

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			
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"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/> Controlled temperature <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						
051199 Other						
05119985 Other						
Commodity	Species	Breed/Category	Identification number	Date of collection/production		
Quantity						

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) is recognised by the World Organisation for Animal Health (OIE) as a country or zone free of foot-and-mouth disease where vaccination is not practised and fulfills the conditions of the most recent Edition of the Terrestrial Animal Health Code of the OIE for a country or zone free of contagious bovine pleuropneumonia.</p> <p>(2) <input type="checkbox"/> II.2. In the case of in vivo derived embryos, the semen used in for artificial insemination met Chilean import requirements for bovine semen (1).]</p> <p>II.3. The donor female:</p> <p>II.3.1. was born or has been resident in the country or region indicated in II.1. during a period of at least 6 months prior to the collection of <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) intended for export to Chile;</p> <p>II.3.2. was inspected on the day of collection and showed no clinical signs nor other evidence of contagious diseases in the last 90 days prior to the collection;</p> <p>II.3.3. comes from a holding 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 Terrestrial Animal Health Code of the OIE;</p> <p>II.3.4. comes from a holding on which there has been no clinical evidence of contagious diseases transmissible to animals of the bovine species in the last 90 days prior to collection, nor was subject to quarantine measures for animal health reasons.</p> <p>II.4. The <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) were collected and processed by an embryo <input type="radio"/> collection (2) / <input type="radio"/> production (2) team which:</p> <p>II.4.1. has been approved in accordance with Chapter I of Annex A to Directive 89/556/EEC (3);</p> <p>II.4.2. carried out the collection, processing, storing and transport of the embryos in accordance with Chapter II of Annex A to Directive 89/556/EEC;</p> <p>II.4.3. is subject to inspection by an official veterinarian at least twice a year;</p> <p>II.4.4. has not carried out activities in an infected area or in an area subject to restrictions for animal health reasons affecting animals of the bovine species.</p> <p>II.5. The storage and transport</p> <p>II.5.1. The <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) were stored in premises approved for that purpose by the Competent Authority, under the supervision of the team veterinarian, who is responsible for collection, processing, storage and transport of the <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) .</p> <p>II.5.2. During handling and processing of the <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) for export to Chile no <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) of a lower health status have been processed.</p> <p>II.5.3. The <input type="radio"/> ova (2) / <input type="radio"/> embryos (2) were put in straws which were identified and stored in sterilized or new containers over fresh nitrogen not used for any other purpose.</p>		

Part II: Certification	II. Health information								
	<p>Notes</p> <p>Part I:</p> <p>Box I.11.: Place of origin: shall correspond to the embryo collection or production teams approved for export to Chile, from which the embryos are dispatched.</p> <p>Box I.20.: Number of packages: shall correspond to the number of containers.</p> <p>Box I.21.: Identification of container and seal number shall be indicated.</p> <p>Box I.25.: Species: Select the species amongst the genus “Bos taurus”, “Bison bison” and “Bubalus bubalis” as appropriate.</p> <p>Donor identity: shall correspond to the official identification of the animal (eartag number).</p> <p>Date of collection: shall be indicated in the following format: dd/mm/yyyy.</p> <p>Approval number of the team shall correspond to the approval number of the embryo collection or production team indicated in Box I. 11.</p> <p>Part II:</p> <p>(1) Please check TRACES Export Health Certificate for frozen bovine semen to Chile.</p> <p>(2) Delete as appropriate.</p> <p>(3) Only embryo collection and production teams, approved as being in compliance with Chapter I of Annex A to Directive 89/556/EEC, and included in the list which can be found on the Commission’s website:</p> <p>http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm.</p> <p>and listed in the register of authorised establishments for export to Chile:</p> <p>http://www.sag.cl/ambitos-de-accion/importaciones-0/115/registros</p> <p>- The signature and the stamp must be in a different colour to that of the printing.</p> <p>- The certificate must be issued in Spanish and in the language of the EU Member State of origin.</p>								
<p>Certifying Officer</p> <table border="0" style="width: 100%;"> <tr> <td style="width: 50%;">Name (in capital letters)</td> <td style="width: 50%;">Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>				Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp	
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