## PART 2

Model certificate applicable from 1 January 2005 to imports and transits of stocks of semen collected, processed and stored before 31 December 2004 in conformity with the provisions of Council Directive 88/407/EEC applying until 1 July 2003, and imported after 31 December 2004 in accordance with Article 2(2) of Directive 2003/43/EC, dispatched from an approved semen collection centre

COUNTRY								Veterinary certific	ate to EU	
	I.1. Consignor  Name  Address  Tel.			I.2. Certificate reference number I.2.a.						
				I.3. Central Competent Authority						
ļ_					I.4. Local Competent Authority					
men	1.5.	I.5. Consignee  Name		I.6. Person responsible for the load in EU						
ligi					Name					
ous		Address Postal code Tel.			Address Postal code Tel.					
þ										
ils of dispatched consignment										
	1.7.	Country of origin ISO coo	e I.8. Region of origin	Code	I.9. Country of destination		ISO code	I.10. Region of destination	Code	
	I.11. Place of origin			•	I.12. Place of destination		n			
Part I: Details		Name Approval number			Name					
1Ξ		Address			Address	Address				
Parl		Name Approval number								
		Address	Address							
		Name	Approval number		Postal code					
		Address	ddress							
	1.13	I.13. Place of loading			I.14. Date of departure					
	1.18	5. Means of transport			I.16. Entry BIP in EU					
		Aeroplane Ship Railway wagon Road vehicle Other I								
						1.17.				
	Do	Documentary references:								
	1.18	3. Description of commodity			I.19. Commodity code (HS code)					
					05 11 10					
						l.20. Quantity				
	1.2	1.				I.22. Number of packages				
	I.23. Identification of container/Seal number							1.24.		
	I.25. Commodities certified for:									
Artificial reproduction ☐										
	I.26. For transit through EU to third Country					I.27. For import or admission into EU				
		Third country	ISO code							
	1.28	3. Identification of the commodi	es							
	Species Identifica (Scientific name)			tion mark			Quantity			

II.6.1.

satisfy the conditions laid down in Annex C of Directive 88/407/EEC;

COUNTRY **Bovine** semen II.a. Certificate reference number II. Health information I, the undersigned official veterinarian, hereby certify that: II.1. (name of exporting country) (2) Part II: Certification has been 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 and no vaccination against these diseases has taken place during the same period. 11.2. The semen described above was collected before 31 December 2004 at the semen collection centre which: II.2.1. meets the conditions laid down in Chapter I of Annex A to Directive 88/407/EEC; 11.2.2. is operated and supervised in accordance with the conditions laid down in Chapter II 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 period commencing 30 days prior to the date of collection of the semen to be exported and the 30 days after collection. 11.4. At the time semen described above was collected, all bovine animals standing at the semen collection centre: 11.4.1. came from herds and/or were born to dams which satisfy the conditions of paragraph 1(b) and (c) of Chapter I of Annex B to Directive 88/407/EEC: had tested negative, within the 30 days preceding the quarantine isolation period, to: 11.4.2. — the tests referred to in points 1(d)(i), (ii) and (iii) of Chapter I of Annex B to Directive 88/407/EEC, and - a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis, and - a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea, deferred until the animal reached the age of six months in the case of younger animals; 11.4.3. had undergone the 30-day quarantine isolation period and had tested negative to the following health tests: — a serological test for brucellosis carried out in accordance with the procedure described in Annex C to Directive 64/432/EEC; - either an immunofluorescent antibody test or a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings, or, in the case of a female animal, a vaginal mucus agglutination test; — a microscopic examination and culture test for Trichomonas foetus on a sample of preputial material or artificial vagina washings, or in the case of a female animal a vaginal mucus agglutination test; 11.4.4. had tested negative, at least once a year, to the routine tests referred to in points 1(a), (b) and (c) of Chapter II of Annex B to Directive 88/407/EEC. II.5. At the time the semen described above was collected, II.5.1. all female bovine animals in the centre had tested negative at least once a year to a vaginal mucus agglutination test for Campylobacter fetus infection, and 11.5.2. all bulls used for semen production had tested negative either to an immunofluorescent antibody test or to a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings carried out in 12 months prior to collection. II.6. The semen to be exported was obtained from donor bulls which

EN

COUNTRY Bovine semen

II.6.2.	
( <sup>1</sup> ) either	[were resident in the exporting country during the six months immediately prior to collection of the semen for export;]
( <sup>1</sup> ) or	[were imported from
II.6.3.	stand in a semen collection centre at which:
( <sup>1</sup> ) either	[all bovine animals were not vaccinated against infectious bovine rhinotracheitis and tested negative at least once a year to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis;]
( <sup>1</sup> ) or	[bovine animals not vaccinated against infectious bovine rhinotracheitis tested negative, at least once a year, to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis, at which testing for infectious bovine rhinotracheitis was not carried out on bulls which had received their first vaccination against infectious bovine rhinotracheitis at the insemination centre after they had tested negative to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis and which had been regularly re-vaccinated at intervals of not more than six months since the first vaccination;]
II.6.4.	
( <sup>1</sup> ) either	[have not been vaccinated against infectious bovine rhinotracheitis,]
( <sup>1</sup> ) or	[have been vaccinated against infectious bovine rhinotracheitis in accordance with point II.6.3,]
II.6.5.	fulfil the import conditions for bovine semen laid down in the Bluetongue Chapter of the Terrestrial Animal Health Code of the OIE, depending on the status of the country or zone of residence;****
II.6.6.	were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist:
II.6.7.	were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist:
II.6.8.	tested negative on two occasions not more than 12 months apart to a serum neutralisation test for Akabane virus carried out in approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen.*
II.7.	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.8.	The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC prior to its amendment by Directive 2003/43/EC.

## Notes

## Part I:

- Box reference 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 reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.23: identification of container and seal number shall be indicated.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: identification mark shall correspond to the identification of the donor animals and the date of collection that must be prior to 31 December 2004.

COUNTRY	Bovine semer
COONTIN	Dovine Seiner

## Part II:

- (1) Delete as necessary.
- (2) Countries listed in Annex I to Decision 2004/639/EC.
- (3) Standards for EHD virus diagnostic tests are described in the Bluetongue Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.
- \*\*\*\* To be used only by Australia, Canada and the USA.
- To be used only by Australia and the USA.
- To be used only by Canada.
- To be used only by Australia.
- The signature and the stamp must be in a different colour to that of the printing.

Official	veterinarian	
слистат	vererinarian	

Name (in capital letters): Qualification and title: Place: Signature: Stamp