EUROPEAN UNION INTRA

	I.1. Consignor				I.2. IMSOC refe	erence	I.2.a. Local reference	
	Name Address						I.3. Central Competent Authority	
	Country ISO Code					I.4. Local Competent Authority		
of consignment	I.5. Consignee Name Address Country		ISO Code		I.6. Operator conducting assembly operations independently of an establishment Name Address Approval Number			
							ISO Code	
tion	I.7. Country of orig	gin	ISO	O Code	I.9. Country of	f destination	ISO Code	
CLID	I.8. Region of origi	in	Со	de	I.10. Region of	f destination	Code	
Part I: Description	I.11. Place of dispa Name Address Approval Number Country	atch	ISO Code		I.12. Place of d Name Address Approval Number Country	lestination	ISO Code	
Ì	I.13. Place of loadi	ng			I.14. Date and	time of departure		
	Name Address Approval Number Country ISO Code							
- 1					I.16. Transporter			
-	Mode	International transport document	Identification		Name Address Approval Number			
					Country		ISO Code	
	I.18. Transport conditions Frozen				I.17. Accompanying documents Document Type Accompanying document reference Date of issue Country Place of issue			
- 1					Ambient \square			
- 1					Transformation			
ļ								
					ISO Code BCP code BCP code			
ļ	I.22. For transit through Member State(s)		I.23. For expo	rt				
	Member State ISO Code		Third country ISO Code Exit point BCP code					
İ	I.24. Estimated journey time				I.25. Journey Log			
İ	I.26. Total number of packages I.27. Total quantity				I.28. Total net weight I.28. Total gross weight			
I.30. Description of consignment						•		
	1. 41 RAW HIDES AND SKINS (OTHER THAN FURSKINS) AND LEATHER							
	4102 Raw skins of sheep or lambs (fresh, or salted, dried, limed, pickled or prepared), whether or not with wool on or split, other than those excluded					reserved, but not tann) to this chapter	ed, parchment-dressed or further	
	#1. Commodity		e of commodity	Category		Manufacturing plant	Quantity	

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	Species	Batch number	Package count	Net weight	
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Part I: Description of consignment					
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EUROPEAN UNION

EU —	ROPEAN UNION		esta	EDIIShed for	r the prevention and contr	of of certain fisted diseases		
	II. Health information							
	I, the undersigned offi	icial veterin	arian. herek	ov certify the	at:	1		
	• (1)either [II.1					kept animals:		
	 (1)either [II.1 The animal by-products described in Part I were obtained from kept animals: (1)either [killed for the purpose of the prevention and control of: 							
		- (1)010101	(1)	l Larbone	_			
Part II: Certification		either			[(introduce the name of the relevant category A disease) following the instructions of the competent authority in accordance with Regulation (EU) 2020/687 and are intended for processing by			
Certifi				o (1) either	[Method 1 to 5.]]]			
ij				o (1) or	[incineration.]]]			
art				○ (1) or	[co-incineration.]]]			
Д			o (1)or	[following		,		
					of Regulation (EU) 2016/429			
				o (1) either	[Method 1 to 5.]]]			
				o (1) or	[incineration.]]]			
				o (1) or	[co-incineration.]]]			
o (1)or [not subject to killing by the competent authority for the pand control of category A diseases or emerging diseases, located in restricted zones established for the prevention diseases in accordance with					eases, kept in establishments ention and control of animal			
				o (1) either	[Delegated Regulation (EU) 2	2020/687,]		
				o (1) or	[temporary special disease of the control of the co	control measures referred to EU) 2016/429,]		
				o (1) or	[emergency measures adopt accordance with Article 259	ted by the Commission in of Regulation (EU) 2016/429,]		
	and the animal by-products are moved from that restricted zones in compliance with the conditions set out in (2), for the							
				o (1) either	from aquatic animals as set	as set out in Chapter II of ilage of by-products obtained out in point K of Section 2 of egulation (EU) No 142/2011.]]]		
				o (1) or	[processing or treatment by Annex X, Annex XI or Annex 142/2011.]]]			
				o (1) or	[production of processed pe referred to in Annex XIII to 142/2011.]]]	tfood, other than raw petfood Regulation (EU) No		
				o (1) or	[transformation into compo Section 1 of Chapter III of An 142/2011.]]]	st or biogas referred to in nnex V to Regulation (EU) No		
	o (1) or [II.1	species for (introduce competent	The animal by-products described in Part I were obtained from wild animals of listed species found dead or killed for the purpose of the prevention and control of (introduce the name of the relevant category A disease) following the instructions of the competent authority (3) in accordance with Article 64(2)(c) of Delegated Regulation (EU) 2020/687 and are intended for processing by					
		o (1) either	[Method 1	to 5.]]				

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2023/1521 (2021/1699) Animal by-products from restricted zones established for the prevention and control of certain listed diseases $\,$

	II. Health information									
			o (1) or	[incineration.]]						
		o (1) or [co-incineration.]]								
ification	(1)	either	ther [II.2 Unprocessed animal by-products other than hides and skins, hides and skins, colostrum, milk and dairy products, indicate as appropriate) obtained from bovine animals kept in vaccination zone I in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, points (3.5) and (3.7), to Commission Delegated Regulation (EU) 2023/361.]							
Part II: Certification	(1)	or	□ [II.2 Unprocessed animal by-products other than hides and skins, hides and skins, colostrum, milk and dairy products, indicate as appropriate) obtained from bovine animals kept in vaccination zone II in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, points (3.6) and (3.7), to Commission Delegated Regulation (EU) 2023/361.]							
	Notes									
	Part I:									
	•	Box refere	nce I.9 and	I.11: delete as appropria	ate.					
	•	Box refere	nce I.12, I.1	3 and I.17: approval nui	mber or registration number.					
	•	Box refere	nce I.14: co	mplete if different from	'I.1. Consignor'.					
	•	Box refere	nce I.25: for	r 'processing', 'treatmen	- 1					
	•			iture of commodity: 'AB Category 1','Category 2',	ABPs referred to in Article 22(5) of Delegated Regulation (EU)					
	Part II:									
	(1)	Delete as a	ppropriate.							
	(2)	Insert the number of the relevant article(s), the title and date of publication in the Official Journal of the European Union of the relevant legal act adopted by the Commission providing those conditions or the reference to the legal act or instruction approved and made public by the competent authority providing for those conditions.								
	(3)	See special legislation on the prevention of transmissible diseases.								
		icer/Official ve	terinarian							
	Name (in capi Date of signate Stamp				Qualification and title Signature					
						1				

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