

Part I : Details of consignment	I.1. Consignor			I.2. IMSOC Reference		
	Name			I.2.a. Local Reference		
	Address					
	Country			ISO Code		
	I.5. Consignee			I.3. Central competent authority		
	Name			I.4. Local competent authority		
	Address					
	Country			ISO Code		
	I.7. Country of origin			I.9. Country of destination		
	ISO Code			ISO Code		
	I.8. Region of origin			I.10. Region of destination		
	Code					
	I.11. Place of Dispatch			I.12. Place of destination		
	Name			Name		
	Address			Address		
Approval Number			Approval Number			
Country			Country			
ISO Code			ISO Code			
I.13. Place of Loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country						
ISO Code						
I.15. Means of Transport			I.16 Entry Point			
Mode	International transport document	Identification				
I.18. Transport conditions			I.17. Accompanying documents			
Ambient <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.19. Container No / Seal No						
I.20. Certified as						
Artificial reproduction <input type="checkbox"/> Breeding <input type="checkbox"/>						
I.21. For transit through a third country <input type="checkbox"/>			I.22. For transit through Member State(s) <input type="checkbox"/>			
Country			Country			
ISO Code			ISO Code			
EU Exit Authority			BCP code			
EU Entry Authority			BCP code			
I.24. Total quantity			I.25. Total gross weight			
I.28. Description of consignment						
1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED						
0511 Animal products not elsewhere specified or included; dead animals of Chapter 1 or 3, unfit for human consumption						
051110 Bovine semen						
05111000 Bovine semen						
Commodity	Species	Identification number	Identification mark	Nature of commodity		
Quantity	Date of collection/production		Manufacturing plant			

Part II: Certification	II. Health information		
	I, the undersigned official veterinarian, hereby certify that:		
II.1	(name of exporting country or part thereof) (2) was free from rinderpest and foot-and-mouth disease during the 12 months immediately prior to collection of the semen for export and until its date of dispatch to Great Britain and no vaccination against these diseases has taken place during the same period.		
II.2	the centre (3) described in box I.11. at which the semen to be exported was collected:		
II.2.1	meets the conditions laid down in Chapter I(1) of Annex A to Directive 88/407/EEC;		
II.2.2	is operated and supervised in accordance with the conditions laid down in Chapter II(1) of Annex A to Directive 88/407/EEC		
II.3	The centre at which the semen to be exported was collected was free from rabies, tuberculosis, Brucellosis, anthrax and contagious bovine pleuropneumonia during the 30 days prior to the date of collection of the semen to be exported and the 30 days after collection (in the case of fresh semen until the day of dispatch to Great Britain).		
II.4	The bovine animals standing at the semen collection centre:		
(8)II.4.1	come from herds which satisfy the conditions of paragraph 1(b) of Chapter I of Annex B to Directive 88/407/EEC;		
II.4.2	come from herds or were born to dams which comply with the conditions of paragraph 1(c) of chapter I of Annex B to Directive 88/407/EEC, or were tested at the age of at least 24 months in accordance with paragraph 1(c) of Chapter II of Annex B to that Directive;		
II.4.3	underwent the tests required in accordance with paragraph 1(d) of Chapter I of Annex B to Directive 88/407/EEC in the 28 days preceding the quarantine isolation period		
II.4.4	have satisfied the quarantine isolation period and testing requirements laid down in paragraph 1(e) of Chapter I of Annex B to Directive 88/407/EEC;		
II.4.5	have undergone, at least once a year, the routine tests referred to in Chapter II of Annex B to Directive 88/407/EEC.		
II.5	The semen to be exported was obtained from donor bulls which:		
II.5.1	satisfy the conditions laid down in Annex C of Directive 88/407/EEC;		
(1)either	○ [II.5.2	have remained in the exporting country for at least the last six months prior to collection of the semen to be exported;]	
(1)or	○ [II.5.2	have remained in the exporting country for at least 30 days prior to the collection of the semen since entry and they were imported from (2) during the period of less than six months prior to the collection of the semen and satisfied the animal health conditions applying to donors of the semen which is intended for export to Great Britain;]	
II.5.3	comply with at least one of the following conditions as regards bluetongue, as detailed in the table in point I.28.:		
(1)either	<input type="checkbox"/> [II.5.3.1	were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]	
(1)and/or	<input type="checkbox"/> [II.5.3.2	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)and/or	<input type="checkbox"/> [II.5.3.3	were kept in a vector-protected establishment for at least 60 days prior to, and during, collection of the semen;]	
(1)and/or	<input type="checkbox"/> [II.5.3.4	were subjected to a serological test for the detection of antibody to the bluetongue virus serogroup, carried out in accordance with the World Organisation for Animal Health (WOAH (formerly OIE)) Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, at least every 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen;]	

Part II: Certification	II. Health information	
	(1)and/or	<input type="checkbox"/> [II.5.3.5 were subjected to an agent identification test for bluetongue virus, carried out in accordance with the World Organisation for Animal Health (WOAH (formerly OIE)) 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, if carried out as polymerase chain reaction (PCR), during collection for this consignment of semen;]
	II.5.4	comply with at least one of the following conditions as regards epizootic haemorrhagic disease (EHD), as detailed in the table in point 1.28:
	(1)either	<input type="checkbox"/> [II.5.4.1 were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]
	(1)(5)and/or	<input type="checkbox"/> [II.5.4.2 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 the following tests carried out in an approved laboratory:
	(1)either	<input type="checkbox"/> [II.5.4.2.1 a serological test (4) for the detection of antibody to the EHD virus serogroup, carried out on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days following collection for this consignment of semen;]
	(1)and/or	<input type="checkbox"/> [II.5.4.2.2 a serological test (4) for the detection of antibody to the EHD virus serogroup, carried out on samples 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)and/or	<input type="checkbox"/> [II.5.4.2.3 an agent identification test (4) carried out on blood samples collected at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days, if carried out as PCR, during collection for this consignment of semen.]]
	II.6	The semen to be exported was collected after the date on which the centre was approved by the competent national authorities of the exporting country.
	II.7	The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC.
Notes		
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) and can be viewed on the UK legislation website (legislation.gov.uk).		
References to Great Britain in this certificate include Channel Islands and Isle of Man.		
Part I:		
Box reference 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 reference I.11:	Place of origin shall correspond to the semen collection centre listed in accordance with Article 9(2) of Directive 88/407/EEC and where the semen was collected.	
Box reference I.22:	Number of packages shall correspond to the number of containers.	
Box reference I.23:	Identification of container and seal number shall be indicated.	
Box reference I.26:	Fill in according to whether it is a transit or an import certificate.	

Part II: Certification	II. Health information		
	Box reference I.27: Fill in according to whether it is a transit or an import certificate.		
	Box reference I.28: Species: select amongst 'Bos taurus', 'Bison bison' or 'Bubalus bubalis' 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. Quantity shall correspond to the number of straws of semen collected on a particular date from an identified donor bull complying with particular conditions for bluetongue and EHD.		
	Part II:		
	(1) Delete as necessary.		
	(2) Only third countries or parts thereof listed in a document relating to 'bovine semen' published on gov.uk, in accordance with Decision 2011/630/EU.(9)		
	(3) Only semen collection centres listed in accordance with Article 9(2) of Directive 88/407/EEC.		
	(4) Standards for EHD virus diagnostic tests are described in the Bluetongue Chapter (2.1.3) of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.		
	(5) Compulsory for Australia, Canada and the United States.		
	(6) Referring to each straw or batch of straws indicate applicable condition (for example II.5.3.1).		
	(7) Referring to each straw or batch of straws indicate applicable condition (for example II.5.4.1 or II.5.4.2.1).		
	(8) For New Zealand, appearing with the entry 'XII' in column 6 of a document relating to 'live ungulates' published on gov.uk, in accordance with Commission Regulation (EU) No 206/2010, officially tuberculosis-free bovine herds shall be considered equivalent to officially tuberculosis-free bovine herds in Great Britain recognised based on the conditions laid down in paragraphs 1 and 2 of Annex A.I to Council Directive 64/432/EEC.(9)		
	(9) Documents relating to 'bovine semen' and 'live ungulates' published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here: EU and EFTA states approved to export animals and animal products to Great Britain - data.gov.uk Non-EU countries approved to export animals and animal products to Great Britain - data.gov.uk		
	The signature and the stamp must be in a different colour to that of the printing.		
	Certifying Officer		
	Name (in capital letters)	Qualification and title	
	Date of signature	Signature	
	Stamp		