

---

**Changes to legislation:** *Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) [View outstanding changes](#)*

---

Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU)

**Changes to legislation:** Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

## ANNEX

Annex E to Directive 92/65/EEC is amended as follows:

- (1) Part 1 is replaced by the following:

## Part 1 —

**Health certificate for trade in animals from holdings (ungulates, birds vaccinated against avian influenza, lagomorphs, dogs, cats and ferrets) 92/65 EI**

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.				
	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 <input type="checkbox"/>  Name Address  Postal code			I.13. Place of destination Holding <input type="checkbox"/> Establishment <input type="checkbox"/> Approved body <input type="checkbox"/>  Name Address  Postal code				
	I.14. Place of loading Postal code			I.15. Date and time of departure				
	I.16. Means of transport  Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification			I.17. Transporter  Name Address  Postal code				
	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 <input type="checkbox"/> Production <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Slaughter <input type="checkbox"/> Pets <input type="checkbox"/> Approved body <input type="checkbox"/>								
I.26. Transit through third country <input type="checkbox"/> Third country Exit point Entry point			I.27. Transit through Member States <input type="checkbox"/> Member State Member State Member State					
I.28. Export <input type="checkbox"/> Third country Exit point			I.29. Estimated journey time					
I.30. Route plan Yes <input type="checkbox"/> No <input type="checkbox"/>								
I.31. Identification of the commodities								
Species (scientific name)		Identification system	Identification number	Sex	Age	Quantity		

**Changes to legislation:** Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

EUROPEAN UNION		92/65 EI Animals from holdings (ungulates, birds <sup>(2)</sup> , lagomorphs, dogs, cats and ferrets)	
		II.a. Certificate reference number	II.b.
Part II: Certification	II.	<b>Health information</b>	
		I, the undersigned official veterinarian <sup>(1)</sup> /veterinarian responsible for the establishment of origin and approved by the competent authority <sup>(1)</sup> certify that:	
	<sup>(1)</sup> either	II.1.	at the time of inspection the above animals were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005.]
	<sup>(1)</sup> or	II.1.	at the time of inspection the dogs <sup>(1)</sup> /cats <sup>(1)</sup> /ferrets <sup>(1)</sup> to be moved for non-commercial purposes in accordance with Commission Regulation (EU) No 388/2010 were fit to travel.]
	<sup>(1)</sup> either	II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the ruminant(s) <sup>(1)</sup> /suidae <sup>(1)</sup> other than that/those covered by Council Directive 64/432/EEC <sup>(1)</sup> or Council Directive 91/68/EEC <sup>(1)</sup> :
			(a) belong(s) to the species .....
			(b) at the time of examination, do(does) not show any clinical sign of any disease to which it/they is/are susceptible;
			(c) come(s) from an officially tuberculosis-free <sup>(1)</sup> /officially brucellosis-free <sup>(1)</sup> or brucellosis-free <sup>(1)</sup> herd <sup>(1)</sup> /holding <sup>(1)</sup> not subject to swine fever restrictions or from a holding where it/they was/were subjected with negative results to the tests laid down in Article 6(2)(b) <sup>(1)</sup> /the test laid down in Article 6(3)(d) <sup>(1)</sup> of Council Directive 92/65/EEC.]
	<sup>(1)</sup> <sup>(2)</sup> or	II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the birds other than those referred to in Council Directive 2009/158/EC:
			(a) conform to Decision 2007/598/EC and were vaccinated against avian influenza on ..... (date) with vaccine ..... (name) and come from a holding on which vaccination against avian influenza was carried out during the past 12 months;
			(b) satisfy the requirements of Article 7 of Council Directive 92/65/EEC;
			(c) at the time of examination do not show any clinical sign of any disease to which they are susceptible.]
	<sup>(1)</sup> or	II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the lagomorphs:
			(a) satisfy the requirements of Article 9 of Council Directive 92/65/EEC;
		(b) at the time of examination do not show any clinical signs of disease to which they are susceptible.]	
<sup>(1)</sup> or	II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the dogs underwent a clinical examination, within 24 hours before dispatch, by a veterinarian authorised by the competent authority, and that examination showed the animals to be in good health, and satisfy, in accordance with Article 10(2) of Council Directive 92/65/EEC, the requirements laid down in Article 5 of Regulation (EC) No 998/2003 of the European Parliament and of the Council,	
and	<sup>(1)</sup> either	[have not been treated against <i>Echinococcus multilocularis</i> .]	
	<sup>(1)</sup> or	[have been treated against <i>Echinococcus multilocularis</i> in accordance with Commission Delegated Regulation (EU) No 1152/2011.]	
<sup>(1)</sup> or	II.2.	the conditions of Article 4 of Council Directive 92/65/EEC are fulfilled and the cats <sup>(1)</sup> /ferrets <sup>(1)</sup> underwent a clinical examination, within 24 hours before dispatch, by a veterinarian authorised by the competent authority, and that examination showed the animals to be in good health, and satisfy, in accordance with Article 10(2) of Council Directive 92/65/EEC, the requirements laid down in Article 5 of Regulation (EC) No 998/2003 of the European Parliament and of the Council.]	
<sup>(1)</sup> or	II.2.	the consignment of more than five dogs to be moved for non-commercial purposes in accordance with Commission Regulation (EU) No 388/2010 underwent a clinical examination, within 24 hours before dispatch, by a veterinarian authorised by the competent authority, and that examination showed the animals to be in good health, and the animals satisfy, in accordance with Article 10(2) of Council Directive 92/65/EEC, the requirements laid down in Article 5 of Regulation (EC) No 998/2003 of the European Parliament and of the Council,	
and	<sup>(1)</sup> either	[their scheduled destination indicated in Box I.10, or in Box I.11 where regionalisation is applied, does not require a treatment against <i>Echinococcus multilocularis</i> in accordance with Commission Delegated Regulation (EU) No 1152/2011.]	
	<sup>(1)</sup> or	[they have been treated against <i>Echinococcus multilocularis</i> in accordance with Article 7 of Commission Delegated Regulation (EU) No 1152/2011.]	

**Changes to legislation:** Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

EUROPEAN UNION		92/65 E1 Animals from holdings (ungulates, birds <sup>(2)</sup> , lagomorphs, dogs, cats and ferrets)	
II.	Health information	II.a. Certificate reference number	II.b.
( <sup>1</sup> ) or	II.2. the consignment of more than five cats ( <sup>1</sup> )/ferrets ( <sup>1</sup> ) to be moved for non-commercial purposes in accordance with Commission Regulation (EU) No 388/2010 underwent a clinical examination, within 24 hours before dispatch, by a veterinarian authorised by the competent authority, and that examination showed the animals to be in good health and the animals satisfy, in accordance with Article 10(2) of Council Directive 92/65/EEC, the requirements laid down in Article 5 of Regulation (EC) No 998/2003 of the European Parliament and of the Council;]		
II.3.	The additional guarantees regarding diseases listed in Annex B <sup>(3)</sup> to Council Directive 92/65/EEC are as follows: ( <sup>1</sup> )		
	Disease	Decision	
	Disease	Decision	
	Disease	Decision	
II.4.	This certificate is valid until ..... ( <sup>4</sup> )		
<b>Notes</b>			
<b>Part I:</b>			
— Box references I.1 to I.4, I.8, I.20, I.25 and I.31: Required for non-commercial movement of more than five dogs, cats and ferrets.			
— Box reference I.6: No(s) of accompanying documents: CITES, if applicable.			
— Box reference I.19: Use the appropriate HS code: 01.06.19, 01.06.31, 01.06.32, 01.06.39.			
— Box reference I.25: Indicate "Pets" only when more than five dogs, cats or ferrets are to be certified for strictly non-commercial movements.			
— Box reference I.31: Identification system: individual identification must be used wherever possible but in the case of small animals, batch identification may be used.			
<b>Part II:</b>			
( <sup>1</sup> ) Delete as necessary.			
( <sup>2</sup> ) Certification requirements only apply to birds that have been vaccinated against avian influenza under a preventive vaccination plan approved by Commission Decision 2007/598/EC.			
( <sup>3</sup> ) As requested by a Member State benefiting from additional guarantees under Union legislation.			
( <sup>4</sup> ) The period of validity of this certificate is 10 days from the date of issue, except for dogs, cats and ferrets moved for non-commercial purposes in accordance with Commission Regulation (EU) No 388/2010, in which case the certificate is valid for a period of four months or until the date of expiry of the anti-rabies vaccination shown in Section IV of the passport, whichever is earlier.			
— 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:		

(2) Part 3 is replaced by the following:

**Changes to legislation:** Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Part 3 —

**Health certificate for trade in animals, semen, ova and embryos from approved bodies, institutes or centres 92/65 EIII**

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.				
	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 Approved body <input type="checkbox"/>  Name Address Postal code			I.13. Place of destination Approved body <input type="checkbox"/>  Name Address Postal code				
	I.14. Place of loading Postal code			I.15. Date and time of departure				
	I.16. Means of transport  Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification			I.17. Transporter  Name Address Postal code				
	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:  Approved body <input type="checkbox"/>							
	I.26. Transit through third country <input type="checkbox"/> Third country Exit point Entry point			ISO code Code BIP No	I.27. Transit through Member States <input type="checkbox"/> Member State Member State Member State			ISO code ISO code ISO code
I.28. Export <input type="checkbox"/> Third country Exit point			ISO code Code	I.29. Estimated journey time				
I.30. Route plan Yes <input type="checkbox"/> No <input type="checkbox"/>								
I.31. Identification of the commodities  Species (scientific name)      Identification system      Identification number      Sex      Age      Quantity								

**Changes to legislation:** Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

EUROPEAN UNION		92/65 EIII Animals from approved bodies, institutes or centres		
Part II: Certification	II. <b>Health information</b>	II.a. Certificate reference number	II.b.	
	I, the undersigned official veterinarian <sup>(1)</sup> /veterinarian responsible for the establishment of origin and approved by the competent authority <sup>(1)</sup> certify that:			
	II.1.	The body, institute or centre of origin is approved in accordance with Annex C to Council Directive 92/65/EEC for the purpose of trade in the animals, semen, ova or embryos described in Box I.18.		
	II.2.	The animals <sup>(1)</sup> /donor animals <sup>(1)</sup> described in this certificate have been examined today <sup>(1)</sup> /on the day of collection <sup>(1)</sup> and found to be healthy and free of clinical signs of infectious diseases including those listed in Annex A to Directive 92/65/EEC and are not subject to any official restrictions and remained in this body, institute or centre either since birth or for the following time ..... (months or years).		
	II.3.	At the time of inspection, the above animals were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005 and IATA requirements and/or CITES guidelines for transport, where applicable.		
	II.4.	The additional guarantees regarding diseases listed in Annex B <sup>(2)</sup> to Council Directive 92/65/EEC are as follows: <sup>(1)</sup>		
		Disease	Decision	
		Disease	Decision	
		Disease	Decision	
	II.5.	Birds conforming to Decision 2007/598/EC were vaccinated against avian influenza on ..... (date) with vaccine ..... (name) and come from an approved body, institute or centre of origin on which vaccination against avian influenza was carried out during the past 12 months.] <sup>(1)</sup>		
<b>Notes</b>				
<b>Part I:</b>				
— Box reference I.6: No(s) of accompanying documents: CITES, if applicable.				
— Box reference I.19: Use the appropriate HS code: 01.06.11, 01.06.19, 01.06.31, 01.06.32, 01.06.39, 05.11.99.85.				
— Box reference I.31: <i>Identification system:</i> individual identification must be used wherever possible but in the case of small animals, batch identification may be used.				
In the case of semen, ova and embryos it shall correspond to the <i>donor identity</i> and the <i>date of collection</i> and shall be indicated in the following format: official identification of the animal/dd/mm/yyyy.				
<i>Age</i> and <i>sex:</i> to be completed only in the case of live animals, if appropriate.				
<i>Quantity:</i> in the case of semen, ova and embryos the number of straws, ampoules or other packaging express as units should be indicated.				
<b>Part II:</b>				
<sup>(1)</sup> Delete as necessary.				
<sup>(2)</sup> As requested by a Member State benefiting from additional guarantees under Union legislation.				
— 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:			

**Changes to legislation:**

Commission Implementing Decision of 17 February 2012 amending Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and animals, semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU) is up to date with all changes known to be in force on or before 01 November 2023. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations.

[View outstanding changes](#)

**Changes and effects yet to be applied to :**

- Decision implicit repeal by [EUR 2016/429](#) Regulation