport safety and security, and physical protection.

A Host State Agreement (HSA) was signed between the IAEA and Kazakhstan on 27 August 2015. That agreement establishes Kazakhstan as the host state for the IAEA LEU Bank, and provides the legal framework for Kazakhstan to ensure that the IAEA LEU Bank will be managed and regulated in accordance with the laws and regulations of Kazakhstan and in compliance with the applicable provisions of the IAEA safety standards and security guidance.

A facility operator agreement was signed between the IAEA and UMP on 27 August 2015. That agreement establishes UMP as the facility where the IAEA LEU Bank will be located, and provides the legal framework for UMP to operate and manage the IAEA LEU Bank, in accordance with its license, the national regulatory framework, and the applicable provisions of the IAEA safety standards and security guidance.

In addition, the IAEA and Kazakhstan Ministry of Energy signed a Technical Agreement on the specific arrangements to be implemented for the establishment of the IAEA LEU Bank in Kazakhstan ('Technical Agreement'). The Technical Agreement ensures that each party provides the necessary resources for the implementation of its respective activities for the timely establishment of the IAEA LEU Bank, including activities to comply with the applicable provisions of the IAEA safety standards and security guidance. Under the Technical Agreement, the parties established a joint coordination committee (JCC) to facilitate implementation of the Technical Agreement and approved a plan of specific activities (PSA), to ensure the establishment and operation of the IAEA LEU Bank in compliance with the applicable provisions of the IAEA safety standards and security guidance. The Technical Agreement requires that activities be completed in two years following the signing of the legal agreements, or by September 2017.

In November 2015, UMP began designing a new IAEA LEU storage facility to house the IAEA LEU Bank. An IAEA mission visited UMP from 29 February to 4 March 2016 to review design progress. The IAEA mission examined whether or not the applicable provisions of the IAEA safety standards and security guidance had been properly taken into account in the design process. The IAEA mission's review focused on five technical areas: building structure, safety analysis, radiation protection, emergency preparedness and response, and nuclear security. The review of the proposed design and relevant supporting documentation led to the general conclusion that the design provides adequate measures to ensure nuclear safety and security guidance.

Following the completion of the design and its review by the IAEA, in May 2016 a partnership agreement between the IAEA and UMP was signed. That agreement provides for the technical and financial terms of the establishment of the IAEA LEU Storage Facility. It marks a significant milestone in the establishment of the IAEA LEU Bank.

In May 2016, the Board of Governors report GOV/INF/2016/8 highlighted the significant progress achieved. It also provides the first baseline comprehensive Project and Financial Plan.

Board of Governors Document GOV/2010/67, both authorised the IAEA Director-General to establish the IAEA LEU Bank, and required that the costs (including the human resources costs) related to the establishment and operation of the IAEA LEU Bank must be exclusively covered by extra-budgetary voluntary contributions, with no impact to the regular budget of the IAEA. To that end, the IAEA LEU Bank project reimburses various divisions of the IAEA for the technical input and support provided to the project experts and specialists through the use of Service Level Agreements (SLAs). Those agreements, which define the services to be provided by the divisions to the project to execute the project plan (including the PSA), as well as the costs for the level of support provided from each Division, were finalised and agreed in March 2016.

From 1 April 2016, a number of Member States, the Commission, the Nuclear Threat Initiative (NTI) and the World Nuclear Transport Institute (WNTI) have pledged funds totalling approximately USD 124 900 000 and EUR 25 000 000, and the contributions received by the Agency as of this date are USD 124 900 000 and EUR 20 000 000. Financial contributions have been provided by the Nuclear Threat Initiative (USD 50 000 000), US (USD 50 000 000), UAE (USD 10 000 000), Norway (USD 5 000 000), Kuwait (USD 10 000 000), WNTI (EUR 10 000) Kazakhstan (USD 400 000). EUR 20 000 000 donated by the Commission is dedicated to purchase of LEU for the IAEA LEU Bank and up to EUR 5 000 000 have been pledged for security related upgrades. The funds for security related upgrades (up to EUR 5 000 000) are the subject to this Annex.

The next key stages for the IAEA LEU Bank are:

(a) Finalisation of the establishment of the IAEA LEU storage facility, including completing construction; confirmation that the building and equipment correspond to the design intent and the applicable safety and security provisions;

(b) Agreement on a cylinder management programme with UMP to ensure long-term safety and security of the cylinders and their readiness for transport;

(c) Commissioning of the facility;

(d) Acquisition of the IAEA LEU and transporting it to the storage facility;

(e) Start of operations.

Objectives of the project

Contributing to the establishment and the secure management of the IAEA LEU Bank, in particular by ensuring high levels of security and safety during transport and storage, in line with the IAEA safety standards and security guidance.

Benefits

The following benefits will be achieved:

(a) enhancing the assurance of supply of nuclear fuel in a secure and safe manner; and

(b) assisting the IAEA with ensuring the security and safety of transport of LEU from procurement to supply, as well as during storage at the site of the LEU Bank.

II. DESCRIPTION OF THE ACTIVITIES

The LEU Bank

The IAEA LEU Bank will consist of a physical stock of roughly 90 tonnes of LEU, the quantity needed for one initial load for a modern light water reactor (equivalent to approximately 3 core reloads) for electricity generation, along with related equipment and services. The stock will be owned by the IAEA. The IAEA LEU Bank will operate in accordance with certain non-discriminatory criteria for the release of the LEU to a recipient country. Those criteria are fully compatible with the IAEA Statute and have been approved by the Board of Governors. The nuclear facility using the LEU must be covered by a safeguards agreement with the IAEA, and be in full compliance with that agreement.

Union support

The Union will support the IAEA LEU Bank in a complementary manner through different instruments. A financial contribution of EUR 20 000 000 for LEU acquisition was already provided in 2011 from the Instrument for Stability.

This Decision will contribute to the safe and secure operation and management of the IAEA LEU Bank. Relevant activities foreseen by the IAEA, to which this Decision will contribute financially, may include:

1. Supporting the Safe and Secure Establishment of storage for the 90 tonnes of LEU

This item covers the cost of implementation of the project plan activities, including the plan of specific activities (PSA) for the year 2017 and follow-up activities in 2018. The PSA, agreed between the IAEA, UMP, and the Committee for Atomic and Energy Supervision and Control of Kazakhstan, is a list of activities considered necessary for the upgrading of facilities, equipment, procedures, and practices to ensure that the IAEA LEU Bank will be established, stored, operated and protected in accordance with the relevant provisions of the IAEA safety standards and security guidance. These activities were developed on the basis of several assessments, conducted from 2012 to 2016. In particular, an Agency mission in January 2016 also identified some further extrinsic equipment that will be required to enable the storage facility to be operated in accordance with IAEA Standards for emergency preparedness and response.

Activities include the development of procedures for safe and secure operations, procurement of emergency preparedness and response and radiation protection equipment, and provision of associated training; conducting workshops on security topics relevant for the IAEA LEU Bank (e.g. nuclear security culture); observation of emergency exercises at the facility; and follow-up assessment activities to validate the compliance of upgrades with the relevant IAEA safety standards and security guidance prior to commissioning.

The LEU will only be placed in the IAEA LEU storage facility when the IAEA is satisfied that the IAEA LEU Bank has been established and complies with the applicable provisions of the IAEA safety standards and security guidance. Therefore, the IAEA will undertake a confirmation mission, planned for the summer of 2017, to confirm that the completed building and its key equipment correspond to the design intent and that the full infrastructure needed to meet the applicable safety and security provisions is in place.

Over a period of two years the budget will include the cost of planning, executing, supporting, and reporting on the project plan and PSA activities, inclusive of costs to cover IAEA human resources to ensure that, in accordance with the mandate of the IAEA LEU Bank project, no regular budget funds are used.

2. Ensuring the secure transport of 90 tonnes of LEU

It is expected that the LEU 90 tonnes will be transported from the facility of the vendor, or vendors, to the Ulba Metallurgical Plant, in Oskemen, Kazakhstan, where the IAEA LEU Bank will be located. Shipments of LEU moving through various jurisdictions, must satisfy all documentation, insurance, transit approval, and marking requirements, including States' requirements related to physical protection which are contained in relevant conventions and recommendations developed under the auspices of the IAEA, and the security requirements of the International Maritime Organisation (IMO). Expenditure associated with the transport of the LEU will depend on the geographic location of the LEU vendor and the maritime and overland transport distances between the vendor and Ulba Metallurgical Plant (UMP) in Kazakhstan, and the number of ports of call and border transits required for completing delivery. The item includes the development of technical specifications with IAEA officials and external experts specialist input; the planning and supervision of the transport; insurance; chartering of a maritime transport vessel for enhanced security; consultation on transport route security risk assessment and planning; as well as guarding during maritime transport, ports of call and transit locations.

3. Ensuring the long term storage of the 90 tonnes of LEU

The LEU will be located at a dedicated storage facility within the UMP territory, located in Oskemen, Kazakhstan. The IAEA LEU will be stored in 30B cylinders. The facility operator, on behalf of the IAEA, will be responsible for storing and protecting the LEU, in compliance with the IAEA safety standards and security guidance. This includes the purchase of the 30B cylinders, which will provide physically safe and secure storage of the LEU. Safety studies indicate that these containment vessels provide robust, safe storage for up to 50 years. In addition to providing safety, the thickness of the walls of the cylinders and the overall design of the cylinders contributes to physical protection from sabotage and theft.

A key activity in ensuring the long term storage of the LEU is the implementation of a cylinder management programme, including routine inspection and recertification of 30B cylinders in storage, for compliance with the ISO 7195 standard, as well as for assurance of readiness for transport to Member States. This requires procurement of services for re-certification of cylinders by authorised inspectors as well as services from UMP staff to support the performance of the testing.

In addition, during long-term storage of the IAEA LEU, the IAEA will undertake routine and ad-hoc monitoring activities, including receipt, review, and verification of annual reports from UMP. This will include annual meetings with UMP to review the safety and security activities related to the IAEA LEU Bank, as well as other types of missions to UMP to ensure that the relevant provisions of the IAEA safety standards and security guidance continue to be applied to the IAEA LEU Bank.

This item will cover a period of 5 years.

III. DURATION

The estimated duration of the implementation period of the project is 60 months, starting with the signing of the financing agreement referred to in Article 3.

IV. BENEFICIARIES

The beneficiaries of the project in this Decision are all eligible recipient states of services of the IAEA LEU Bank, fulfilling the conditions for access to the LEU Bank as established by the IAEA Board of Governors.

V. IMPLEMENTING ENTITY

The IAEA will be entrusted with the technical implementation of the project, as described above, under the control of the HR. The project will be implemented directly by staff of the IAEA, experts from other national nuclear authorities and contractors. In the case of contractors, the procurement of any goods, works or services by the IAEA in the context of this Decision will be carried out as detailed in the financing agreement to be concluded by the Commission with the IAEA.

VI. REPORTING

The implementing entity will prepare:

(a) regular reports on the implementation of the project;

(b) a final report not later than two months after the end of the implementation of the project.

Reports will be sent to the HR.

COMMISSION IMPLEMENTING DECISION (EU) 2016/2002

of 8 November 2016

amending Annex E to Council Directive 91/68/EEC, Annex III to Commission Decision 2010/470/EU and Annex II to Commission Decision 2010/472/EU concerning trade in and imports into the Union of ovine and caprine animals and semen of animals of the ovine and caprine species in relation to the rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies

(notified under document C(2016) 7026)

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 91/68/EEC of 28 January 1991 on animal health conditions governing intra-Community trade in ovine and caprine animals ([1]), and in particular Article 14(2) thereof,

Having regard to Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC ([2]), and in particular the fourth indent of Article 11(2), Article 17(2)(b), the first indent of Article 18(1) and the introductory phrase and point (b) of Article 19 thereof,

Whereas:

(1) Directive 91/68/EEC lays down the animal health conditions governing intra-Union trade in ovine and caprine animals. It provides, inter alia, that ovine and caprine animals must be accompanied during transportation to their destination by a health certificate conforming to Model I, II or III set out in Annex E thereto.

(2) Regulation (EC) No 999/2001 of the European Parliament and of the Council ([3]) lays down rules for the prevention, control and eradication of transmissible spongiform encephalopathies (TSEs) in bovine, ovine and caprine animals. Annex VII to that Regulation sets out the measures for the control and eradication of TSEs. In addition, Chapter A of Annex VIII to that Regulation lays down, inter alia, the conditions for intra-Union trade in live animals.

(3) Regulation (EC) No 999/2001 was recently amended by Commission Regulation (EU) 2016/1396 ([4]). Those amendments provide, inter alia, for an exemption from the conditions set out in point 4.1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, which are aimed at preventing the spread of classical scrapie in farmed animals kept on holdings, for ovine and caprine animals moved exclusively between approved bodies, institutes or centres as defined in Article 2(1)(c) of Directive 92/65/EEC.

(4) Regulation (EU) 2016/1396 also introduces specific conditions for intra-Union trade in ovine and caprine animals of rare breeds which do not comply with the requirements of point 4.1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001. Those specific conditions were introduced to maintain a possibility for regular exchange of such animals between Member States in order to avoid inbreeding and to preserve the genetic diversity in rare breed populations.

([1]) OJ L 46, 19.2.1991, p. 19.
([2]) OJ L 268, 14.9.1992, p. 54.
([3]) Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 147, 31.5.2001, p. 1).
([4]) Commission Regulation (EU) 2016/1396 of 18 August 2016 amending certain Annexes to Regulation (EC) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 225, 19.8.2016, p. 76).

(5) The health certificates conforming to Models II and III set out in Annex E to Directive 91/68/EEC should therefore be amended in order to reflect the requirements relating to intra-Union trade in ovine and caprine animals of rare breeds or of those moved between approved bodies, institutes or centres laid down in Regulation (EC) No 999/2001, as amended by Regulation (EU) 2016/1396.

(6) In addition, some Member States notified the Commission of problems related to additional administrative work caused by the obligation to provide in point I.31 of health certificates conforming to Models I, II and III set out in Annex E to Directive 91/68/EEC details such as breed and quantity of animals forming the consignment. To reduce administrative burden for the official veterinarians, it is appropriate to remove from point I.31 of those model health certificates information on the breed, as such information is not necessary in relation to the health status of the animals in the consignment, and on the quantity of those animals, as such information is already stated in point I.20 and an official identification number of each individual animal must be provided in point I.31.

(7) Furthermore, in order to state more precisely the conditions for individual identification of the animals in points II.5 and II.6 of the health certificates conforming to Models II and III in Annex E to Directive 91/68/EEC, it is necessary to introduce in those points a reference to Council Regulation (EC) No 21/2004 ([1]).

(8) Directive 91/68/EEC should therefore be amended accordingly.

(9) Directive 92/65/EEC lays down conditions applicable to trade in and imports into the Union, inter alia, of semen of animals of the ovine and caprine species.

(10) Annex III to Commission Decision 2010/470/EU ([2]) lays down model health certificates for trade within the Union in consignments of semen of animals of the ovine and caprine species. Part A of that Annex sets out the model health certificate for semen collected after 31 August 2010 and dispatched from an approved semen collection centre of origin of the semen.

(11) Annex II to Commission Decision 2010/472/EU ([3]) lays down, inter alia, model health certificates for the imports into the Union of consignments of semen of animals of the ovine and caprine species. Section A of Part 2 of that Annex sets out the model health certificate for semen dispatched from an approved semen collection centre of origin of the semen.

(12) Point 4.2 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 sets out the scrapie-related conditions to be fulfilled for intra-Union trade in semen of ovine and caprine animals. Chapter H of Annex IX to Regulation (EC) No 999/2001 sets out the scrapie-related conditions to be fulfilled for imports of semen of ovine and caprine animals.

(13) Regulation (EU) 2016/1396 introduces specific conditions for semen collection centres amongst the conditions for a holding to be recognised as having a negligible risk or a controlled risk of classical scrapie in points 1.2 and 1.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, given that the risk of spreading scrapie via male ovine and caprine animals kept at semen collection centres approved and supervised in accordance with the conditions set out in Annex D to Directive 92/65/EEC is limited. A reference to those specific conditions is also introduced in the conditions for trade in and import of semen of ovine and caprine animals set out in Annexes VIII and IX to Regulation (EC) No 999/2001 respectively.

(14) The model health certificate for intra-Union trade in consignments of semen of animals of the ovine and caprine species set out in Part A of Annex III to Decision 2010/470/EU and the model health certificate for imports into the Union of consignments of semen of animals of the ovine and caprine species set out in Section A of Part 2 of Annex II to Decision 2010/472/EU should therefore be amended in order to reflect the requirements relating to semen collection centres laid down in Regulation (EC) No 999/2001, as amended by Regulation (EU) 2016/1396.

([1]) Council Regulation (EC) No 21/2004 of 17 December 2003 establishing a system for the identification and registration of ovine and caprine animals and amending Regulation (EC) No 1782/2003 and Directives 92/102/EEC and 64/432/EEC (OJ L 5, 9.1.2004, p. 8).

([2]) Commission Decision 2010/470/EU of 26 August 2010 laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (OJ L 228, 31.8.2010, p. 15).

([3]) Commission Decision 2010/472/EU of 26 August 2010 on imports of semen, ova and embryos of animals of the ovine and caprine species into the Union (OJ L 228, 31.8.2010, p. 74).

(15) In addition, Chapter H of Annex IX to Regulation (EC) No 999/2001, as amended by Regulation (EU) 2016/1396, provides that meat-and-bone meal should be understood as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE) ([1]), rather than in point 27 of Annex I to Commission Regulation (EC) No 142/2011 ([2]).

(16) Therefore, point II.4.10.4 of the model health certificate for imports into the Union of consignments of semen of animals of the ovine and caprine species set out in Section A of Part 2 of Annex II to Decision 2010/472/EU should be amended according to the amended provisions of Chapter H of Annex IX to Regulation (EC) No 999/2001.

(17) Decisions 2010/470/EU and 2010/472/EU should therefore be amended accordingly.

(18) Regulation (EU) 2016/1396 provides that the amendments made to Annex IX to Regulation (EC) No 999/2001 and related to imports of certain commodities are to apply from 1 July 2017. In addition, to avoid any disruption of imports into the Union of consignments of semen of ovine and caprine animals, the use of certificates issued in accordance with Decision 2010/472/EU as applicable prior to the amendments being introduced by this Decision should be authorised during a transitional period subject to certain conditions.

(19) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS DECISION:

Article 1

Annex E to Directive 91/68/EEC is amended in accordance with Annex I to this Decision.

Article 2

Annex III to Decision 2010/470/EU is amended in accordance with Annex II to this Decision.

Article 3

Annex II to Decision 2010/472/EU is amended in accordance with Annex III to this Decision.

Article 4

Article 3 of this Decision shall apply from 1 July 2017.

For a transitional period until 31 December 2017, consignments of semen of ovine and caprine animals, accompanied by a health certificate issued in accordance with the model set out in Section A of Part 2 of Annex II to Decision 2010/472/EU, as applicable before the amendments made by this Decision, shall be authorised for importation into the Union provided that the certificate was issued no later than 30 November 2017.

([1]) http://www.oie.int/index.php?id=169&L=0&htmfile=glossaire.htm

([2]) Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (OJ L 54, 26.2.2011, p. 1).

Article 5

This Decision is addressed to the Member States.

Done at Brussels, 8 November 2016.

For the Commission
Vytenis ANDRIUKAITIS
Member of the Commission

ANNEX I

Annex E to Directive 91/68/EEC is replaced by the following:

'ANNEX E

MODEL I

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6. No(s) of related original certificates	No(s) of accompanying documents
		I.7. Dealer Name	Approval number
I.8. Country of origin / ISO code	I.9. Region of origin / Code	I.10. Country of destination / ISO code	I.11. Region of destination / Code
I.12. Place of origin Holding ☐ Assembly centre ☐ Dealer's premises ☐ Name Approval number Address Postal code		I.13. Place of destination Holding ☐ Assembly centre ☐ Dealer's premises ☐ Name Approval number Address Postal code	
I.14. Place of loading Postal code		I.15. Date and time of departure	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification: Number(s):		I.17. Transporter Name Approval number Address Postal code Member State	
I.18. Description of commodity		I.19. Commodity code (CN code)	
		I.20. Quantity	

I.21.	I.22. Number of packages
I.23. Seal/Container No	I.24.
I.25. Commodities certified for: Slaughter ☐	
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code
I.28. Export ☐ Third country ISO code Exit point Code	I.29. Estimated journey time
I.30. Route plan Yes ☐ No ☐	
I.31. Identification of the commodities Species (Scientific name) / Official individual identification / Age / Sex	

European Union **91/68 EI Ovine/Caprine for slaughter**

II.	Health information	II.a. Certificate reference number	II.b. Local reference number

Part II: Certification

I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:

([1]) *either* [II.1. The animals were born and have been reared since birth on Union territory.]

([1]) *or* [II.1. The animals were imported from a third country in accordance with Commission Regulation (EU) No 206/2010 at least 30 days prior to loading.]

II.2. The animals:

II.2.1. have been inspected today (within 24 hours prior to loading) and show no clinical sign of disease;

II.2.2. are not animals which are to be destroyed under a scheme to eradicate a contagious or infectious disease;

II.2.3. come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of rabies, for the last 15 days in the case of anthrax, and, have not been in contact with animals from holdings which did not comply with these conditions;

II.2.4. do not come from a holding and have not been in contact with animals from holdings in a protection zone which has been set up under Union legislation and from which animals are prohibited from leaving;

II.2.5. are not the subject of animal health measures pursuant to Union legislation on foot-and-mouth disease and have not been vaccinated against foot-and-mouth disease.

II.3. Based on the written declaration made by the keeper or an examination of the holding register and movement documents kept in accordance with Council Regulation (EC) No 21/2004, in particular in Sections B and C of the Annex to that Regulation:

II.3.1. the animals have remained on a single holding of origin for a period of at least the last 21 days, or on the holding of origin since birth where the animals are less than 21 days old, and no biungulate animal imported from a third country has been introduced into the holding of origin during the last 30 days, unless those animals were introduced in accordance with Article 4a(2) of Council Directive 91/68/EEC, and

([1]) *either* [have remained on a single holding of origin into which no animals of the ovine or caprine species have been introduced, unless those animals were introduced in accordance with Article 4a(1) of Directive 91/68/EEC, during the last 21 days.]

([1]) *or* [are to be consigned directly from a single holding to the slaughterhouse of destination.]

II.4.1. The animals were transported using means of transport and containment which had, before-hand, been cleansed and disinfected using an officially approved disinfectant, and in such a way as to provide effective protection of the animals' health status.

II.4.2. Based on the official documentation accompanying the animals, the consignment covered by this health certificate is due to start the journey on ..... (*insert date*) ([2]).

II.4.3. At the time of inspection the animals covered by this health certificate were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005 ([3]) ([4]).

II.5. This certificate

([1]) *either* [is valid for 10 days from the date of inspection on the holding of origin, or in the approved assembly centre or approved dealer's premises in the Member State of origin;]

([1]) *or* [expires in accordance with Article 9(6) of Directive 91/68/EEC on ..... (*insert date*)] ([5]).

European Union **91/68 EI Ovine/Caprine for slaughter**

II. Health information	II.a. Certificate reference number	II.b. Local reference number

Notes

Part I:

— Box reference I.19: Use the appropriate CN code under the following headings: 0104 10 or 0104 20.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) must be included.

— Box reference I.31: *Identification system:* The animals must bear: An individual number which permits tracing of their premises of origin, in accordance with Council Regulation (EC) No 21/2004.

Age: (months).

Sex: (M = male, F = female, C = castrated).

Part II:

([1]) Delete where not applicable.

([2]) In the case where a consignment is grouped in an assembly centre and comprises animals that were loaded on different dates, the date on which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the holding of origin.

([3]) This statement does not exempt transporters from their obligations in accordance with Union rules in force in particular regarding the fitness of animals to be transported.

([4]) To be completed in case of consignment grouped in an approved assembly centre or in approved dealer's premises.

([5]) To be completed in case of consignment grouped in an approved assembly centre located in the Member State of transit.

— The colour of the stamp and the signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local Veterinary Unit: LVU No:

Date: Signature:

Stamp:

MODEL II

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6. No(s) of related original certificates	No(s) of accompanying documents
		I.7. Dealer Name	Approval number
I.8. Country of origin / ISO code / I.9. Region of origin / Code		I.10. Country of destination / ISO code / I.11. Region of destination / Code	
I.12. Place of origin Holding ☐ Assembly centre ☐ Name Approval/registration number Address Postal code		I.13. Place of destination Holding ☐ Assembly centre ☐ Dealer's premises ☐ Name Approval number Address Postal code	
I.14. Place of loading Postal code		I.15. Date and time of departure	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification: Number(s):		I.17. Transporter Name Approval number Address Postal code Member State	
I.18. Description of commodity		I.19. Commodity code (CN code)	
		I.20. Quantity	

I.21.	I.22. Number of packages
I.23. Seal/Container No	I.24.
I.25. Commodities certified for: Fattening ☐	
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code
I.28. Export ☐ Third country ISO code Exit point Code	I.29. Estimated journey time
I.30. Route plan Yes ☐ No ☐	
I.31. Identification of the commodities Species (Scientific name) / Official individual identification / Age / Sex	

European Union **91/68 EII Ovine/Caprine for fattening**

II.	Health information	II.a. Certificate reference number	II.b. Local reference number

Part II: Certification

I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:

([1]) *either* [II.1. The animals were born and have been reared since birth on Union territory.]

([1]) *or* [II.1. The animals were imported from a third country in accordance with Commission Regulation (EU) No 206/2010 at least 30 days prior to loading.]

II.2. The animals:

II.2.1. have been inspected today (within 24 hours prior to loading) and show no clinical sign of disease;

II.2.2. are not animals which are to be destroyed under a scheme to eradicate a contagious or infectious disease;

II.2.3. come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of rabies, for the last 15 days in the case of anthrax, and, have not been in contact with animals from holdings which did not comply with these conditions;

II.2.4. do not come from a holding and have not been in contact with animals from holdings in a protection zone which has been set up under Union legislation and from which animals are prohibited from leaving;

II.2.5. are not the subject of animal health measures pursuant to Union legislation on foot-and-mouth disease and have not been vaccinated against foot-and-mouth disease.

II.3. Based on the written declaration made by the keeper or an examination of the holding register and movement documents kept in accordance with Council Regulation (EC) No 21/2004, in particular in Sections B and C of the Annex to that Regulation, the animals have remained on a single holding of origin for a period of at least the last 30 days, or on the holding of origin since birth where the animals are less than 30 days old, and no animal of the ovine and caprine species has been introduced into the holding of origin during the last 21 days and no biungulate animal imported from a third country has been introduced into the holding of origin during the last 30 days, unless those animals were introduced in accordance with Article 4a(1) of Council Directive 91/68/EEC.

([1]) [II.4. The animals comply with the additional guarantees provided for in Articles 7 or 8 of Directive 91/68/EEC and laid down for the Member State of destination or part of its territory ..... (*insert Member State or part of its territory*) in Commission Decision …/…/… (*insert number*).]

II.5. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is officially brucellosis-free (*B. melitensis*):

([1]) *either* [the holding of origin is situated in a Member State or part of its territory ..... (*insert name of Member State or part of its territory*) which is recognised as being officially brucellosis-free in accordance with Commission Decision …/…/… (*insert number*).]

([1]) *or* [they come from an officially brucellosis-free (*B. melitensis*) holding.]

([1]) *or* [they come from a brucellosis-free (*B. melitensis*) holding, and

(i) are identified individually in accordance with Council Regulation (EC) No 21/2004,

(ii) have never been vaccinated against brucellosis or have not been vaccinated against brucellosis in the last two years or the animals are females over two years old which were vaccinated against brucellosis before the age of seven months,

(iii) were isolated under official supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]

European Union **91/68 EII Ovine/Caprine for fattening**

II.	Health information	II.a. Certificate reference number	II.b. Local reference number

II.6. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is brucellosis-free (*B. melitensis*):

([1]) *either* [they come from an officially brucellosis-free (*B. melitensis*) holding.]

([1]) *and/or* [they come from a brucellosis-free (*B. melitensis*) holding.]

([1]) *and/or* [until the qualifying date under eradication plans approved pursuant to Council Decision 90/242/EEC, they originate from a holding other than officially brucellosis-free or brucellosis-free and comply with the following conditions:

(i) are identified individually in accordance with Council Regulation (EC) No 21/2004,

(ii) originate from a holding in which all the animals of species susceptible to brucellosis (*B. melitensis*) have been free of clinical symptoms or any other symptoms of brucellosis for at least the last 12 months; and

([1]) *either* [have not been vaccinated against brucellosis (*B. melitensis*) in the last two years, and were isolated under veterinary supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]]

([1]) *or* [were vaccinated with Rev. 1 vaccine before the age of seven months, but not later than 15 days before their introduction into the holding of destination.]]

([1]) [II.7. The animals are intended for a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 of the European Parliament and of the Council as having a negligible risk status for classical scrapie, or for a Member State listed in point 3.2 of that Section as having an approved national scrapie control programme, and

([1]) *either* [come from a holding situated in a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]]

([1]) *and/or* [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1 of that Section.]]

([1]) *and/or* [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species of the ARR/ARR prion protein genotype.]]

([1]) *and/or* [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]]

([1]) *or* [comply with the conditions set out in point 4.1(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

II.8.1. The animals were transported using means of transport and containment which had, before-hand, been cleansed and disinfected using an officially approved disinfectant, and in such a way as to provide effective protection of the animals' health status.

II.8.2. Based on the official documentation accompanying the animals, the consignment covered by this health certificate is due to start the journey on ..... (*insert date*) ([2]).

II.8.3. At the time of inspection the animals covered by this health certificate were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005 ([3]).

European Union **91/68 EII Ovine/Caprine for fattening**

II. Health information	II.a. Certificate reference number	II.b. Local reference number

Notes

Part I:

— Box reference I.19: Use the appropriate CN code under the following headings: 0104 10 or 0104 20.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) must be included.

— Box reference I.31: *Identification system:* The animals must bear: An individual number which permits tracing of their premises of origin, in accordance with Council Regulation (EC) No 21/2004.

Age: (months).

Sex: (M = male, F = female, C = castrated).

Part II:

([1]) Delete where not applicable.

([2]) In the case where a consignment is grouped in an assembly centre and comprises animals that were loaded on different dates, the date on which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the holding of origin.

([3]) This statement does not exempt transporters from their obligations in accordance with Union rules in force in particular regarding the fitness of animals to be transported.

— This certificate is valid for 10 days.

— The colour of the stamp and the signature must be different from that of the other particulars in this certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local Veterinary Unit: LVU No:

Date: Signature:

Stamp:

MODEL III

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6. No(s) of related original certificates	No(s) of accompanying documents
		I.7. Dealer Name	Approval number
I.8. Country of origin / ISO code	I.9. Region of origin / Code	I.10. Country of destination / ISO code	I.11. Region of destination / Code
I.12. Place of origin Holding ☐ Assembly centre ☐ Name Approval/registration number Address Postal code		I.13. Place of destination Holding ☐ Assembly centre ☐ Dealer's premises ☐ Name Approval number Address Postal code	
I.14. Place of loading Postal code		I.15. Date and time of departure	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification: Number(s):		I.17. Transporter Name Approval number Address Postal code Member State	
I.18. Description of commodity		I.19. Commodity code (CN code)	
		I.20. Quantity	

I.21.	I.22. Number of packages
I.23. Seal/Container No	I.24.

I.25. Commodities certified for:

Breeding ☐

I.26. Transit through third country ☐		I.27. Transit through Member States ☐	
Third country	ISO code	Member State	ISO code
Exit point	Code	Member State	ISO code
Entry point	BIP No	Member State	ISO code

I.28. Export ☐		I.29. Estimated journey time
Third country	ISO code	
Exit point	Code	

I.30. Route plan

Yes ☐ No ☐

I.31. Identification of the commodities

Species (Scientific name)	Official individual identification	Age	Sex

European Union **91/68 EIII Ovine/Caprine for breeding**

II.	Health information	II.a. Certificate reference number	II.b. Local reference number

Part II: Certification

I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:

([1]) *either* [II.1. The animals were born and have been reared since birth on Union territory.]

([1]) *or* [II.1. The animals were imported from a third country in accordance with Commission Regulation (EU) No 206/2010 at least 30 days prior to loading.]

II.2. The animals:

II.2.1. have been inspected today (within 24 hours prior to loading) and show no clinical sign of disease;

II.2.2. are not animals which are to be destroyed under a scheme to eradicate a contagious or infectious disease;

II.2.3. come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of rabies, for the last 15 days in the case of anthrax, and, have not been in contact with animals from holdings which did not comply with these conditions;

II.2.4. do not come from a holding and have not been in contact with animals from holdings in a protection zone which has been set up under Union legislation and from which animals are prohibited from leaving;

II.2.5. are not the subject of animal health measures pursuant to Union legislation on foot-and-mouth disease and have not been vaccinated against foot-and-mouth disease.

II.3. Based on the written declaration made by the keeper or an examination of the holding register and movement documents kept in accordance with Council Regulation (EC) No 21/2004, in particular in Sections B and C of the Annex to that Regulation, the animals have remained on a single holding of origin for a period of at least the last 30 days, or on the holding of origin since birth where the animals are less than 30 days old, and no animal of the ovine and caprine species has been introduced into the holding of origin during the last 21 days and no biungulate animal imported from a third country has been introduced into the holding of origin during the last 30 days, unless those animals were introduced in accordance with Article 4a(1) of Council Directive 91/68/EEC.

([1]) [II.4. The animals comply with the additional guarantees provided for in Articles 7 or 8 of Directive 91/68/EEC and laid down for the Member State of destination or part of its territory ..... (*insert Member State or part of its territory*) in Commission Decision …/…/… (*insert number*).]

II.5. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is officially brucellosis-free (*B. melitensis*):

([1]) *either* [the holding of origin is situated in a Member State or part of its territory ..... (*insert name of Member State or part of its territory*) which is recognised as being officially brucellosis-free in accordance with Commission Decision …/…/… (*insert number*).]

([1]) *or* [they come from an officially brucellosis-free (*B. melitensis*) holding.]

([1]) *or* [they come from a brucellosis-free (*B. melitensis*) holding, and

(i) are identified individually in accordance with Council Regulation (EC) No 21/2004,

(ii) have never been vaccinated against brucellosis or have not been vaccinated against brucellosis in the last two years or the animals are females over two years old which were vaccinated against brucellosis before the age of seven months,

(iii) were isolated under official supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]

European Union **91/68 EIII Ovine/Caprine for breeding**

II.	Health information	II.a. Certificate reference number	II.b. Local reference number

II.6. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is brucellosis-free (*B. melitensis*):

([1]) *either* [they come from an officially brucellosis-free (*B. melitensis*) holding.]

([1]) *or* [they come from a brucellosis-free (*B. melitensis*) holding.]

([1]) *or* [until the qualifying date under eradication plans approved pursuant to Council Decision 90/242/EEC, they originate from a holding other than officially brucellosis-free or brucellosis-free and satisfy the following conditions:

(i) are identified individually in accordance with Council Regulation (EC) No 21/2004,

(ii) originate from a holding in which all the animals of species susceptible to brucellosis (*B. melitensis*) have been free of clinical symptoms or any other symptoms of brucellosis for at least the last 12 months; and

([1]) *either* [have not been vaccinated against brucellosis (*B. melitensis*) in the last two years, and were isolated under veterinary supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]]

([1]) *or* [were vaccinated with Rev. 1 vaccine before the age of seven months, and were not vaccinated in the 15 days before the date of emission of this health certificate.]]

([1]) [II.7. They are uncastrated breeding rams and:

(i) come from a holding on which no case of contagious epididymitis of rams (*B. ovis*) has been recorded in the last 12 months,

(ii) have been kept permanently on that holding for the last 60 days,

(iii) have undergone, within the last 30 days, with a negative result, a test to detect contagious epididymitis of rams (*B. ovis*) in accordance with Annex D to Directive 91/68/EEC.]

II.8. To the best of the knowledge of the undersigned and according to the written declaration made by the owner, the animals were not obtained from a holding and have not been in contact with animals from a holding in which the following diseases have been clinically detected:

(i) within the last six months, contagious agalactia of sheep (*Mycoplasma agalactiae*) and contagious agalactia of goats (*Mycoplasma agalactiae, M. capricolum, M. mycoides subsp. mycoides large colony*),

(ii) within the last 12 months, paratuberculosis or caseous lymphadenitis,

(iii) within the last three years, pulmonary adenomatosis, maedi/visna or caprine viral arthritis/encephalitis. However, this time limit is reduced to 12 months if animals affected by maedi/visna or caprine viral arthritis/encephalitis have been slaughtered and the remaining animals have reacted negatively to two tests.

([1]) *either* [II.9. The animals are intended for a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or for a Member State listed in point 3.2 of that Section as having an approved national scrapie control programme, and

([1]) *either* [come from a holding situated in a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]]

([1]) *and/or* [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1 of that Section.]]

European Union **91/68 EIII Ovine/Caprine for breeding**

II. Health information	II.a. Certificate reference number	II.b. Local reference number

([1]) *and/or* [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species of the ARR/ARR prion protein genotype.]]

([1]) *and/or* [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]]

([1]) *or* [comply with the conditions set out in point 4.1(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

([1]) *or* [II.9. The animals are intended for a Member State or zone of a Member State other than those listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or other than those listed in point 3.2 of that Section as having an approved national scrapie control programme, and

([1]) *either* [come from a holding situated in a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]]

([1]) *and/or* [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1 of that Section.]]

([1]) *and/or* [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species of the ARR/ARR prion protein genotype.]]

([1]) *and/or* [come from a holding recognised as having a controlled risk of classical scrapie in accordance with point 1.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1 of that Section.]]

([1]) *and/or* [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]]

([1]) *or* [comply with the conditions set out in point 4.1(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

II.10.1. The animals were transported using means of transport and containment which had, before-hand, been cleansed and disinfected using an officially approved disinfectant, and in such a way as to provide effective protection of the animals' health status.

II.10.2. Based on the official documentation accompanying the animals, the consignment covered by this health certificate is due to start the journey on ..... (*insert date*) ([2]).

II.10.3. At the time of inspection the animals covered by this health certificate were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005 ([3]).

Notes

Part I:

— Box reference I.19: Use the appropriate CN code under the following headings: 0104 10 or 0104 20.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) must be included.

— Box reference I.31: *Identification system*: The animals must bear an individual number which permits tracing of their premises of origin in accordance with Council Regulation (EC) No 21/2004.

Age: (months).

Sex: (M = male, F = female, C = castrated).

European Union **91/68 EIII Ovine/Caprine for breeding**

II. Health information	II.a. Certificate reference number	II.b. Local reference number

Part II:

([1]) Delete where not applicable.

([2]) In the case where a consignment is grouped in an assembly centre and comprises animals that were loaded on different dates, the date at which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the holding of origin.

([3]) This statement does not exempt transporters from their obligations in accordance with Union rules in force in particular regarding the fitness of animals to be transported.

— This certificate is valid for 10 days.

— The colour of the stamp and the signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local Veterinary Unit: LVU No:

Date: Signature:

Stamp: ’

ANNEX II

In Annex III to Decision 2010/470/EU, Part A is replaced by the following:

'Part A

Model health certificate IIIA for trade within the Union in consignments of semen of animals of the ovine and caprine species collected in accordance with Council Directive 92/65/EEC after 31 August 2010 and dispatched from an approved semen collection centre of origin of the semen

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin / ISO code / I.9. Region of origin / Code		I.10. Country of destination / ISO code / I.11. Region of destination / Code	
I.12. Place of origin Semen centre ☐ Name / Approval number Address Postal code		I.13. Place of destination Semen centre ☐ Holding ☐ Name / Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:		I.17.	
I.18. Description of commodity		I.19. Commodity code (CN code) **05 11 99 85**	
			I.20. Quantity

I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages
I.23. Seal/Container No	I.24. Type of packaging
I.25. Commodities certified for: Artificial reproduction ☐	
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code
I.28. Export ☐ Third country ISO code Exit point Code	I.29.
I.30.	

I.31. Identification of the commodities

Species (Scientific name)	Breed	Donor identity	Date of collection	Approval number of the centre	Quantity

European Union **Ovine and caprine semen — Part A**

II.	Health information	II.a. Certificate reference No	II.b.

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

II.1. The semen described above:

II.1.1. was collected, processed and stored in a semen collection centre ([2]) approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;

II.1.2. comes from donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.1.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III(I) of Annex D to Directive 92/65/EEC;

([1]) *either* [II.1.4. was collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or controlled risk of classical scrapie according to point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3(c)(iv) of that Section;]

([1]) *or* [II.1.4. was collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which has/have complied for the last three years before the collection with the requirements laid down in points 1.3(a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3(c)(iv) of that Section;]

([1]) *or* [II.1.4. was collected from animals which have been kept continuously since birth in a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie;]

([1]) *or* [II.1.4. was collected from ovine animals of the ARR/ARR prion protein genotype;]

II.1.5. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

([1]) *either* [II.2. No antibiotics or no mixture of antibiotics were added to the semen.]

([1]) *or* [II.2. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than ([3]):

.....]

Notes

Part I:

Box I.12: *Place of origin* shall correspond to the semen collection centre of origin of the semen.

Box I.13: *Place of destination* shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: *Donor identity* shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected.

European Union **Ovine and caprine semen — Part A**

II. Health information	II.a. Certificate reference No	II.b.

Part II:

([1]) Delete as appropriate.

([2]) Only approved semen collection centres listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animals/live_animals/approved-establishments/index_en.htm].

([3]) Insert names and concentrations.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp: ’

ANNEX III

In Part 2 of Annex II to Decision 2010/472/EU, Section A is replaced by the following:

'Section A

Model 1 — Health certificate for semen dispatched from an approved semen collection centre of origin of the semen

COUNTRY: **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.	I.2. Certificate reference No / I.2.a. I.3. Central competent authority I.4. Local competent authority
I.5. Consignee Name Address Postal code Tel.	I.6. Person responsible for the load in EU Name Address Postal code Tel.
I.7. Country of origin / ISO code / I.8. Region of origin / Code	I.9. Country of destination / ISO code / I.10. Region of destination / Code
I.11. Place of origin Name / Approval number Address Name / Approval number Address Name / Approval number Address	I.12. Place of destination Name Address Postal code
I.13. Place of loading	I.14. Date of departure
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	I.16. Entry BIP in EU I.17.
I.18. Description of commodity	I.19. Commodity code (HS code) **05 11 99 85** I.20. Quantity

I.21.	I.22. Number of packages
I.23. Seal/Container No	I.24.
I.25. Commodities certified for: Artificial reproduction ☐	
I.26. For transit through EU to third country ☐ Third country ISO code	I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Breed	Donor identity	Date of collection	Approval number of the centre	Quantity

COUNTRY **Ovine and caprine semen — Section A**

II. **Health information**	II.a. **Certificate reference No**	II.b.

Part II: Certification

I, the undersigned, official veterinarian, hereby certify that:

II.1. The exporting country .....
(*name of exporting country*) ([2])

II.1.1. has been free from rinderpest, peste des petits ruminants, sheep and goat pox, contagious caprine pleuropneumonia and Rift Valley fever during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to the Union and no vaccination against these diseases took place during that period;

II.1.2. has been free from foot-and-mouth disease during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to the Union and no vaccination against this disease took place during that period.

II.2. The semen collection centre described in Box I.11 and at which the semen to be exported was collected and stored:

II.2.1. meets the conditions for the approval of semen collection centres laid down in Chapter I(I)(1) of Annex D to Directive 92/65/EEC;

II.2.2. is operated and supervised in accordance with the conditions applicable to semen collection centres and storage centres laid down in Chapter I(II)(1) of Annex D to Directive 92/65/EEC.

II.3. The ovine ([1])/caprine ([1]) animals standing at the semen collection centre:

II.3.1. prior to their stay in the quarantine accommodation described in point II.3.3,

([1]) ([4]) *either* [II.3.1.1. originate from the territory described in Box I.8, which has been recognised as officially brucellosis (*B. melitensis*)-free,]

([1]) *or* [II.3.1.1. have belonged to a holding which has obtained and maintained its officially brucellosis (*B. melitensis*)-free status in accordance with Directive 91/68/EEC,]

([1]) *or* [II.3.1.1. originate from a holding, where in respect of brucellosis (*B. melitensis*) all susceptible animals have been free from clinical or any signs of this disease for the last 12 months, none of the ovine and caprine animals have been vaccinated against this disease, save those vaccinated with Rev. 1 vaccine more than two years ago, and all ovine and caprine animals over six months of age have been subjected to at least two tests ([3]), carried out with negative results on samples taken on ..... (*date*) and on ..... (*date*) at least six months apart, the latter being within 30 days before entry into the quarantine accommodation,]

and have not been kept previously in a holding of a lower status;

II.3.1.2. have been kept continuously for at least 60 days on a holding where no case of contagious epididymitis (*Brucella ovis*) has been diagnosed in the last 12 months,

([1]) *and* [they are animals of the ovine species and have undergone during the 60 days prior to their stay in the quarantine accommodation described in point II.3.3 a complement fixation test, or any other test with an equivalent documented sensitivity and specificity, to detect contagious epididymitis with result of less than 50 ICFTU/ml;]

II.3.1.3. to the best of my knowledge do not come from holdings and have not been in contact with animals of a holding, in which, based on the official notification system and according to the written declaration made by the owner, any of the following diseases has been clinically detected within the periods referred to in points (a) to (d) prior to their stay in the quarantine accommodation described in point II.3.3.

(a) contagious agalactia of sheep or goats (*Mycoplasma agalactiae, Mycoplasma capricolum, Mycoplasma mycoides var. mycoides* “large colony”), within the last six months,

(b) paratuberculosis and caseous lymphadenitis, within the last 12 months,

(c) pulmonary adenomatosis, within the last three years;

COUNTRY **Ovine and caprine semen — Section A**

II.	**Health information**	II.a. **Certificate reference No**	II.b.

(1) *either* [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last three years;]

(1) *or* [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last 12 months, and all the infected animals were slaughtered and remaining animals subsequently reacted negatively to two tests carried out at least six months apart;]

II.3.2. have undergone the following tests carried out on a blood sample collected within the 28 days preceding the commencement of the period of quarantine specified in point II.3.3 for:

— brucellosis (*B. melitensis*), with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

— contagious epididymitis (*Brucella. ovis*), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

— border disease in accordance with point 1.4(c) of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.3. have satisfied the quarantine isolation period of at least 28 days in a quarantine accommodation specifically approved for the purpose by the competent authority and during that period:

II.3.3.1. only animals of at least the same health status were present in the quarantine accommodation;

II.3.3.2. the animals have undergone the following tests, carried out by the laboratory approved by the competent authority of the exporting country on samples taken not earlier than 21 days after the animals were admitted to the quarantine accommodation, for:

— brucellosis (*B. melitensis*) with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

— contagious epididymitis (*Brucella ovis*), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

— border disease in accordance with point 1.6 of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.4. have undergone at least once a year the routine tests for:

— brucellosis (*B. melitensis*) with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

— contagious epididymitis (*Brucella ovis),* in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

— border disease in accordance with point 5(c) of Chapter II(II) of Annex D to Directive 92/65/EEC.

II.4. The semen to be exported was obtained from donor rams (1)/bucks (1) which:

II.4.1. were admitted to the approved semen collection centre with the express permission of the centre veterinarian.

II.4.2. show no clinical signs of disease on the day of admission to the approved semen collection centre and on the day the semen was collected;

(1) *either* [II.4.3. have not been vaccinated against foot-and-mouth disease during the 12 months prior to collection of the semen;]

COUNTRY **Ovine and caprine semen — Section A**

II.	**Health information**	II.a. **Certificate reference No**	II.b.

(1) *or*	[II.4.3.	have been vaccinated against foot-and-mouth disease at least 30 days prior to the collection, and 5 % (with a minimum of five straws) of each collection have been submitted to a virus isolation test for foot-and-mouth disease with negative results;]
	II.4.4.	have been kept at an approved semen collection centre for a continuous period of at least 30 days immediately prior to collection of the semen, in the case of collections of fresh semen;
	II.4.5.	have not served naturally after their entry to the quarantine accommodation described in point II.3.3 and up to and including the day of semen collection;
	II.4.6.	have been kept at approved semen collection centres:
	II.4.6.1.	which have been free from foot-and-mouth disease for at least three months prior to collection of the semen and 30 days after collection or, in the case of fresh semen, until the date of dispatch, and which are situated in the centre of an area of 10 kilometres radius in which there has been no case of foot-and-mouth disease for at least 30 days prior to collection of the semen;
	II.4.6.2.	which have been free, during the period commencing 30 days prior to collection and ending 30 days after collection of the semen or, in the case of fresh semen, until the date of dispatch, from brucellosis (*B. melitensis*), contagious epididymitis (*Brucella. ovis*), anthrax and rabies;
(1) *either*	[II.4.7.	have remained in the exporting country for at least the past six months prior to collection of the semen to be exported;]
(1) *or*	[II.4.7.	during the last six months prior to collection of the semen they complied with the animal health conditions applying to donors of the semen which is intended for export to the Union and they have been imported into the exporting country at least 30 days prior to collection of the semen from ..... (2);]
(1) *either*	[II.4.8.	were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]
(1) *or*	[II.4.8.	were kept during a bluetongue virus seasonally free period in a seasonally free zone for at least 60 days prior to, and during collection of the semen;]
(1) *or*	[II.4.8.	were kept in a vector-protected establishment for at least 60 days prior to, and during collection of the semen;]
(1) *or*	[II.4.8.	were subjected to a serological test for the detection of antibody to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, on blood samples taken at least every 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen;]
(1) *or*	[II.4.8.	were subjected to an agent identification test for bluetongue virus, carried out in accordance with the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals with negative results on blood samples taken at commencement and final collection for this consignment of semen and at least every 7 days (virus isolation test) or at least every 28 days (PCR test) during collection for this consignment of semen;]
(1) (5) *either*	[II.4.9.	were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]
(1) *or*	[II.4.9.	were resident in the exporting country in which according to official findings the following serotypes of epizootic haemorrhagic disease (EHD) exist: ..... and were subjected with negative results in each case to:
	(1) *either*	[a serological test (6) for the detection of antibody to the EHDV group carried out in an approved laboratory on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days after the final collection for this consignment of semen.]]

COUNTRY **Ovine and caprine semen — Section A**

II.	**Health information**	II.a. **Certificate reference No**	II.b.

(1) *or* [a serological test (6) for the detection of antibody to the EHDV group, carried out in an approved laboratory on samples of blood taken at intervals of not more than 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen.]]

(1) *or* [an agent identification test (6) carried out in an approved laboratory on samples of blood taken at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days (PCR test) during collection for this consignment of semen.]]

II.4.10. have been kept continuously since birth in a country where the following conditions are fulfilled:

II.4.10.1. classical scrapie is compulsorily notifiable;

II.4.10.2. an awareness, surveillance and monitoring system is in place;

II.4.10.3. ovine and caprine animals affected with classical scrapie are killed and completely destroyed;

II.4.10.4. the feeding to ovine and caprine animals of meat-and-bone meal, or greaves of ruminant origin, as defined in the OIE Terrestrial Animal Health Code, has been banned and effectively enforced in the whole country for a period of at least the last seven years;

(1) *either* [II.4.11. have been kept continuously for a period of the last three years preceding the date of the collection of the semen to be exported in a holding or holdings which has/have fulfilled during that period all the requirements set out in points 1.3(a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3(c)(iv) of that Section;]

(1) *or* [II.4.11. are ovine animals of ARR/ARR prion protein genotype.]

II.5. The semen to be exported:

II.5.1. was collected after the date on which the semen collection centre was approved by the competent authority of the exporting country;

II.5.2. was collected, processed, preserved, stored and transported in accordance with the requirements applicable to semen laid down in Chapter III(I) of Annex D to Directive 92/65/EEC;

II.5.3. was sent to the place of loading in a sealed container in accordance with the requirements for semen to be subject to trade laid down in point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

(1) *either* [II.6. No antibiotics were added to the semen.]

(1) *or* [II.6. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than (7):

.....]

Notes

Part I:

Box I.6: *Person responsible for the load in EU*: this box is to be filled in only if it is a certificate for transit commodity.

Box I.11: *Place of origin* shall correspond to the approved semen collection centre in which the semen was collected and listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animals/semen/ovine_caprine/index_en.htm

Box I.22: Number of packages shall correspond to the number of containers.

COUNTRY **Ovine and caprine semen — Section A**

II. **Health information**	II.a. **Certificate reference No**	II.b.

Box I.23: Identification of container and seal number shall be indicated.

Box I.26: Fill in according to whether it is a transit or an import certificate.

Box I.27: Fill in according to whether it is a transit or an import certificate.

Box I.28: *Species*: select amongst "*Ovis aries*" or "*Capra hircus*" as appropriate.

Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd.mm.yyyy.

Approval number of the centre shall correspond to the approval number of the semen collection centre indicated in Box I.11.

Part II:

(1) Delete as necessary.

(2) Only third countries listed in Annex I to Decision 2010/472/EU.

(3) Tests shall be carried out in accordance with Annex C to Directive 91/68/EEC.

(4) Only for the territory appearing with the entry "V" in column 6 of Part 1 of Annex I to Commission Regulation (EU) No 206/2010 (OJ L 73, 20.3.2010, p. 1.).

(5) See remarks for exporting country concerned in Annex I to Decision 2010/472/EU.

(6) Standards for EHD virus diagnostic tests are described in Chapter 2.1.7 of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

(7) Insert names and concentrations.

— The signature and the stamp must be in a different colour to that of the printing.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp: '

COMMISSION DECISION (EU) 2016/2003

of 14 November 2016

amending Decisions 2009/300/EC, 2011/263/EU, 2011/264/EU, 2011/382/EU, 2011/383/EU, 2012/720/EU and 2012/721/EU in order to prolong the period of validity of the ecological criteria for the award of the EU Ecolabel to certain products

(notified under document C(2016) 7218)

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 66/2010 of the European Parliament and of the Council of 25 November 2009 on the EU Ecolabel ([1]), and in particular Article 8(3)(c) thereof,

After consulting the European Union Eco-Labelling Board,

Whereas:

(1) Commission Decision 2009/300/EC ([2]) expires on 31 December 2016.

(2) Commission Decision 2011/263/EU ([3]) expires on 31 December 2016.

(3) Commission Decision 2011/264/EU ([4]) expires on 31 December 2016.

(4) Commission Decision 2011/382/EU ([5]) expires on 31 December 2016.

(5) Commission Decision 2011/383/EU ([6]) expires on 31 December 2016.

(6) Commission Decision 2012/720/EU ([7]) expires on 14 November 2016.

(7) Commission Decision 2012/721/EU ([8]) expires on 14 November 2016.

(8) An assessment has been carried out confirming the relevance and appropriateness of the current ecological criteria, as well as of the related assessment and verification requirements, established by Decisions 2009/300/EC, 2011/263/EU, 2011/264/EU, 2011/382/EU, 2011/383/EU, 2012/720/EU and 2012/721/EU. As a new revision of the current ecological criteria and the related assessment and verification requirements set out in those Decisions has still not been finalised, it is appropriate to prolong the periods of validity of those ecological criteria and those related assessment and verification requirements until 31 December 2017. Decisions 2009/300/EC, 2011/263/EU, 2011/264/EU, 2011/382/EU, 2011/383/EU, 2012/720/EU and 2012/721/EU should therefore be amended accordingly.

([1]) OJ L 27, 30.1.2010, p. 1.
([2]) Commission Decision 2009/300/EC of 12 March 2009 establishing the revised ecological criteria for the award of the Community Eco-label to televisions (OJ L 82, 28.3.2009, p. 3).
([3]) Commission Decision 2011/263/EU of 28 April 2011 on establishing the ecological criteria for the award of the EU Ecolabel to detergents for dishwashers (OJ L 111, 30.4.2011, p. 22).
([4]) Commission Decision 2011/264/EU of 28 April 2011 on establishing the ecological criteria for the award of the EU Ecolabel for laundry detergents (OJ L 111, 30.4.2011, p. 34).
([5]) Commission Decision 2011/382/EU of 24 June 2011 on establishing the ecological criteria for the award of the EU Ecolabel to hand dishwashing detergents (OJ L 169, 29.6.2011, p. 40).
([6]) Commission Decision 2011/383/EU of 28 June 2011 on establishing the ecological criteria for the award of the EU Ecolabel to all-purpose cleaners and sanitary cleaners (OJ L 169, 29.6.2011, p. 52).
([7]) Commission Decision 2012/720/EU of 14 November 2012 establishing the ecological criteria for the award of the EU Ecolabel for Industrial and Institutional Automatic Dishwasher Detergents (OJ L 326, 24.11.2012, p. 25).
([8]) Commission Decision 2012/721/EU of 14 November 2012 establishing the ecological criteria for the award of the EU Ecolabel for Industrial and Institutional Laundry Detergents (OJ L 326, 24.11.2012, p. 38).

(9) The measures provided for in this Decision are in accordance with the opinion of the Committee set up by Article 16 of Regulation (EC) No 66/2010,

HAS ADOPTED THIS DECISION:

Article 1

Article 3 of Decision 2009/300/EC is replaced by the following:

'*Article 3*

The ecological criteria for the product group "televisions" and the related assessment and verification requirements, shall be valid until 31 December 2017.'

Article 2

Article 4 of Decision 2011/263/EU is replaced by the following:

'*Article 4*

The ecological criteria for the product group "detergents for dishwashers" and the related assessment and verification requirements shall be valid until 31 December 2017.'

Article 3

Article 4 of Decision 2011/264/EU is replaced by the following:

'*Article 4*

The ecological criteria for the product group "laundry detergents" and the related assessment and verification requirements shall be valid until 31 December 2017.'

Article 4

Article 4 of Decision 2011/382/EU is replaced by the following:

'*Article 4*

The ecological criteria for the product group "hand dishwashing detergents" and the related assessment and verification requirements shall be valid until 31 December 2017.'

Article 5

Article 4 of Decision 2011/383/EU is replaced by the following:

'*Article 4*

The ecological criteria for the product group "all-purpose cleaners and sanitary cleaners" and the related assessment and verification requirements, shall be valid until 31 December 2017.'

Article 6

Article 3 of Decision 2012/720/EU is replaced by the following:

'*Article 3*

The ecological criteria for the product group "industrial and institutional automatic dishwasher detergents" and the related assessment and verification requirements shall be valid until 31 December 2017.'

Article 7

Article 3 of Decision 2012/721/EU is replaced by the following:

'*Article 3*

The ecological criteria for the product group "industrial and institutional laundry detergents" and the related assessment and verification requirements shall be valid until 31 December 2017.'

Article 8

This Decision is addressed to the Member States.

Done at Brussels, 14 November 2016.

For the Commission
Karmenu VELLA
Member of the Commission

COMMISSION IMPLEMENTING DECISION (EU) 2016/2004

of 14 November 2016

amending Implementing Decision 2013/780/EU providing for a derogation from Article 13(1)(ii) of Council Directive 2000/29/EC in respect of bark-free sawn wood of *Quercus* L., *Platanus* L. and *Acer saccharum* Marsh. originating in the United States of America

(notified under document C(2016) 7181)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 2000/29/EC of 8 May 2000 on protective measures against the introduction into the Community of organisms harmful to plants or plant products and against their spread within the Community ([1]), and in particular the second indent of Article 15(1) thereof,

Whereas:

(1) Directive 2000/29/EC provides for protective measures against the introduction into the Union from third countries of organisms that are harmful to plants or plant products. Article 1 of Commission Implementing Decision 2013/780/EU ([2]) provides for a derogation from the protective measures under Directive 2000/29/EC and authorises Member States to allow for the introduction into their territory of bark-free sawn wood of *Quercus* L., *Platanus* L. and *Acer saccharum* Marsh., originating in the United States, without being accompanied by a phytosanitary certificate, until 30 November 2016, provided that such wood complies with the conditions set out in the Annex to that Implementing Decision.

(2) Commission Implementing Decision (EU) 2015/893 ([3]) lays down the requirements for the introduction into the Union of bark-free sawn wood of *Platanus* L. and *Acer* spp. originating in third countries. Those requirements are considered necessary for the phytosanitary protection of Union territory from *Anoplophora glabripennis* (Motschulsky) and no derogation from them should be considered justified. Those species of bark-free sawn wood should therefore no longer be subject to the derogation provided for in Implementing Decision 2013/780/EU.

(3) In light of the information submitted periodically by Member States to the Commission, it may be concluded that the application of the specific conditions laid down in the Annex to Implementing Decision 2013/780/EU is sufficient to prevent the introduction of harmful organisms into the Union. Member States should continue to apply those conditions with regard to bark-free sawn wood of *Quercus* L. originating in the United States. The assessment of the technical information submitted by the United States shows that the Kiln Drying Sawn Hardwood Lumber Certification Program, referred to in Article 2(3) of Implementing Decision 2013/780/EU, is functioning effectively.

(4) Therefore the authorisation for the derogation in respect of bark-free sawn wood of *Quercus* L. originating in the United States should be extended until 31 December 2026 in order to avoid any unnecessary disruptions of trade with regard to that wood.

(5) Implementing Decision 2013/780/EU should therefore be amended accordingly.

(6) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

([1]) OJ L 169, 10.7.2000, p. 1.

([2]) Commission Implementing Decision 2013/780/EU of 18 December 2013 providing for a derogation from Article 13(1)(ii) of Council Directive 2000/29/EC in respect of bark-free sawn wood of *Quercus* L., *Platanus* L. and *Acer saccharum* Marsh. originating in the United States of America (OJ L 346, 20.12.2013, p. 61).

([3]) Commission Implementing Decision (EU) 2015/893 of 9 June 2015 as regards measures to prevent the introduction into and the spread within the Union of *Anoplophora glabripennis* (Motschulsky) (OJ L 146, 11.6.2015, p. 16).

HAS ADOPTED THIS DECISION:

Article 1

Amendment of Implementing Decision 2013/780/EU

Implementing Decision 2013/780/EU is amended as follows:

(1) Article 1 is replaced by the following:

'By way of derogation from Article 13(1)(ii) of Directive 2000/29/EC, Member States shall be authorised to allow for the introduction into their territory of bark-free sawn wood of *Quercus* L. originating in the United and covered by one of the CN codes and the descriptions set out in Section I(6) of Part B of Annex V to that Directive without being accompanied by a phytosanitary certificate, provided that such wood complies with the conditions set out in the Annex to this Decision.';

(2) in Article 3, the date '30 November 2016' is replaced by the date '31 December 2026'.

Article 2

Addressees

This Decision is addressed to the Member States.

Done at Brussels, 14 November 2016.

For the Commission
Vytenis ANDRIUKAITIS
Member of the Commission

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