Official certificate to the EU

	I.1. Consignor			I.2. Certificate	reference	I.2.a. IMSC	OC reference	
	Name			I.3. Central co	mpetent authority	Specimen n	ot to be used for imports in	nto the
	Address				petent authority	EU	ot to be used for imports in	ito die
	Country	ISO Cod	P		,			
	country	100 000						
	I.5. Consignee	I.6. Responsible for the consignment in EU						
ا۔	Name	Name						
副	Address			Address				
וַ		100.0-4	1_			TC	0.0-4-	
31	Country	ISO Cod	le	Country		15	O Code	
: Details of consignment	I.7. Country of origin ISO Co	ode I.8. Region	of origin Code	I.9. Country of destination	f ISO Co	de I.10. R	egion of destination	Code
ಶ∤	I.11. Place of dispatch	I		I.12. Place of o	lestination			1
5	_				icotinution			
2	Name			Name				
ਰ∣	Address			Address				
۲	Approval Number			Approval Nu	mber			
-	Country	ISO	Code	Country			ISO Code	
۱.								
4	I.13. Place of loading			I.14. Date and	time of departure			
4	Name							
	Address							
	Approval Number							
	Country	ISO	Code					
	Country	130	code					
İ	I.15. Means of Transport			I.16. Entry poi	nt			
		Identification		into Entry po				
	Mode International transport	iueittiiicati	Ш					
	document							
ŀ	I.18. Transport conditions			I 17 Accompa	nying documents			
	Ambient Chilled	П	Frozen 🗆	Туре	nymg documents			
	Alibient 🗀 Cimed		FIOZEII 🗀	Number				
ŀ	I 10 Container No / Cool No			Mulliber				
	I.19. Container No / Seal No							
ŀ	I.20. Certified as							
	Products for human consumption	۰п						
	Froducts for fidinali consumption							
ŀ	I.21. For transit			I.22. For inter	nal market			
	Non-EU	ISO	Code	I.23. For re-en				
- 1	I.24. Total number of packages	I.25. Total q		I.26. Total net		126 T	otal gross weight	
	1.24. Total number of packages	1.25. Total q	uantity	1.20. Total fiet	weigitt	1.20. 1	otai gross weight	
ŀ	I.27. Description of consignment							
	•							
	1. 04 DAIRY PRODUCE; BIRDS' EC					WHERE SPI	ECIFIED OR INCLUDED	
	0401 Milk and cream, not cond	centrated nor co	ntaining added sugar or o	other sweetenir	ng matter			
	Commodity Sp.	ecies	Cold store		Identification mar	·k	Package count	
	The state of the s							
							l	
- 1		eatment type	Nature of com	nmodity	Batch number		Final consumer	
	Net weight Tre							
	Net weight 110				Dlant / F	stablishme	nt / Centre	
			Manufacturing plant				,	
	Date of collection/production		Manufacturing plant		Flaitt / L			
•			Manufacturing plant		Flatt / L			
-			Manufacturing plant		Fidit / L			
=			Manufacturing plant		Flattt/L			
=			Manufacturing plant		FIGHT / L			
=			Manufacturing plant		FIGHT			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
=			Manufacturing plant		Figure			
-			Manufacturing plant		Figure 1			
			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure			
-			Manufacturing plant		Figure 1			

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	II. Health information		
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Part II: Certification			
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Country

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	II. Health info	rmation							
	□ II.1.	Public heal	th attestatio	on [Delete when the Uni	on is not the final destination	of the raw milk]			
tion	European I Council , Re of the Euro	Parliament a egulation (E pean Parlia	and of the Co C) No 853/20 ment and of	ouncil , Regulation (EC) 004 of the European Par f the Council and Comm	t requirements of Regulation (No 852/2004 of the European cliament and of the Council an dission Implementing Regulati I in accordance with these req	Parliament and of the d Regulation (EU) 2017/625 on (EU) 2019/627 and hereby			
Part II : Certification		(a)	it comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Articles 49 and 50 of Implementing Regulation (EU) 2019/62						
		(b)	it was produced, collected, cooled, stored and transported in accordance with the hy- conditions laid down in Annex III, Section IX, Chapter I, to Regulation (EC) No 853/20						
		(c)	it meets the plate and somatic cell count criteria laid down in Annex III, Section IX, Chapte I, to Regulation (EC) No 853/2004;						
		(d) it comes from animals belonging to herds free or officially free of brucellosis and tuberculosis;							
		(e) the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Article 29 of Council Directive 96/23/EC, are fulfilled and milk is listed in Commission Decision 2011/163/EU for the concerned country of origin;							
		(f)	operator in point 4, to residues of	n accordance with the re Regulation (EC) No 853/2	antibacterial drugs carried ou equirements of Annex III, Sect 2004, complies with the maxir y medicinal products laid dow 7/2010;	ion IX, Chapter I, Part III, num residue limits for			
		(g)	levels for p and of the	esticides laid down in F	tions guaranteeing compliance Regulation (EC) No 396/2005 of num levels for contaminants l	the European Parliament			
	II.2.		nal health attestation [Delete when the raw milk is derived from solipeds, leporidae or other d mammals others than ungulates]						
	The raw m	The raw milk described in Part I:							
	II.2.1.	health/offic XVII to Con infection w	ttes from the zone with code: (2) which, at the date of issue of this animal official certificate is authorised for the entry into the Union of milk and listed in Part 1 of Annex Commission Implementing Regulation (EU) 2021/404, and in which foot and mouth disease and on with rinderpest virus have not been reported for the period of 12 months prior to the date of g, and vaccination against these diseases has not been carried out during the same period.						
	II.2.2.		een obtained from animals of the species \square [Bos Taurus,](1) \square [Ovis aries,](1) \square [Capra hircus,](1) balus bubalis,](1) \square [Camelus dromedarius](1) that:						
		(1) □ either	[have remained in the zone referred to under point II.2.1. since birth, or for the periodeast 3 months prior to the date of milking;]						
		(1) □ and/or			red to under point II.2.1. fron				
			(1) □ either	into the Union of milk,	or territory, or zone thereof vectors or colostrum or colostrum-based period of at least 3 months p	d products and the animals			
			(1) □ and/or	[a Member State;]]					
	II.2.3. has been obtained from animals coming from establishments:								
		(a) registered by and under the control of the competent authority of the third country or territory and have a system in place to maintain and to keep records in accordance with Article 8 of Commission Delegated Regulation (EU) 2020/692;							

Country

II II albi to						
II. Health info	rmation					
	(b)	which receive regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex I to Commission Delegated Regulation (EU) 2020/692 and emerging diseases;				
	(c)	which were not subject to national restriction measures for animal health reasons, including the relevant listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 an emerging diseases, at the date of milking.				
Notes						
from the E Protocol or	uropean U n Ireland/N	he Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland Union and the European Atomic Energy Community, and in particular Article 5(4) of the Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union i Ficial certificate include the United Kingdom in respect of Northern Ireland.				
		fficial certificate is intended for the entry into the Union of milk, including when the Union is ion of such milk.				
		fficial certificate shall be completed in accordance with the notes for the completion of for in Annex I, Chapter 4, to Implementing Regulation (EU) 2020/2235.				
Box reference I.8:	Provide the code of the zone as appearing in column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.					
Box reference I.11:	Name, ad	ldress and approval number of the establishment of dispatch.				
Box reference I.15:	Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (vessel) must be provided. In the case of unloading and reloading, the consignor must inform the border control post of the entry into the Union.					
Box reference I.19:	For the co	or the containers or boxes, the container number and the seal number (if applicable) shall be included.				
Box reference I.27:	Use the a	ppropriate Harmonised System (HS) code under the following headings: 04.01; 04.02 or 04.03.				
		Description of consignment:				
		"Manufacturing plant": Introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to the European Union.				
Part II:						
(1)	-	appropriate.				
(2)	Regulatio	he zone in accordance with column 2 of the table in Part 1 of Annex XVII to Implementing on (EU) 2021/404.				
(3)	to be sigr	•				
-		an official veterinarian when Part II.2 Animal health attestation is not deleted,				
-		a certifying officer or an official veterinarian when Part II.2 Animal health attestation is deleted.				
Official veteri	narian or Off	icial inspector				
Name (in cap Date of signa Stamp		Qualification and title Signature				

en 4/4