

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

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[^{F1}ANNEX E

Textual Amendments

- F1** Substituted by [Commission Decision of 26 April 2007 amending Annex E to Council Directive 92/65/EEC to include additional health measures for the trade in live bees, and to update the health certificates models \(notified under document number C\(2007\) 1811\) \(Text with EEA relevance\) \(2007/265/EC\)](#).

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[^{F2}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
I.12. Place of origin Holding <input type="checkbox"/> Name Approval/registration number Address Postal code				I.13. Place of destination Holding <input type="checkbox"/> Establishment <input type="checkbox"/> Approved body <input type="checkbox"/> 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 <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 Approval number Address Postal code					
I.18. Description of commodity						I.19. Commodity code (CN code)			
								I.20. Quantity	

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EUROPEAN UNION		92/65 EI	Animals from holdings (ungulates, birds ⁽²⁾ , lagomorphs, dogs, cats and ferrets)
II. Health information		II.a. Certificate reference No	II.b.
I, the undersigned official veterinarian ⁽¹⁾ /veterinarian responsible for the holding of origin and approved by the competent authority ⁽¹⁾ certify that:			
Part II: Certification	II.1.	the animals described in Box I.31 comply with the conditions of Article 4 of Council Directive 92/65/EEC and at the time of inspection were fit to be transported for the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005.	
	⁽¹⁾ either II.2.	the ruminant(s) ⁽¹⁾ / <i>suidae</i> ⁽¹⁾ other than that/those covered by Council Directive 64/432/EEC ⁽¹⁾ or Council Directive 91/68/EEC ⁽¹⁾	
	(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 ⁽¹⁾ /officially brucellosis-free ⁽¹⁾ or brucellosis-free ⁽¹⁾ herd ⁽¹⁾ /holding ⁽¹⁾ 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) ⁽¹⁾ /the test laid down in Article 6(3)(d) ⁽¹⁾ of Council Directive 92/65/EEC.]	
	⁽¹⁾ ⁽²⁾ or II.2.	the birds other than those referred to in Council Directive 2009/158/EC	
	(a)	at the time of examination do not show any clinical sign of any disease to which they are susceptible;	
	(b)	satisfy the requirements of Article 7 of Council Directive 92/65/EEC;	
	(c)	conform to Commission 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.]	
	⁽¹⁾ or II.2.	the lagomorphs	
(a)	at the time of examination do not show any clinical signs of disease to which they are susceptible;		
(b)	satisfy the requirements of Article 9 of Council Directive 92/65/EEC.]		
⁽¹⁾ or II.2.	the dogs		
(a)	at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;		
(b)	are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;		
⁽¹⁾ either [(c)	were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination];		
⁽¹⁾ or [(c)	are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and		
(i)	the Member State of destination has informed the public in accordance with point (b) of Article 37(2) of Regulation (EU) No 576/2013 of the European Parliament and of the Council that it authorises the movement of such animals into its territory; and they are accompanied by		

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EUROPEAN UNION

92/65 EI Animals from holdings (ungulates, birds ⁽²⁾, lagomorphs, dogs, cats and ferrets)

II. Health information	II.a. Certificate reference No	II.b.
<p>(¹) either</p> <p>(¹) or</p> <p>(d)</p> <p>(¹) and</p>	<p>[(ii)</p> <p>[(ii)</p> <p></p> <p>[(e)</p>	<p>a declaration of the owner ⁽³⁾, attached to this certificate, stating that from birth until the time of dispatch the animals have had no contact with wild animals of species susceptible to rabies];</p> <p>their mother, on whom they still depend, and from the passport of their mother, it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council];</p> <p>are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No 577/2013;</p> <p>due to their scheduled destination ⁽⁴⁾ indicated in Box I.10, or in Box I.11 where regionalisation is applied, have been treated against <i>Echinococcus multilocularis</i> in accordance with Commission Delegated Regulation (EU) 2018/772];</p>
<p>(¹) or</p> <p>[(¹) either</p> <p>(¹) or</p> <p>(¹) either</p> <p>(¹) or</p> <p>(¹) either</p> <p>(¹) or</p> <p>(d)</p>	<p>II.2.</p> <p>the cats ⁽¹⁾/ferrets ⁽¹⁾</p> <p>[(c)</p> <p>[(c)</p> <p>(i)</p> <p>[(ii)</p> <p>[(ii)</p> <p></p> <p>(d)</p>	<p>the cats ⁽¹⁾/ferrets ⁽¹⁾</p> <p>at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;</p> <p>are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;</p> <p>were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination];</p> <p>are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and</p> <p>the Member State of destination has informed the public in accordance with point (b) of Article 37(2) of Regulation (EU) No 576/2013 of the European Parliament and of the Council that it authorises the movement of such animals into its territory; and they are accompanied by</p> <p>a declaration of the owner ⁽³⁾, attached to this certificate, stating that from birth until the time of dispatch the animals have had no contact with wild animals of species susceptible to rabies];</p> <p>their mother, on whom they still depend, and from the passport of their mother, it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council];</p> <p>are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No 577/2013.]</p>
<p>(¹) or</p> <p>(a)</p> <p>(b)</p>	<p>II.2.</p> <p>the dogs ⁽¹⁾/cats ⁽¹⁾/ferrets ⁽¹⁾ are destined for a body, institute or centre described in Box I.13 and approved in accordance with Annex C to Council Directive 92/65/EEC, and</p> <p></p> <p></p>	<p>the dogs ⁽¹⁾/cats ⁽¹⁾/ferrets ⁽¹⁾ are destined for a body, institute or centre described in Box I.13 and approved in accordance with Annex C to Council Directive 92/65/EEC, and</p> <p>at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;</p> <p>are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;</p>

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EUROPEAN UNION 92/65 EI **Animals from holdings (ungulates, birds ⁽²⁾, lagomorphs, dogs, cats and ferrets)**

II. Health information	II.a. Certificate reference No	II.b.						
<p>(c) are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No 577/2013.]</p> <p>II.3. The additional guarantees regarding diseases listed in Annex B ⁽⁵⁾ to Council Directive 92/65/EEC are as follows ⁽¹⁾:</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 60%;">Disease</td> <td>Decision</td> </tr> <tr> <td>Disease</td> <td>Decision</td> </tr> <tr> <td>Disease</td> <td>Decision</td> </tr> </table>	Disease	Decision	Disease	Decision	Disease	Decision		
Disease	Decision							
Disease	Decision							
Disease	Decision							
Notes								
Part I:								
Box I.6: No(s) of accompanying documents: CITES, if applicable.								
Box I.19: Use the appropriate CN code: 01.06.19, 01.06.31, 01.06.32, 01.06.39.								
Box 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 dogs, cats and ferrets, select passport.								
<i>Identification number:</i> in the case of dogs, cats and ferrets, indicate the alphanumeric code of the tattoo or transponder.								
<i>Passport number:</i> in the case of dogs, cats and ferrets, indicate the unique alphanumeric code of the passport.								
Part II:								
⁽¹⁾ Delete as necessary.								
⁽²⁾ 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.								
⁽³⁾ The declaration referred to in point II.2 to be attached to the certificate shall be drawn up in accordance with Annex I to Commission Implementing Regulation (EU) No 577/2013.								
⁽⁴⁾ Member States or parts thereof listed in the Annex to Commission Implementing Regulation (EU) 2018/878.								
⁽⁵⁾ 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.								
This certificate is valid for 10 days from the date of signature of the official veterinarian or of the veterinarian responsible for the holding of origin and approved by the competent authority.								
Official veterinarian								
Name (in capital letters):	Qualification and title:							
Local veterinary unit:	LVU No:							
Date:	Signature:							
Stamp:								

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Textual Amendments

- F2** Substituted by [Commission Implementing Decision \(EU\) 2019/1206](#) of 12 July 2019 amending Part 1 of Annex E to Council Directive 92/65/EEC as regards the animal health certificate for trade in dogs, cats and ferrets (notified under document C(2019) 5210) (Text with EEA relevance).

[^{F3}Part 2 —

Health certificate for trade in bees and bumble bees

92/65 EII]

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EUROPEAN UNION

Intra trade certificate

Part I: Details of consignment presented	I.1. Consignor Name		I.2. Certificate reference number	I.2.a. Local reference number:			
	Address Postal code		I.3. Central Competent Authority				
			I.4. Local Competent Authority				
	I.5. Consignee Name		I.6.				
	Address Postal code					I.7.	
	I.8. Country of origin	ISO code	I.9.		I.10. Country of destination		
	I.12. Place of origin/Place of harvest Holding <input type="checkbox"/> Other <input type="checkbox"/> Name Approval number Address Postal code			I.13. Place of origin/Place of harvest Holding <input type="checkbox"/> Other <input type="checkbox"/> 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 <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification:			I.17.			
	I.18. Animal species/product						I.19. Commodity code (CN code) 01.06.90
					I.20. Number/quantity		
I.21.			I.22. Number of packages				
I.23. Identification of container/seal number			I.24.				
I.25. Animals certified as/products certified for: Breeding <input type="checkbox"/> Transhumance <input type="checkbox"/> <input checked="" type="checkbox"/> Production (pollination) <input type="checkbox"/>							
I.26. Transit through third country <input type="checkbox"/> Third country ISO code Exit point Code Entry point BIP unit no.:			I.27. Transit through Member States <input type="checkbox"/> Member State ISO code Member State ISO code Member State ISO code				
I.28. Export <input type="checkbox"/> Third country ISO code Exit point Code			I.29.				
I.30.							
I.31. Identification of the animals Species (Scientific name) Quantity Batch number <input checked="" type="checkbox"/> Nature of commodity queens, packages of bees, nucleus colonies, colonies <input type="checkbox"/>							

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COUNTRY		92/65 EII Bees (<i>Apis mellifera</i>) and bumble bees (<i>Bombus</i> spp.)	
II. Health information		II.a. Certificate reference number	II.b.
Part II: Certification	I, the undersigned certify that:		
	II.1		
	either ⁽²⁾	[(a) the bees/bumble bees ⁽²⁾ come from an area which is not subject of the prohibition order associated with an occurrence of American foulbrood (the period of prohibition has been continued for at least 30 days following the last recorded case and the date of which all hives within a radius of three kilometres have been checked by the competent authority and all infected hives burned or treated and inspected to the satisfaction of the said competent authority);]	
	or ⁽²⁾	[(a) the bumble bees come from an environmentally isolated structure recognised by and under the supervision of the competent authority of the Member State which is free of American foulbrood and was inspected immediately prior to dispatch and all bumble bees and breeding stock show no clinical signs or suspicion of the disease;]	
	and	(b) the bees/bumble bees ⁽²⁾ come from an area of at least 100 km radius which is not the subject of any restrictions associated with the suspicion or confirmed occurrence of the small hive beetle (<i>Aethina tumida</i>) or the Tropilaelaps mite (<i>Tropilaelaps</i> spp.), and where these infestations are absent;	
	▶ ⁽¹⁾ or		
		(b) the consignment consists only of cages of queen bees each containing one single queen with a maximum of 20 accompanying attendants and comes from an area of at least 100 km radius which is not the subject of any restrictions associated with the suspicion or confirmed occurrence of the Tropilaelaps mite (<i>Tropilaelaps</i> spp.) and from an establishment that fulfils all the following requirements:	
		— it is situated at least 30 km distance from the limits of a protection zone of at least 20 km in radius around confirmed occurrence(s) of the small hive beetle, and	
		— it is situated outside of a zone restricted by protective measures established by the Union due to the occurrence of small hive beetle, and	
		— it is situated in an area where annual surveillance for the detection of small hive beetle by the competent authority is ongoing to provide a confidence level of at least 95 % of detecting small hive beetle if at least 2 % of the apiaries were infested, and	
	— it is inspected every month by the competent authority with negative results to provide a confidence level of at least 95 % of detecting small hive beetle if at least 2 % of the hives were infested, and		
	— where each cage or the whole consignment is covered by a fine mesh of maximum 2 mm pore size immediately after the visual examination for the health certification;		
	or		
		(b) the bumble bees come from an environmentally isolated structure recognised by and under the supervision of the competent authority, which is free of small hive beetle; ◀	
	and	(c) the bees/bumble bees ⁽²⁾ as well as their packaging have undergone a visual examination to detect the occurrence of the small hive beetle (<i>Aethina tumida</i>) or their eggs and larvae, or other infestations, in particular the Tropilaelaps mite (<i>Tropilaelaps</i> spp.), affecting bees.	
	II.2	the additional guarantees regarding diseases listed in Annex B ⁽¹⁾ to Directive 92/65/EEC are as follows ⁽²⁾ :	
	Disease	Decision	
	Disease	Decision	
	Disease	Decision	
Notes			
Part I:			
— Box reference I.31: Species: introduce <i>Apis mellifera</i> or <i>Bombus</i> spp. Quantity: provide the number of colonies. Batch number: provide the number of seals where applicable.			
Part II:			
⁽¹⁾ As requested by a Member State benefiting from additional guarantees under Union legislation.			
⁽²⁾ Delete as necessary.			
— The colour of the stamp and signature must be different from that of the other particulars in the certificate.			
Approved veterinarian or approved official			
Name (in capital letters):		Qualification and title:	
Date:		Signature:	
Stamp:			

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Textual Amendments

- F3** Substituted by Commission Decision of 6 May 2010 amending Parts 1 and 2 of Annex E to Council Directive 92/65/EEC as regards the model health certificates for animals from holdings and for bees and bumble bees (notified under document C(2010) 2624) (Text with EEA relevance) (2010/270/EU).

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[^{F4}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.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 Approval number				I.13. Place of destination Approved body <input type="checkbox"/> Name Address Postal code Approval number			
	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 Approval number			
	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				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								

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EUROPEAN UNION	92/65 EIII Animals from approved bodies, institutes or centres								
Part II: Certification	II. Health information	<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%;">II.a. Certificate reference number</td> <td style="width: 50%;">II.b.</td> </tr> </table>	II.a. Certificate reference number	II.b.					
	II.a. Certificate reference number	II.b.							
<p>I, the undersigned official veterinarian ⁽¹⁾/veterinarian responsible for the establishment of origin and approved by the competent authority ⁽¹⁾ certify that:</p> <p>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.</p> <p>II.2. The animals ⁽¹⁾/donor animals ⁽¹⁾ described in this certificate have been examined today ⁽¹⁾/on the day of collection ⁽¹⁾ 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 (<i>months or years</i>).</p> <p>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.</p> <p>II.4. The additional guarantees regarding diseases listed in Annex B ⁽²⁾ to Council Directive 92/65/EEC are as follows: ⁽¹⁾</p> <table style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%;">Disease</td> <td style="width: 50%;">Decision</td> </tr> <tr> <td>Disease</td> <td>Decision</td> </tr> <tr> <td>Disease</td> <td>Decision</td> </tr> </table> <p>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.] ⁽¹⁾</p> <p>Notes</p> <p>Part I:</p> <p>— Box reference I.6: No(s) of accompanying documents: CITES, if applicable.</p> <p>— 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.</p> <p>— 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.</p> <p style="padding-left: 20px;">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.</p> <p style="padding-left: 20px;"><i>Age and sex:</i> to be completed only in the case of live animals, if appropriate.</p> <p style="padding-left: 20px;"><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.</p> <p>Part II:</p> <p>⁽¹⁾ Delete as necessary.</p> <p>⁽²⁾ As requested by a Member State benefiting from additional guarantees under Union legislation.</p> <p>— The colour of the stamp and signature must be different from that of the other particulars in the certificate.</p>		Disease	Decision	Disease	Decision	Disease	Decision		
Disease	Decision								
Disease	Decision								
Disease	Decision								
<p>Official veterinarian or official inspector</p> <table style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%;">Name (in capital letters):</td> <td style="width: 50%;">Qualification and title:</td> </tr> <tr> <td>Local veterinary unit:</td> <td>LVU No:</td> </tr> <tr> <td>Date:</td> <td>Signature:</td> </tr> <tr> <td>Stamp:</td> <td></td> </tr> </table>		Name (in capital letters):	Qualification and title:	Local veterinary unit:	LVU No:	Date:	Signature:	Stamp:	
Name (in capital letters):	Qualification and title:								
Local veterinary unit:	LVU No:								
Date:	Signature:								
Stamp:									

Textual Amendments

- F4** Substituted by [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,](#)

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semen, ova and embryos from approved bodies, institutes or centres (notified under document C(2012) 860) (Text with EEA relevance) (2012/112/EU).