

**COUNTRY: Countries subject to  
transitional import arrangements (\*)**

**Health certificate to Great Britain,  
Channel Islands and Isle of Man**

<b>Part I: Details of dispatched consignment</b>	I.1. Consignor Name Address				I.2. Certificate reference number		I.2.a. UNN	
	Tel.				I.3. Central Competent Authority			
	I.5. Consignee Name Address				I.6. Person responsible for the load in Great Britain, Channel Islands and Isle of Man Name Address			
	Postal Code Tel.				Postal Code Tel.			
	I.7. Country of origin		ISO code	I.8. Region of origin		Code	I.9. Country of destination	
							I.10. Region of destination	
	I.11. Place of origin Name Address				I.12. Place of destination Name Address			
	Approval number				Postal Code			
	Name Address				Approval number			
Name Address				Approval number				
I.13. Place of loading				I.14. Date of departure				
I.15. Means of transport <input type="checkbox"/> Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/>				I.16. Entry BCP in Great Britain, Channel Islands or Isle of Man				
Identification: Documentary references:				I.17.				
I.18. Description of commodity						I.19. Commodity code (HS code) 05 11 99 85		
I.21.				I.20. Quantity		I.22. Number of packages		
I.23. Seal/Container No.						I.24.		
I.25. Commodity certified for: Artificial Reproduction <input type="checkbox"/>								
I.26. For transit through Great Britain, Channel Islands and Isle of Man to third country <input type="checkbox"/> Third country ISO code				I.27. For import or admission into Great Britain, Channel Islands and Isle of Man <input type="checkbox"/>				
I.28. Identification of the commodities								
Species (Scientific Name)	Breed	Category	Dam identity	Sire identity	Date of freezing	Approval number of the team	Quantity	

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**In vitro produced bovine embryos**

**Part II: Certification**

II. Health information	II.a. Certificate reference no	II.b. UNN
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I, the undersigned, official veterinarian of the ..... certify that:  
(exporting country) <sup>(2)</sup>

II.1. The embryos to be exported:

II.1.1. were produced in the exporting country, which according to official findings:

II.1.1.1. was free from rinderpest during the 12 months immediately prior to their production;  
(<sup>1</sup>) either [II.1.1.2. was free from foot-and-mouth disease and lumpy skin disease during the 12 months immediately prior to their production and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

(<sup>1</sup>) or [II.1.1.2. was not free from foot-and-mouth disease or lumpy skin disease during the 12 months immediately prior to their production or carried out vaccination against foot-and-mouth disease or lumpy skin disease during that period, and

- The embryos were produced without penetration of the zona pellucida,
- The embryos were stored under approved conditions for at least 30 days immediately after their production,
- The donor females come from holdings on which no animal was vaccinated against foot-and-mouth disease or lumpy skin disease during the 30 days prior to collection and no animal of a susceptible species showed clinical signs of foot-and-mouth disease or lumpy skin disease during the 30 days prior to, and at least the 30 days after, the embryos were collected.]

II.1.2. were collected by the embryo collection team <sup>(3)</sup> which:

- has been approved in accordance with Chapter I of Annex A to Directive 89/556/EEC;
- carried out the production, processing, storing and transport in accordance with Chapter II of Annex A to Directive 89/556/EEC;
- is subject to inspection by an official veterinarian at least twice a year.

II.2. The oocytes used in the production of the embryos to be exported were collected on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease in the 30 days immediately prior to their collection and until their dispatch to Great Britain, , in the case of fresh embryos, or during the 30 days after collection, in the case of embryos subject to mandatory storage for at least 30 days in accordance with point II.1.1.2.

II.3. From the time of collection of the oocytes until 30 days thereafter or, in the case of fresh embryos, until the day of their dispatch, the embryos to be exported were stored on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease.

II.4 The donors of oocytes used in the production of the embryos to be exported:

II.4.1. were located, during the 30 days immediately prior to collection of the oocytes, on premises situated in an area of at least 10-km radius on which, according to official findings, there was no occurrence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease;

II.4.2. showed no clinical signs of disease on the day of collection;

II.4.3. spent the six months immediately prior to collection within the territory of the exporting country in no more than two herds:

- which, according to official findings, were free from tuberculosis during that time,
- which, according to official findings, were free from brucellosis during that time,
- which were free from enzootic bovine leukosis or in which no bovine animal showed clinical signs of enzootic bovine leukosis during the previous three years,
- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months;

(<sup>1</sup>) either [II.4.4. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the oocytes.]

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(<sup>1</sup>) or [II.4.4. were kept during a seasonally free period or protected from the vector for at least 60 days prior to, and during, the collection of the oocytes, and the embryos were produced without penetration of the *zona pellucida*, except if the donors underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results and the embryos were stored for at least 30 days.]

(<sup>1</sup>) or [II.4.4. underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results, and the embryos were stored for at least 30 days.]

(<sup>1</sup>) or [II.4.4. underwent an agent identification test, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals on a blood sample taken on the day of collection or the day of slaughtering and giving negative results - the embryos having been produced, in the latter case, without penetration of the *zona pellucida*.]

II.5. The embryos to be exported were conceived by *in vitro* fertilisation using semen coming from semen collection or storage centres(<sup>4</sup>):

(<sup>1</sup>) either [II.5.1. approved in accordance with Article 5(1) of Directive 88/407/EEC and located in a Great Britain, and the semen complies with the requirements of Directive 88/407/EEC.]

(<sup>1</sup>) or [II.5.1. approved in accordance with Article 9(1) of Directive 88/407/EEC and located in a third country or part thereof listed in Annex 1 to Implementing Decision 2011/630/EU, and the semen complies with the requirements set out in section A of Part 1 of Annex 2 to that Decision.]

**Notes**

(\*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

**Part I:**

Box I.6: Person responsible for the load in Great Britain: 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 embryo production team from which the embryos are dispatch to Great Britain and listed in accordance with Article 8(2) of Directive 89/556/EEC

Box I.16: Do not use this box until the end of the transitional staging period

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

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 '*Bos taurus*', '*Bison bison*' or '*Bubalus bubalis*' as appropriate.  
*Category*: select '*in vivo* produced embryos'.  
*Dam identity* shall correspond to the official identification of the animal.  
*Sire identity* shall correspond to the official identification of the animal.  
*Date of freezing* shall be indicated in the following format: dd.mm.yyyy  
*Approval number of the team*: shall correspond to the embryo production team by which the embryos were produced, processed and stored; and listed in accordance with Article 8(2) of Directive 89/556/EEC

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**In vitro produced bovine embryos**

II. Health information	II.a. Certificate reference no	II.b. UNN						
<p><b>Part II:</b></p> <p>(1) Delete as appropriate.</p> <p>(2) Only third countries listed in Annex 1 to Decision 2006/168/EC.</p> <p>(3) Only embryo production teams listed in accordance with Article 8(2) of Directive 89/556/EEC</p> <p>(4) Only semen collection centres listed in accordance with Article 5(2) and Article 9(2) of Directive 88/407/EEC</p> <p>- The signature and the stamp must be in a different colour to that of the printing.</p>								
<p>Official Veterinarian</p> <table border="0"><tr><td>Name (in capital letters):</td><td>Qualification and title:</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:	Date:	Signature:	Stamp:	
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