Export Health Certificate

	01061300	Kamele (Cameno					
•		ERE					
- 1	I.23. Gesamtanzahl an Packungen I.24. Gesamtmenge				I.25. Nettogesamtgewicht	I.25. Bruttogesamtg	gewicht
	EU Entry EU Entry Authority BCP code BCP code			Country	Ländercode		
	Country ISO- Ländercode				Country	ISO-	
	Breeding □ Vermittlung □ Technische Verwendung □ I.21. Für die Durchfuhr durch ein Drittland □				Futtermittel Künstliche Vermehrung I.22. Für die Durchfuhr durch Mitgliedstaaten		
	I.20. Waren zertifiz Production Sonstiges		Pharmazeutische Ver		Mast □ Menschlicher Verzehr □	Breeding and prod Schlachtung □	uction \square
	I.19. Containernum	nmer/Plombennun	nmer		Land	ort	
	I.18. Beförderungsl Umgebungstemper				I.17. Begleitdokumente Bezugsnum mer des Handelspapi ers	Ausstellungs datum Ausstellungs	
	I.15. Transportmitt	el Dokument	Identifikation		I.16 Entry Point		
_	Zulassungsnumme Land	er	ISO- Ländercode				
	I.13. Ladeort Name Adresse				I.14. Datum und Uhrzeit des Ab	otransports	
	Name Adresse Zulassungsnumme Land	er	ISO- Ländercode		Name Adresse Zulassungsnummer Land ISO- Ländercode		
	I.8. Ursprungsregio	n		Code	I.10. Region des Bestimmungso I.12. Bestimmungsort	orts	
TETT	I.7. Ursprungsland			ISO- Ländercode	I.9. Bestimmungsland		ISO- Ländercode
-	Name Adresse Land		ISO- Ländercode		I.4. Zuständige örtliche Behörde		
	I.5. Empfänger				I.3. Zentrale zuständige Behörde		
	Adresse Land		ISO- Ländercode				
	Name				I.2. IMSOC-Bezugsnummer I.2.a. Lokale Bezugsnummer		

EUROPÄISCHE UNION (GB) Model RUM-A / GBHC124E (v								
	II. Gesundhei	tsinformatione	en					
	II.1.	Animal he	alth attestat	ion				
Part II: Certification			I, the undersigned official veterinarian responsible for the approved body, institute or centre/holding (1) of origin certify that the animals described in Part I meet the following requirements:					
		II.1.1.	They come from the country, territory or part thereof described in Box I.7.:					
			(a)	where the diseases refe	rred to in this certificate are	notifiable,		
			(b)	which at the date of issurinderpest.	uing this certificate has been	free for 12 months from		
		II.1.2.	They come	e from the body, institute	or centrer/holding (1) descri	bed in Box I.11;		
			(a)		rding to the requirements ar ulation (EU) No 206/2010;	nd conditions set out in Part 3		
	the control of infectious diseases to which the anima are susceptible;							
			(c)	where there have been animals referred to in B	no clinical cases of the follov ox I.28. are susceptible:	ving diseases to which the		
				- anthrax for	the last 30 days;			
				stomatitis, r disease, pes	outh disease, bluetongue, Rift abies, contagious bovine ple te des petits ruminants, shee uropneumonia for the past 6	uropneumonia, lumpy skin p pox, goat pox, contagious		
			(d)	where there have been brucellosis for the past	no clinical or non-clinical ca 6 months;	ses of tuberculosis and		
			(e)	case of the following dis susceptible: foot-and-me	a of 10 km radius for the last seases to which the animals nouth disease, vesicular stoma e des petits ruminants, sheep nia;	referred to in Box I.28. are attitis, contagious bovine		
			(f)	case of the following dis	seases to which the animals i	st 30 days, there has been no referred to in Box I.28. are ease, Rift valley fever, lumpy		
	(1)	o either	[(g)	in which they have rem dispatch to Great Britain	ained since birth or for the pn.]	past 6 months before		
		o Or (1)	[(h)	that at that date was au animals has not been in	I.2.1, from the EU territory w	als into Great Britain and the en-hoofed animals from		
		II.1.3.	They:					
			(a)	same health requiremen	tion from the approved body	icate for the last 30 days and		
			(b)		fficial veterinarian within 24 of disease and are fit for the			
			(c)	are not animals to be ki diseases.	lled under a national progra	mme for the eradication of		
	II.1.4. Foot-and-Mouth Disease							
		o either(1)	[(a)		st 12 months from foot-and-	of described in Box I.7 which mouth disease with or		
		or(1)	[(a)	They have been subject	ed to the following tests:			

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	_	ROPAISCHE UNION			(GD) Model	RUM-A / GBHC124E (V1.2)	
		II. Gesundheitsinformatione	n				
	ľ			infection ca for internat Tests and V	al test for evidence of foot-and arried out in accordance with tional trade laid down in the accines for Terrestrial Anima we results, taken within 10 da	one of the prescribed tests OIE Manual of Diagnostic ls (OIE Terrestrial Manual),	
	Part II: Certification			infection ca the OIE Ter days prior t occasions 1	bang test for evidence of foot arried out in accordance with restrial Manual with negative to dispatch to Great Britain] (2 5 days apart, the second of wo or to dispatch to Great Britain	the procedures described in e results], (1)(3) \circ [taken 10 l)(4) \circ [taken on two hich must have been taken	
	Pa	(1)	(b)	they have not been vac	cinated against foot-and-mou	th disease.	
		II.1.5.	Bluetongu	e and Epizootic haemorr	hagic disease (EHD)		
		o either(1)	[They come from the country, territory or part thereof described in Box I.7 which has been free for 24 months from bluetongue/EHD in accordance with the OIE Terrestrial Animal Health Code (OIE Terrestrial Code).]				
		o or(1)	centre/hol according	ding (1) for at least 30 da to the OIE Terrestrial Ma	ted facility in the approved bys prior to shipment and wer nual, with negative results, c d body, institute or centre.]	e subjected to a serology test	
			o or(1)	centre/holding (1) for a PCR test according to th	ector-protected facility in the t least 30 days prior to shipm ne OIE Terrestrial Manual, wi or introduction into the appro	ent and were subjected to a th negative results, carried	
			o or(1)	to an serology test acco	sonally free area and were su rding to the OIE Terrestrial M lays after introduction into th	Ianual, with negative results,	
			o or(1)	to a PCR test according	sonally free area and were su to the OIE Terrestrial Manua lays after introduction into th	l, with negative results,	
		II.1.6.	Rift valley	fever			
		o either(1)			tory or part thereof described fever and have not been vac		
	or(1) [They were held in a vector-prote centre/holding (1) for at least 30 d no clinical signs of Rift valley feve protected facility and the place of shipment.]		ys prior to shipment during vertical and were protected from vertical from vertical from vertical and were protected from vertical from the contract of the co	which the animals showed ctors between the vector-			
	or(1) [They have been subjected to a vir of Rift valley fever, as laid down a Terrestrial Manual, taken at the b days later on, the second of which Great Britain.]			ey fever, as laid down ar l Manual, taken at the be on, the second of which	nd prescribed for internationa ginning of the isolation/quara	al trade by the OIE antine period and at least 42	
		II.1.7. Brucellosis					
	either(1) [They come from a country, territoring free for the past 12 months from b that disease;]			e past 12 months from bi			
	o or(1) [They have been subjected to a tes the OIE Terrestrial Manual, in the						
	or(1) [They are castrated males of any a			castrated males of any ag	ge].		

	II. Gesundheitsinformation	ion.		\ ,			
	ii. Gesululieitsiilioi liiation	ien					
	II.1.8.	Other va	ccinations				
		(a)	They have not been va	ccinated against vesicular sto	matitis,		
	(5)	(b)	They have been vaccin	They have been vaccinated against:			
 ¤		(1) 🗆	[anthrax on the vaccine(s)	(dd/mm/yyyy)(date(s)) (name of vaccine(s) used)],	with the following		
Part II: Certification		(1) 🗆		(dd/mm/yyyy)(date(s)) v f vaccine(s) used) and a blood yyyy)(date(s)) shows a protect			
.	II.1.9.	Parasite	treatment				
Part II:		They have been treated at least twice during the 40 days prior to dispatch to Great Britain against internal and external parasites with the following product(s) . Specify the active ingredients and the doses of the products used					
	II.1.10.	Loading	g on the means of transport				
		They have been loaded for dispatch to Great Britain on (dd/mm/yyyy)(6) in the means of transport described in Box I.15. that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.					

II. Gesundheitsinformationen

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway 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.

This certificate is to be used for live animals listed in the note for Box 1.25 coming from an approved body, institute or centre in a third country, territory of part thereof, and destined to an approved body, institute or centre situated within Great Britain. Use one certificate per species.

Part I:

Part II:

Box Registration number (railway wagons or container and lorries), flight number (aircraft) or reference name (ship) is to be provided. In case of unloading and reloading, the consignor shall inform

I.15.: the BCP of entry into Great Britain.

Box Do not use this box until the end of the transitional staging period.

reference

Box Use appropriate HS code: 010613 or 010619.

reference I.28.:

Box Identification system: Specify the identification system (tag, tattoos, brand, chip,

reference transponder). The identifier shall include the ISO code of the exporting country and permit

I.28.: tracing of their premises of origin.

Age: months.

Sex (M = male, F = female, C = castrated).

Species: Select the species amongst those listed below:

Order Family Genera/species Artiodacty Antilocapr Antilocapra la idae

Bovidae

Addax ssp., Aepyceros ssp., Alcelaphus ssp., Ammodorcas ssp., Ammotragus ssp., Antidorcas ssp., Antilope ssp., Bison ssp., Bos ssp.(including Bibos, Novibos, Poephagus), Boselaphus ssp., Bubalus ssp. (including anoa), Budorcas ssp., Capra ssp., Cephalophus ssp., Connochaetes ssp., Damaliscus ssp.(including Beatragus), Dorcatragus ssp., Gazella ssp., Hemitragus ssp., Hippotragus ssp., Kobus ssp., Litocranius ssp., Madoqua ssp., Naemorhedus ssp. (including Nemorhaedus and Capricornis), Neotragus ssp., Oreamnos ssp., Oreotragus ssp., Oryx ssp., Ourebia ssp., Ovibos ssp., Ovis ssp., Patholops ssp., Pelea ssp., Procapra ssp., Pseudois ssp., Pseudoryx ssp., Raphicerus ssp., Redunca ssp., Rupicapra ssp., Saiga ssp., Sigmoceros-Alecelaphus ssp., Sylvicapra ssp., Syncerus ssp., Taurotragus ssp., Tetracerus ssp., Tragelaphus ssp. (including Boocerus).

Camelidae Camelus ssp., Lama ssp., Vicugna ssp.

Cervidae Alces ssp., Axis-Hyelaphus ssp., Blastocerus ssp., Capreolus ssp., Cervus-Rucervus

ssp., Dama ssp., Elaphurus ssp., Hippocamelus ssp., Hydropotes ssp., Mazama ssp., Megamuntiacus ssp., Muntiacus ssp., Odocoileus ssp., Ozotoceros ssp., Pudu

ssp., Rangifer ssp.

Giraffidae Giraffa ssp., Okapia ssp.

Moschidae Moschus ssp.

Tragulida Hyemoschus ssp., Tragulus-Moschiola ssp.

e

Part II:

(1) Keep as appropriate.

(2) This attestation is only applicable to Bovidae and Cervidae.

			(02) 1110 1101	ROWITT GETTOTE IE (VIII)			
	II. Gesundheitsinfor	rmationen					
		L					
		This attestation is only applicable to Bovidae and Cervidae other than African buffalo (Syncerus caffer).					
	(4) This	s attestation is only applicable to African bu	ffalo (Syncerus caffer).				
		Vaccination is not compulsory, but if the animals have been vaccinated, information on the vaccine(s) used and the time of vaccination shall be filled in.					
Part II: Certification	prio ther	Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to Great Britain of the third country, territory or part thereof described in Boxes I.7. and I.8., or during a period where restrictive measures have been adopted by Great Britain against imports of these animals from that country, territory or part thereof.					
ert	(7) Code	le of the territory as it appears in Part 1 of A	nnex 1 to Regulation (EU) No	206/2010.			
I: C	Certifying Officer	,					
rt I	Name (in capital le	etters)	Qualification and title				
Pa			Unterschrift				
	Stempel						