

Part I : Details of consignment	I.1. Consignor Name Address Country		ISO Code		I.2. IMSOC Reference I.2.a. Local Reference	
	I.5. Consignee Name Address Country		ISO Code		I.3. Central competent authority	
					I.4. Local competent authority	
	I.7. Country of origin		ISO Code		I.9. Country of destination	
					ISO Code	
	I.8. Region of origin		Code		I.10. Region of destination	
	I.11. Place of Dispatch Name Address Approval Number Country		ISO Code		I.12. Place of destination Name Address Approval Number Country	
					ISO Code	
	I.13. Place of Loading Name Address Approval Number Country		ISO Code		I.14. Date and time of departure	
	I.15. Means of Transport				I.16 Entry Point	
	Mode	International transport document	Identification			
	I.18. Transport conditions Frozen <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Ambient <input type="checkbox"/> Chilled <input type="checkbox"/>				I.17. Accompanying documents Accompanying document reference Date of issue Country Place of issue	
I.19. Container No / Seal No						
I.20. Certified as Pharmaceutical use <input type="checkbox"/> Other <input type="checkbox"/> Slaughter <input type="checkbox"/> Relaying <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Fattening <input type="checkbox"/> Production <input type="checkbox"/> Breeding <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Animal Feedingstuff <input type="checkbox"/> Human consumption <input type="checkbox"/> Technical use <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	ISO Code	Country	ISO Code			
EU Exit Authority	BCP code					
EU Entry Authority	BCP code					
I.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	I.25. Total gross weight			
I.28. Description of consignment 1. 35 ALBUMINOIDAL SUBSTANCES; MODIFIED STARCHES; GLUES; ENZYMES 3501 Casein, caseinates and other casein derivatives; casein glues						
#1.	Commodity	Quantity	Net weight	Package count		
	Species	Identification number	Identification system			

Part II: Certification	II. Health information		
	<p>I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council, and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Section 4 of Chapter II of Annex X, and Chapter I of Annex XIV thereto, and certify that the milk(2), the milk-based products (2) and milk-derived products(2) referred to in box I.28 comply with the following conditions:</p> <p>II.1. they were produced and derived in (insert name of exporting country)(3), (insert name of region)(3), which is listed in Part I of Annex II to Commission Regulation (EU) No 605/2010, and which has been free from foot-and-mouth disease (FMD) and rinderpest for a period of 12 months immediately prior to export and has not practised vaccination against rinderpest during that period;</p> <p>II.2. they were produced from raw milk derived from animals which at the time of milking did not show clinical signs of any disease transmissible through milk to humans or animals, and which had been kept for a period of at least 30 days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;</p> <p>II.3. they are milk or milk products that:</p> <p>(2) <input type="radio"/> either [have undergone one of the treatments or combinations thereof described in point II.4;]</p> <p>(2) <input type="radio"/> or [comprise whey to be fed to animals of species susceptible to foot-and-mouth disease, and that whey was collected from milk subjected to one of the treatments described in point II.4 and:</p> <p>(2) <input type="radio"/> either [the whey was collected at least 16 hours after clotting and has a pH below 6;]</p> <p>(2)(5) <input type="radio"/> or [the whey has been produced at least 21 days before the shipping and during that period no cases of FMD have been detected in the exporting country;]</p> <p>(2)(5) <input type="radio"/> or [the whey has been produced on / / , this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border control post of the point of entry into Great Britain, Channel Islands or Isle of Man;]</p> <p>II.4. they have been subject to one of the following treatments:</p> <p>(2) <input type="radio"/> either [high temperature short time pasteurisation at 72oC for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test in bovine milk, in combination with:</p> <p>(2) <input type="radio"/> either [a subsequent second high temperature short time pasteurisation at 72oC for at least 15 seconds or an equivalent pasteurisation which itself achieves a negative reaction to a phosphatase test in bovine milk;]</p> <p>(2) <input type="radio"/> or [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72oC or higher;]</p> <p>(2) <input type="radio"/> or [a subsequent process by which the pH is reduced and kept for at least one hour at a level below 6;]</p> <p>(2)(5) <input type="radio"/> or [the condition that the milk/milk product has been produced at least 21 days prior to the date of shipping and during that period no cases of FMD have been detected in the exporting country;]</p> <p>(2)(5) <input type="radio"/> or [the milk/milk product has been produced on .../.../...(insert the date), this date, in consideration of the foreseen voyage duration, being at least 21 days prior to the date that the consignment is presented to a border control post of the point of entry into Great Britain, Channel Islands or Isle of Man;]</p> <p>(2) <input type="radio"/> or [sterilisation at a level of at least F03;]</p> <p>(2) <input type="radio"/> or [ultra high temperature treatment at 132oC for at least one second in combination with:</p> <p>(2) <input type="radio"/> either [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72'C or higher;]</p> <p>(2) <input type="radio"/> or [a subsequent process by which the pH is reduced and kept for at least one hour at a level below 6;]</p> <p>(2)(5) <input type="radio"/> or [the condition that the milk/milk product has been produced at least 21 days prior to the date of shipping and during that period no cases of FMD has been</p>		

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			detected in the exporting country;]
	(2)(5)	○ or	[the milk/milk product has been produced on .../.../...(insert the date), this date, in consideration of the foreseen voyage the duration, being at least 21 days prior to the date that consignment is presented to a border control post of the point of entry into Great Britain, Channel Islands or Isle of Man;]]
	II.5.		every precaution was taken to avoid contamination of the milk/milk-based product/milk-derived product after processing;
	II.6.		the milk/milk-based product/milk-derived product was packed:
	(2)	○ either	[in new containers;]
	(2)	○ or	[in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]
			and the containers are marked so as to indicate the nature of the milk/milk-based product/milk-derived product and bear labels indicating that the product is Category 3 material and not intended for human consumption;
	II.7.		the milk, milk-based products and milk-derived products described above:
	(2)	○ either	[does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for farmed animals, other than fur animals.]
(2)	○ or	[contains milk or milk products of ovine or caprine animal origin and is intended for feed for farmed animals, other than fur animals, and the milk or milk products:	
	(a)	are derived from ovine and caprine animals which have been kept continuously since birth in a country where the following conditions are fulfilled:	
	(i)	classical scrapie is compulsorily notifiable;	
	(ii)	an awareness, surveillance and monitoring system is in place for classical scrapie;	
	(iii)	official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classical scrapie;	
	(iv)	ovine and caprine animals affected with classical scrapie are killed and destroyed;	
	(v)	the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;	
	(b)	originate from holdings where no official restrictions are imposed due to a suspicion of TSE;	
	(c)	originate from holdings where no case of classical scrapie has been diagnosed during a period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:	
	(2)	○ either [all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]	
	(2)	○ or [all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001 , of all of the following animals which are over the age of 18 months, except ovine animals	

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		<p>of the ARR/ARR genotype:</p> <ul style="list-style-type: none"> - animals which have been slaughtered for human consumption; and - animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]] 	
	Notes		
	(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.		
	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).		
	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, Channel Islands or Isle of Man: this box is required to be filled in only if it is a certificate for a commodity to be transited through Great Britain, Channel Islands or Isle of Man; it may be filled in if the certificate is for a commodity to be imported into the Great Britain, Channel Islands or Isle of Man.	
	- 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.15:	Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading, the consignor must inform the border control post of the point of entry into Great Britain, Channel Islands or Isle of Man .	
	- Box reference I.16:	Do not use this box until the end of the transitional staging period.	
	- Box reference I.19:	use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.01; 04.02; 04.03; 04.04; 23.09.10, 23.09.90, 35.01, 35.02 or 35.04.	
	- Box reference I.23:	for bulk containers, the container number and the seal number (if applicable) must be included.	
	- Box reference I.25:	technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.	
	- Box reference I.26 and I.27:	fill in according to whether it is a transit or an import certificate.	
	- Box reference I.28:	'Manufacturing plant': provide the registration number of treatment or processing establishment.	
	Part II:		
	(2)	Delete as appropriate.	
	(3)	For completion if the authorisation to import into or transit through Great Britain, Channel Islands or Isle of Man is restricted to certain regions of the third country concerned.	
	(5)	this condition applies only to third countries listed in column 'A' of Annex I to Regulation (EU) No	

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	<p>605/2010.</p> <ul style="list-style-type: none"> - The signature and the stamp must be in a different colour to that of the printing. - Note for the person responsible for the consignment in Great Britain, Channel Islands or Isle of Man: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border control post. 			
	Certifying Officer Name (in capital letters) Date of signature Stamp		Qualification and title Signature	