

Veterinary certificate to EU

Part I : Details of consignment presented

I.1. Consignor Name Address Postal code / Region Country	I.2. Certificate reference number	I.2.a. TRACES reference number :
	I.3. Central Competent Authority	
	I.4. Local Competent Authority	
	I.6 Person responsible for load in EU	
I.5. Consignee Name Address Postal code / Region Country		I.6 Person responsible for load in EU
I.7. Country of origin, ISO code	I.8. Region of origin, Code	I.9. Country of destination ISO code I.10. Region of destination Code
I.11. Place of origin Holding <input type="checkbox"/> Semen centre <input type="checkbox"/> Establishment <input type="checkbox"/> Name Approval number Address Name Approval number Address Name Approval number Address		I.12. Place of destination
I.13 Place of loading Address Approval number		I.14 Date of departure
I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification:: Document:		I.16. Entry BIP in EU Name BIP unit no.: I.17. No.(s) of CITES
I.18. Description of commodity		I.19. Commodity code (HS code)
I.21 Temperature of products Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/>		I.20. Total Gross weight
I.22. Number of packages		I.24. Type of packaging
I.23. Identification of container/Seal number		I.25. Commodity certified as: Breeding <input type="checkbox"/> Fattening <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved bodies <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Quarantine <input type="checkbox"/> Game restocking <input type="checkbox"/> Registered horses <input type="checkbox"/> Pets <input type="checkbox"/> Circus <input type="checkbox"/> Relaying <input type="checkbox"/> Other <input type="checkbox"/> Humane consumption <input type="checkbox"/> Animal feedingstuff <input type="checkbox"/> Further process <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Technical use <input type="checkbox"/>
I.26. For transit to 3rd Country by EU		I.27. For import or admission into EU Definitive import <input type="checkbox"/> <input type="checkbox"/> Horses Re-entry <input type="checkbox"/> <input type="checkbox"/> Temporary admission horses <input type="checkbox"/> <input type="checkbox"/>
I.28. Identification of the commodity Species (scientific name) Treatment type Slaughterhouse Cutting/manufacturing plant Cold store Number of packages Type of packaging Batch number Nature of commodities Net weight		

II. Health information	II.a. Certificat reference number	II.b. TRACES reference number		
<p>1. Animal Health attestation</p> <p>I, the undersigned official veterinarian, hereby certify :</p> <p>1.1. that the heat-treated milk / milk-based product made from heat-treated milk / heat-treated milk-based product (3.) (3.) described above has been obtained from animals :</p> <ul style="list-style-type: none"> (a) under the control of the official veterinary service, (b) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and, <ul style="list-style-type: none"> I subject to regular veterinary inspectors inspections to ensure that they satisfy the animal health conditions laid down in Annex A, Chapter I of Directive 92/46/EEC ; <p>1.2. that I am familiar with the animal health requirements of Directive 92/46/EEC.</p> <p>2. Public Health attestation</p> <p>I, the undersigned official inspector, hereby certify :</p> <p>2.1. that the heat-treated milk / milk-based product made from heat-treated milk / heat-treated milk-based product (3.) (3.) described above :</p> <p>2.1. that the raw milk product described above:</p> <ul style="list-style-type: none"> (a) was manufactured from raw milk : <ul style="list-style-type: none"> (i) not, according to the findings of monitoring plans at least equivalent to those provided for in Directive 92/46/EEC, containing residues of anti-microbial substances in excess of the limits laid down in Annexes I and III to Regulation (EEC) No 2377/90, as amended, (ii) not, according to the findings of monitoring plans at least equivalent to those provided for in Directive 92/46/EEC, containing pesticide residues in excess of the maximum levels laid down in Annex II to Directive 86/363/EEC, as amended, (iii) not, according to the findings of monitoring plans at least equivalent to those provided for in Directive 92/46/EEC, containing contaminants in excess of the maximum tolerances laid down in the Community list provided for in Article 2 (3) of Regulation (EEC) No 315/93, (iv) which comes from registered and checked holdings meeting the hygiene conditions laid down in Chapter II of Annex A to Directive 92/46/EEC, (v) which was obtained, collected, cooled, stored and transported in accordance with the specific hygiene conditions laid down in Chapter III of Annex A to Directive 92/46/EEC, (vi) which meets the plate and somatic cell count standards laid down in Chapter IV of Annex A to Directive 92/46/EEC, and (vii) which, where necessary, was collected and standardized in accordance with the hygiene conditions laid down in Chapters I, III and IV of Annex B to Directive 92/46/EEC; (b) comes from a processing establishment offering equivalent guarantees to those laid down in Chapter II of Directive 92/46/EEC shown on the list of establishments authorized to export to the European Community and which is subject to supervision by the competent authority in accordance with the provisions of Chapter VI of Annex C to Directive 92/46/EEC; <p>I has undergone, prior to import into the territory of the Community :</p> <ul style="list-style-type: none"> (i) either a sterilization process, to achieve an F0 value equal to or greater than three , (4.) or (ii) an ultra high temperature (UHT) treatment at 132 °C for at least one second , (4.) or (iii) a high temperature short time pasteurisation treatment at 72 °C for at least 15 seconds or equivalent pasteurisation effect achieving a negative reaction to a phosphatase test (HTST) applied twice to milk with a Ph equal to or above 7,0 , (4.) or (iv) a HTST treatment of milk with a Ph below 7,0 , (4.) or (v) a HTST treatment combined with another physical treatment by : <ul style="list-style-type: none"> (v)(1) either lowering the Ph below 6 for one hour , (4.) or (v)(2) additional heating to 72 °C or more, combined with dissaccation; (4.) (4.) (g) was transported, where appropriate, in tanks as described in Article 16 (2) of Directive 92/46/EEC; <p>2.2. that I am aware of the provisions contained in Directive 92/46/EEC, Annexes I and III to Regulation (EEC) No 2377/90, Annex II to Directive 86/363/EEC and Regulation (EEC) No 315/93.</p>				
<p>(1) Issued by the competent authority.</p> <p>(2) Country and ISO code of the territory as appearing in Annex I to Commission Decision 2004/438/EC (as last amended).</p> <p>(3) To be specified if the authorization to import into the Community is restricted to certain regions of the third country concerned.</p> <p>(4) Complete as appropriate.</p> <p>(5) The registration number(s) of rail-wagon or lorry and the name of the ship should be given as appropriate. If known, the flight number of the aircraft. In case of transport in containers or boxes, the total number, their registration and seal numbers, if present, should be indicated under points 1.20 and 1.23.</p> <p>(6) Complete if appropriate.</p> <p>(7) The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.</p>				
<p>Official veterinarian or official inspector</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;"> Name (in Capital): Local Veterinary Unit: Date: Stamp </td> <td style="width: 50%; border: none;"> Qualification and title: LVU N°: Signature: </td> </tr> </table>			Name (in Capital): Local Veterinary Unit: Date: Stamp	Qualification and title: LVU N°: Signature:
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