

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			
Chilled <input type="checkbox"/>			Commercial document reference			
Frozen <input type="checkbox"/>			Date of issue			
Ambient <input type="checkbox"/>			Country			
Controlled temperature <input type="checkbox"/>			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		
	<p>I, the undersigned official veterinarian, certify the following:</p> <p>II.1. (insert EU Member State of origin or a region thereof)</p> <p>is recognised by the World Organisation for Animal Health (OIE) as a country or zone free of footand-mouth disease where vaccination is not practised and fulfills the conditions of the most recent Edition of the OIE Terrestrial Animal Health Code for a country or zone free of contagious bovine Pleuropneumonia</p> <p>II.2. The semen collection centre(1) at which the semen to be exported was collected:</p> <p>II.2.1. is approved and supervised by the Competent Authority in accordance with the EU legislation in force;</p> <p>II.2.2. is approved by SAG to export bovine semen to Chile and complies with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code;</p> <p>II.2.3. is under direct supervision of a veterinarian employed by the centre;</p> <p>II.2.4. maintains daily records of the resident animals' health;</p> <p>II.2.5. maintains records of daily production of semen that ensure its traceability;</p> <p>II.2.6. is physically isolated from other stockfarming establishments and meets biosecurity conditions;</p> <p>II.2.7. the staff working in the centre has no contact with animals outside the centre that could pose a health hazard to the species concerned.</p> <p>II.3. Admission of bulls in the centre: The centre only admits animals that:</p> <p>II.3.1. come from holdings free from bovine brucellosis (<i>Brucella abortus</i>) and bovine tuberculosis (<i>Mycobacterium bovis</i>) in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code;</p> <p>(2)either ○ [II.3.2. come from a country/region(2) free of bluetongue;]</p> <p>(2)or ○ [II.3.2. come from holdings under a surveillance program for bluetongue;]</p> <p>II.3.3. come from holdings that have not been subjected to health restrictions for infectious-contagious diseases notifiable for the species in the last 24 months.</p> <p>II.3.4. were subjected to a pre-entry quarantine of at least 28 days, which complies with the provisions set by the competent authority of the country and within that period have undergone with negative results(3), the routine diagnostic tests that are performed at the centre and which are in line with those set out in the most recent Edition of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE.</p> <p>II.4. The donor animals</p> <p>II.4.1. meet the requirements laid down in point II.3;</p> <p>II.4.2. were born in the country or region thereof mentioned in point II.1 or comes from a country with the same health status as described under point II.1;</p> <p>II.4.3. were examined on the day the semen was collected and there was no evidence of contagious diseases affecting the species, and have not been exposed to these diseases in the last 90 days prior to the collection of the semen.</p> <p>II.5. The animals standing permanently at the semen collection centre</p> <p>(2)either ○ [II.5.1. come from a country/region(2) free of bovine brucellosis;]</p> <p>(2)or ○ [II.5.1. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(4) for bovine brucellosis.]</p> <p>(2)either ○ [II.5.2. come from a country/region(2) free of bovine tuberculosis;]</p> <p>(2)or ○ [II.5.2. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(4) for bovine tuberculosis(5).]</p> <p>(2)either ○ [II.5.3. come from a country/region(2) free of bluetongue.]</p>		

Part II: Certification	II. Health information		
	(2)or	○ [II.5.3. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(4) for bluetongue.]	
	II.5.4.	have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code to a diagnostic test(4) for bovine viral diarrhoea(6).	
	(2)either	○ [II.5.5. come from a country/region(2) free of infectious bovine rhinotracheitis (IBR).]	
	(2)or	○ [II.5.5. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(4) for infectious bovine rhinotracheitis (IBR).]	
	(2)either	○ [II.5.6. come from a country/region(2) free of trichomonosis.]	
	(2)or	○ [II.5.6. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(4) for trichomonosis(5).]	
	(2)either	○ [II.5.7. come from a country/region(2) free of bovine genital campylobacteriosis.]	
	(2)or	○ [II.5.7. have been annually subjected in accordance with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code with negative results to a diagnostic test(3) for bovine genital campylobacteriosis(6).]	
	II.6.	The semen to be exported:	
II.6.1.	was collected, processed and stored in accordance with with the recommendations of the most recent Edition of the OIE Terrestrial Animal Health Code;		
II.6.2.	was diluted using sterile diluents to which the following antibiotics were added: ;(indicate the type of antibiotics and concentrations used)		
II.6.3.	has been stored only in sterilised vials over fresh nitrogen not used for any other purpose;		
II.6.4.	after collection and until its departure was kept in containers exclusively for export to Chile or with semen that meets at least the same requirements as those that must be achieved to export to Chile, and separate from any other semen.		
Notes			
Part I:			
·	Box I.11.: Place of origin: shall correspond to the semen collection or storage centres approved for export to Chile, from which the semen is dispatched.		
·	Box I.20.: Number of packages: shall correspond to the number of containers.		
·	Box I.21.: Identification of container and seal number shall be indicated.		
	Box I.25.: Donor identity: shall correspond to the official identification of the animal (eartag number).		
Date of admission to the centre: shall be indicated in the following format: dd/mm/yyyy.			
Approval number of the centre shall correspond to the approval number of the semen collection centre indicated in Box I.11.			
Part II:			
(1)	Only semen collection centres listed in accordance with Article 5(2) of Directive 88/407/EEC on the Commission website:		
	http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm		
	and listed in the register of authorized establishments for export to Chile:		
	http://www.sag.gob.cl/opendocs/asp/pagDefault.asp?boton=Doc51&argInstanciaId=51&argCarpetaId=1394&argTreeNodosAbiertos=(1394)(-51)&argTreeNodoActual=1394&argTreeNodoSel=8		
(2)	Delete as necessary.		
(3)	Seropositive bulls for bovine viral diarrhoea are accepted to the centre provided that their semen is tested for virus with negative results.		
(4)	Diagnostic tests must be carried out in official laboratories or authorised for this purpose by the competent authority of the exporting country.		
(5)	Relates only to donors or animals having contact with donors.		

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	(6)	<p>Animals negative to previous serological tests are retested to confirm absence of antibodies. Should an animal become serologically positive, every ejaculate of that animal collected since the last negative test has been either discarded or tested for virus with negative results. Prior to the initial dispatch of semen from BVD serologically positive bulls, a semen sample from each animal has been subjected to a virus isolation or virus antigen test for BVD. In the event of positive result, the bull is removed from the centre and all of its semen destroyed.</p> <ul style="list-style-type: none"> The signature and the stamp must be in a different colour to that of the printing. The certificate must be issued in Spanish and in the language of the EU Member State of origin. 		
	Certifying Officer			
	Name (in capital letters)	Qualification and title		
	Date of signature	Signature		
	Stamp			