	UNITED KINGDOM		TAG III I E		cial certificate to the EU
	I.1 Consignor/Exporter		I.2 Certificate refe		I.2a
	Name			4.	/
	Address		I.3 Central Compe	•	/
			DEPARTMENT FO FOOD & RURAL	R ENVIRONMENT, AFFAIRS	
	•		I.4 Local Compete		/
	Country	ISO country code	ANIMAL AND PL	ANT HEALTH AGENCY	
	I.5 Consignee/Importer		I.6 Operator respo	onsible for the consignment	
	Name		Name		
ii.	Address		Address		
Part I: Description of consignment	, Q <sup>2</sup>				
on (	Country	ISO country code	Country		O country code
ripti	I.7 Country of origin	ISO country code	I.9 Country of des	tination	ISO country code
Desc	I.8 Region of origin	Code	I.10 Region of dest	tination	Code
<b>:</b>	I.11 Place of dispatch	Registration/Approval No	I.12 Place of destin	antion ]	Registration/Approval No
Part	1.11 I face of dispatch	registration/Approvar 110	1.12 I late of uesti	iation	Negisuauon/Appiovai 110
	Name		Name		
	Address		Address		
		(			
	Country	ISO country code	Country		O country code
	I.13 Place of loading		I.14 Date and time	of departure	
	I.15 Means of transport		I.16 Entry Border	Control Post	
	☐ Aircraft	□ Vessel	I.17 Accompanyin	g documents	
					,
	□ Railway	☐ Road vehicle	Type	Coo	
	□ Kanway	Li Roau venicie	Country	ISC	country code
	Identification		Commercial docum	nent reference	
ŀ	I.18 Transport conditions	☐ Ambient	☐ Chilled	☐ Froze	n
Ī	I.19 Container number/Seal number				
	Container No		Seal No	•	
	I.20 Certified as or for				
	☐ Products for human consu	mption			
	I.21		I.22	internal market	
	Third country	ISO country code	I.23		
•	I.24 Total number of packages	1.25		I.26 Total net weight/gross	weight (kg)

τ	NITED KINGD	OOM				II.a Certificate		
I.	27 Description	of consignment				<u> </u>		
_1	CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type	Final consumer
	Nature of commodity	Number of packages	Batch No	Date of collection/ production	Manufacturing plant			
2	CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type	Final consumer
	Nature of commodity	Number of packages	Batch No	Date of collection/ production	Manufacturing plant			
3	CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type	Final consumer
	Nature of commodity	Number of packages	Batch No	Date of collection/ production	Manufacturing plant			
4	CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type	Final consumer
	Nature of commodity	Number of packages	Batch No	Date of collection/ production	Manufacturing plant			
5	CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type	Final consumer
	Nature of commodity	Number of packages	Batch No	Date of collection/ production	Manufacturing plant			

II.a Certificate reference

# UNITED KINGDOM

### II. Health information

II.1. Public health attestation [Delete when the Union is not the final destination of the meat products]

I, the undersigned, declare that I am aware of the relevant requirements of Regulation (EC) No 999/2001 of the European Parliament and of the Council, Regulation (EC) No 178/2002 of the European Parliament and of the Council, Regulation (EC) No 852/2004 of the European Parliament and of the Council, Regulation (EC) No 853/2004 of the European Parliament and of the Council and Regulation (EU) 2017/625 of the European Parliament and of the Council and hereby certify that the meat products (2), including rendered animal fats and greaves, meat extracts and treated stomachs, bladders and intestines others than casings, described in Part I were produced in accordance with these requirements, in particular that:

- II.1.1. they come from (an) establishment(s) applying general hygiene requirements and implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004, regularly audited by the competent authorities, and is listed as a Union approved establishment;
- [the animals from which the meat products were derived have passed *ante-mortem* and *post-mortem* inspections;] [the wild game from which the meat products were derived have passed *post-mortem* inspection;]
- II.1.3. they have been produced from raw materials which met the requirements of Sections I to VI of Annex III to Regulation (EC) No 853/2004;
- II.1.4 they have been marked with an identification mark in accordance with Section I of Annex II to Regulation (EC) No 853/2004;
- II.1.5. the label(s) affixed on the packaging of meat products described in Part I, bear(s) an identification mark to the effect that the meat products come wholly from fresh meat from establishments (slaughterhouses, game handling establishment and cutting plants) approved for entry into the Union;
- II.1.6. they satisfy the relevant criteria laid down in Commission Regulation (EC) No 2073/2005;
- II.1.7. the guarantees covering live animals and products thereof provided by the control plan submitted in accordance with Article 6(2) of Commission Delegated Regulation (EU) 2022/2292 are fulfilled and the concerned animals and products are listed in Annex -I to Commission Implementing Regulation (EU) 2021/405 for the concerned third country or territory;
- II.1.8. the means of transport and the loading conditions of meat products of this consignment meet the hygiene requirements laid down as regards the entry into the Union;
- (1) [II.1.9.1. if obtained from meat of domestic porcine animals, this meat fulfills the requirements of Commission Implementing Regulation (EU) 2015/1375, and in particular:
  - (1) either [has been subjected to an examination by a digestion method for Trichinella with negative results;]
  - (1) or [has been subjected to a freezing treatment in accordance with Annex II to Implementing Regulation (EU) 2015/1375:1
  - (1) (10) or [in the case of meat from demestic porcine animals kept solely for fattening and slaughter, comes from a holding or category of holdings that has been officially recognised by the competent authorities as free from *Trichinella* in accordance with Annex IV to implementing Regulation (EU) 2015/1375 or not weaned and less than five weeks of age;]]
- (1) [II.1.9.2. if obtained from meat of solipeds or wild boar, this meat fulfils the requirements of Implementing Regulation (EU) 2015/1375, and in particular, has been subjected to an examination by a digestion method for *Trichinella* with negative results;]
- (1) [II.1.9.3. the treated stomachs, bladders and intestines and meat extracts have been produced in accordance with Section XIII of Annex III to Regulation (EC) No 853/2004.]
- (1) [II.1.9.4. the rendered animal fats and greaves have been produced in accordance with Section XII of Annex III to Regulation (EC) No 853/2004.]
- (II) [II.1.10. if containing material from bovine, ovine or caprine animals, with regard to bovine spongiform encephalopathy (BSE): [the country or region of origin is classified in accordance with Commission Decision 2007/453/EC as a country or region posing a negligible BSE risk, and
  - (1) either [the animals from which the meat products are derived were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there have been no BSE indigenous cases;]]]
  - (1) and/or [the animals from which the meat products are derived originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there has been at least one BSE indigenous case, and the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;]]]
  - (1) and/or [the animals from which the meat products are derived originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a controlled BSE risk, and:
    - the meat products do not contain and are not derived from specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;
    - the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;
    - (iii) the animals from which the meat products are derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;]]]

NITED KINGDOM				II.a Certificate reference
	(1) and/or	-		om which the meat products are derived originate from a country or region classified in accordance 2007/453/EC as a country or region posing an undetermined BSE risk, and:
		(i)	the me	at products do not contain and are not derived from specified risk material as defined in point 1 of V to Regulation (EC) No 999/2001;
		(ii)		at products do not contain and are not derived from mechanically separated meat obtained from of bovine, ovine and caprine animals;
•		(iii)	of gas stunnin	mals from which the meat products are derived have not been slaughtered after stunning by means injected into the cranial cavity or killed by the same method or slaughtered by laceration after ng of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cavity;
* ~C		(iv)		mals from which the meat products are derived have not been fed with meat-and-bone meal or s, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;
O		(v)		at products were produced and handled in a manner which ensures that they do not contain and ot contaminated with nervous and lymphatic tissues exposed during the deboning process;]]]
(1) or	[the countrolled			gin is classified in accordance with Decision 2007/453/EC as a country or region posing a
	O	(a)	of gas stunning cranial	mals from which the meat products are derived have not been slaughtered after stunning by means injected into the cranial cavity or killed by the same method or slaughtered by laceration after ng of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cavity;
	(1) either	[(p)		at products do not contain and are not derived from:
			(i) (ii)	specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;
	(1) and/or	[(b)	(ii)	mechanically separated meat obtained from bones of bovine, ovine and caprine animals.] at products contain and are derived from treated intestines sourced from animals which were born,
	una/or	[(U)	continu 2007/4	uously reared and slaughtered in a country or region classified in accordance with Decision 153/EC as a country or region posing a negligible BSE risk in which there have been no BSE nous cases;
	(1) and/or	[(b)	from a	at products contain and are derived from treated intestines sourced from animals which originate country or region classified in accordance with Decision 2007/453/EC as a country or region a negligible BSE risk in which there has been at least one BSE indigenous case, and:
	(1)	either	[(i)	the animals were born after the date from which the ban on the feeding of ruminants with meat- and-bone meal and greaves derived from ruminants has been enforced;]]
	(1)	and/or	· [(ii)	the treated intestines of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001.]]
	<sup>(1)</sup> either	[(c)		mals from which the meat products are derived originate from a country or region classified in ance with Decision 2007/453/EC as a country or region posing a negligible or a controlled BSE
	(1) and/or	[(c)		mals from which the meat products are derived originate from a country or region classified in ance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and:
			(i)	the animals from which the meat products are derived have not been fed with meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;
			(ii)	the meat products were produced and handled in a manner which ensures that they do not contain and were not contaminated with nervous and lymphatic tissues exposed during the deboning process;]]]
<sup>(1)</sup> or	[the countrundetermine		_	gin is classified in accordance with Decision 2007/453/EC as a country or region with an
	(a) th	e animals	from wh	nich the meat products are derived have not been:
	(i)	sla	aughtered	I after stunning by means of gas injected into the cranial cavity or killed by the same method or I by laceration after stunning of central nervous tissue by means of an elongated rod-shaped introduced into the cranial cavity;
	(i			nd-bone meal or greaves derived from ruminants, as defined in the Terrestrial Animal Health Code
<sup>(1)</sup> either	[(b) th			ld Organisation for Animal Health; o not contain and are not derived from:
enner	(i) (i	-		isk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;
	(i.	-		lly separated meat obtained from bones of bovine, ovine and caprine animals;
	(:			d lymphotic ticgues symposed dyning the deboning process 111

nervous and lymphatic tissues exposed during the deboning process.]]]

(iii)

TED K	INGDOM				II.a Certificate reference
	(1) and/or	[(b)	continu	at products contain and are derived from treated intestines sociously reared and slaughtered in a country or region classified or region posing a negligible BSE risk in which there have be	l in accordance with Decision 2007/453/EC as a
	(1) and/or	[(b)	the mea	at products contain and are derived from treated intestines sourt or region classified in accordance with Decision 2007/453/E which there has been at least one BSE indigenous case, and:	urced from animals which originate from a
7	•	(1) either	[(i)	the animals were born after the date from which the ban on meal and greaves derived from ruminants has been enforced	
	<b>♦</b>	(1) and/or	[(ii)	the treated intestines of bovine, ovine and caprine animal or specified risk material as defined in point 1 of Annex V to I	•
•	(1) [II.1.11	_	_	terial from domestic solipeds, the fresh meat used in the prep ds which immediately prior to the date of their slaughter had b	1
	(1) eithe	anothe	<b>\</b>	nonths in the third country of slaughter, if born in that third c untry which is listed for the concerned animals and products i	•
	$^{(1)}or$			ntry of slaughter, since birth, if slaughtered at an age of less t	<del>-</del>
	<sup>(1)</sup> or	soliped	ds for foo	ntry of slaughter for six months or less if they entered that this production, and	ird country from a Member Stateas domestic
				y or territory of slaughter in which:	
		(a)		ninistration to domestic solipeds of:	
			(i) (ii)	substances listed in Table 2 of the Annex to Commission Re thyrostatic substances, stilbenes, stilbene derivatives, their s derivatives is prohibited;	
			(iii)	other substances having oestrogenic, androgenic or gestages for:	nic action and of beta-agonists is only allowed
		_ (1)	either	[therapeutic treatment as defined in Article 1(2), point (b), conformity with Article 4(2) of that Directive,]	of Council Directive 96/22/EC, where applied
		_ (1)	or	[zootechnical treatment as defined in Article 1(2), poin conformity with Article 5 of that Directive,]	at (c), of Directive 96/22/EC, where applied
		(b)	by the	nestic solipeds fulfilled, at least curing the six months prior to control plan submitted in accordance with the Article 6(2) of ned animals and products are listed in Annex -I to Implementationary or territory.]	Delegated Regulation (EU) 2022/2292 and the
	(1) [II.1.12	. (1) (11)	either	[if containing material from farmed cervidae, the product of excluding offal and spinal cord, of farmed cervid animals we disease by histopathology, immunohistochemistry or other authorities with negative results and is not derived from an idisease has been confirmed or is officially suspected.]]	hich have been examined for chronic wasting diagnostic method recognised by the competer
		(1) (12)	or	[if containing material from wild cervidae, the product cont excluding offal and spinal cord, of wild cervid animals white disease by histopathology, immunohistochemistry or other cauthorities with negative results and is not derived from ani wasting disease has been confirmed in the last three years p health/official certificate or is officially suspected.]]	ch have been examined for chronic wasting diagnostic method recognised by the competen mals coming from a region where chronic
(	<sup>1) (13)</sup> [ <b>II.1.a.</b>		ntion as remeat prod	egards Commission Delegated Regulation (EU) 2023/905 [ucts]	Delete when the Union is not the final destinat
		and of render Part I v have n medici infection Regula	the Counted animal were product been ad animal productions in humation (EU	ed, declare that I am aware of the relevant requirements of Recil and Commission Delegated Regulation (EU) 2023/905 and fats and greaves, meat extracts and treated stomachs, bladder uced in accordance with these requirements, and in particular dministered antimicrobial medicinal products for growth pronects containing an antimicrobial that is included in the list of a mans laid down in Commission Implementing Regulation (EU) 2023/905 and originate from a third country or region therecolation (EU) 2023/905.]	d hereby certify that the meat products, includers and intestines other than casings described in the theat is derivative, that the animals from which the meat is derivation or yield increase or antimicrobial antimicrobials reserved for the treatment of certification. We will be set to the treatment of the control of

other than ungulates and leporidae]

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II.a Certificate reference

		at product, including rendered animal fats and greaves, meat extracts and treated stomachs, bladders and intestines others than described in Part I:
	II.2.	1. has been processed in and dispatched from the <b>zone</b> with code:
	(i) either [II.2	2.2. has been processed from fresh meat from <b>only one species of animals</b> , with code <sup>(4)</sup> , and the fresh meat used for the processing of the meat product has undergone the specific treatment <sup>(5)</sup> , which is specifically assigned in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404 to the species of origin of the fresh meat and to the zone referred to in point II.2.1. and has been obtained from animals originating from:
1	(1) eithe	[the zone referred to in point II.2.1.;]]
	(I) or	[the zone with code
		(1) either [Part 1 of Annex XIII to Implementing Regulation (EU) 2021/404, in case of fresh meat of ungulates;]]] (7)
	(1) or	[Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404, in case of fresh meat of poultry and game birds;]]] [a Member State;]]
	(1) or [II.2	
	<i>о</i> г (п.2	of fresh meat of poultry where there has been a case or an outbreak of highly pathogenic avian influenza or infection with Newcastle disease virus and the fresh meat used for the processing of the meat product has undergone at least the specific treatment "D" [5];
	(1) or [II.2	2.2. has been processed mixing fresh meat from different species of animals, with codes,, (4), and such fresh
	(1) eithe	meat:  er [II.2.2.1. has been <b>mixed before the final treatment</b> and, after mixing, has undergone the specific treatment <sup>(5)</sup> , as it
	· euni	is the most severe of the treatments specifically assigned in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404 to the different species of origin of the fresh meat and to the zone referred to in point II.2.1., and has been obtained from animals originating from:
		(1) either [the zone referred to in point II.2.1.;]]
		(1) or [the zone with:
		[code
		[code
		(1) or [a Member State;]]
	<sup>(1)</sup> or [II.2.2	.1. has been <b>mixed after the final treatment</b> and, before the mixing, has undergone the specific treatment(s)
		originating from:
		(1) either [the zone referred to in point II.2.1.;]]
		(1) or [the zone with: (1) [code
		XIII to Implementing Regulation (EU) 2021/404 for entry into the Union of fresh meat of the species from which the meat product has been processed;] (7)
		[code6] which, at the date of issue of this animal health/official certificate, is listed in Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404 for entry into the Union of fresh meat of the species from which the meat product has been processed;]]]
		(1) or [a Member State.]]
	(1) or [II.2	2.2. has:
		(a) been processed from fresh meat from <b>one species of animals or mixing fresh meat from different species of animals</b> , with codes,,(4);
		been processed from fresh meat obtained from animals originating from the zone/s with code/s,,,
		(c) undergone the specific treatment "B" (5);]
	II.2.3.	has been processed from fresh meat obtained from:

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[animals kept in an establishment that was not subject to national restriction measures for animal health reasons, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species of origin of the meat products and emerging diseases at the date of dispatch of the animals to the slaughterhouse and in and around the establishment, in an area of 10 km radius, including where appropriate the territory of a neighbouring country, such diseases have not been reported during the last 30 days prior to the date of slaughter of the animals;]

[wild animals which originate from a place in and round which, in an area of 10 km radius, including where appropriate the territory of a neighbouring country, none of the listed diseases reffered to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species of origin of the meat products and emerging diseases, has been reported during the last 30 days prior to the date of killing of the animals;],

II.2.4. after processing, has been handled until packaging in a way to prevent cross contamination that could introduce animal health risk;

is intended for a Member State or zone thereof which has been granted the status free from infection with Newcastle disease virus without vaccination in accordance with Commission Delegated Regulation (EU) 2020/689, and has been obtained from poultry that have not been vaccinated against infection with Newcastle disease virus with a live vaccine during the last 30 days prior to the date of their slaughter.]

## II.3. Animal welfare attestation [Delete when the Union is not the final destination]

I, the undersigned official veterinarian, hereby certify, that the meat products described in Part I derive from animals which have been treated in the slaughterhouse in accordance with the requirements of the Union legislation on the protection of animals at the time of killing or at least equivalent requirements.

#### Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health/official certificate include the United Kingdom in respect of Northern Ireland.

This animal health/official certificate is intended for the entry into the Union of meat products from zones not authorised to enter fresh meat of the relevant species and therefore are required to undergo a specific risk-mitigating treatment, including when the Union is not the final destination of such meat products.

This animal health/official certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

## Part I:

Box reference I.27.: Description of consignment:

"Slaughterhouse": Slaughterhouse or game handling establishment

## Part II:

- (1) Delete if not applicable.
- (2) Meat product as defined in point 7.1 of Annex I to Regulation (EC) No 853/2004.
- (3) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404.
- BOV= domestic bovine animals; OVI= domestic ovine animals and caprine animals; POR= domestic porcine animals; RUF= animals of the family *Bovidae* (other than domestic bovine, ovine and caprine animals), camelid animals and cervid animals kept as farmed game; RUW= wild animals of the family *Bovidae* (other than domestic bovine, ovine and caprine animals), wild camelid animals and wild cervid animals; SUF= animals kept as farmed game of wild breeds of porcine animals and animals of the family *Tayassuidae*; SUW= wild animals of wild breeds of porcine animals and animals of the family *Tayassuidae*; POU= poultry other than ratites; RAT= ratites; GB= game birds.
- Treatment as defined in Annex XXVI to Delegated Regulation (EU) 2020/692.
- Code of the zone in accordance with column 2 of the table in Part 1 of Annex XIII or column 2 of the table in Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404.
- Not for zones with the entry related to specific conditions "Maturation, pH and de-boning" in column 5 of the table in Part 1 of Annex XIII to Implementing Regulation (EU) 2021/404.
- (8) Specify the combination of treatments referred to in note (5) and species set out in note (4), as follows: letter of treatment code(s) of species (X-YYY, X-YYY, X-YYY).
- (9) This guarantee is required only for consignments intended for a Member State or zone thereof which has been granted the status free from infection with Newcastle disease virus without vaccination in accordance with Delegated Regulation (EU) 2020/689.
- The derogation for domestic porcine animals coming from a holding officially recognised as applying controlled housing conditions, may only be applied in countries listed in Annex VII to Implementing Regulation (EU) 2015/1375.
- Applicable when the meat has been obtained from a country mentioned in Chapter F, point 1, of Annex IX to Regulation (EC) No 999/2001.
- Applicable when the meat has been obtained from a country mentioned in Chapter F, point 2, of Annex IX to Regulation (EC) No 999/2001.
- (13) Applicable to consignments entering the Union as from 3 September 2026.

## Certificate model MPST

NITED KINGDOM	II.a Certificate reference	
Official veterinarian		
Name (in capital letters)		
Date	Qualification and title	
7		
Stamp	Signature	
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170		
<b>40</b>		
U.C		
O		
	0%	