

Part I : Details of consignment	I.1. Consignor Name Address Country <span style="float: right;">ISO Code</span>		I.2. IMSOC Reference <b>Specimen not to be used for exports from EU</b> I.2.a. Local Reference	
	I.5. Consignee Name Address Country <span style="float: right;">ISO Code</span>		I.3. Central competent authority I.4. Local competent authority	
	I.7. Country of origin <span style="float: right;">ISO Code</span>		I.9. Country of destination <span style="float: right;">ISO Code</span>	
	I.8. Region of origin <span style="float: right;">Code</span>		I.10. Region of destination <span style="float: right;">Code</span>	
	I.11. Place of Dispatch Name Address Approval Number Country <span style="float: right;">ISO Code</span>		I.12. Place of destination Name Address Approval Number Country <span style="float: right;">ISO Code</span>	
	I.13. Place of Loading Name Address Approval Number Country <span style="float: right;">ISO Code</span>		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 Ambient <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>		I.17. Accompanying documents Commercial document reference <span style="float: right;">Date of issue</span> Country <span style="float: right;">Place of issue</span>	
	I.19. Container No / Seal No			
I.20. Certified as Transhumance <input type="checkbox"/> Pollination <input type="checkbox"/> Animal Feedingstuff <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Game Restocking <input type="checkbox"/> Fattening <input type="checkbox"/> Registered equidae <input type="checkbox"/> Quarantine <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Training <input type="checkbox"/> Further process <input type="checkbox"/> Racing <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Competition <input type="checkbox"/> Storage <input type="checkbox"/> Breeding <input type="checkbox"/> Ornamental use/research <input type="checkbox"/> Laboratory <input type="checkbox"/> Rodent food <input type="checkbox"/> Pets <input type="checkbox"/> Other <input type="checkbox"/> Production <input type="checkbox"/> Technical use <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Relaying <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved Bodies <input type="checkbox"/> Human consumption <input type="checkbox"/> Pet food <input type="checkbox"/> Sales <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Circus exhibition <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>				
I.21. For transit through a third country <input type="checkbox"/> Country <span style="float: right;">ISO Code</span> EU Exit Authority <span style="float: right;">BCP code</span> EU Entry Authority <span style="float: right;">BCP code</span>		I.22. For transit through Member State(s) <input type="checkbox"/> Country <span style="float: right;">ISO Code</span>		
I.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	I.25. Total gross weight	

Part I : Details of consignment	I.28. Description of consignment					
	<b>1. 30 PHARMACEUTICAL PRODUCTS</b>					
	<b>3002</b> Human blood; animal blood prepared for therapeutic, prophylactic or diagnostic uses; antisera and other blood fractions and modified immunological products, whether or not obtained by means of biotechnological processes; vaccines, toxins, cultures of micro-organisms (excluding yeasts) and similar products					
	Antisera, other blood fractions and immunological products, whether or not modified or obtained by means of biotechnological processes					
	<b>300219</b> Other					
	<b>30021900</b> Other					
	Commodity		Species	Quantity	Batch number	Manufacturing plant
	Cold store		Cutting plant	Date of freezing	Date of production	Date of slaughter
	Net weight		Product Description	Package count	Identification mark	
	SPECIMEN					

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 8 (c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto, and certify that:</p> <p>II.1. the blood products described above consist of blood products that satisfy the requirements below;</p> <p>II.2. they consist exclusively of blood products not intended for human or animal consumption;</p> <p>II.3. they have been prepared and stored in a plant supervised by the competent authority, exclusively with the following animal by-products:</p> <p>(2) <input type="checkbox"/> either [- blood of slaughtered animals, which is fit for human consumption in accordance with retained EU law, but is not intended for human consumption for commercial reasons;]</p> <p>(2) <input type="checkbox"/> and/or [- blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with retained EU law, but which did not show any signs of diseases communicable to humans or animals, derived from carcasses that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]</p> <p>(2) <input type="checkbox"/> and/or [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]</p> <p>(2) <input type="checkbox"/> and/or [- blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals;]</p> <p>(2) <input type="checkbox"/> and/or [- blood and blood products derived from the production of products intended for human consumption;]</p> <p>(2) <input type="checkbox"/> and/or [- animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1 (2)(d) of Council Directive 96/22/EC or Article 2(b) of Council Directive 96/23/EC;]</p> <p>(2) <input type="checkbox"/> and/or [- animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by retained EU law or, in the absence thereof, in national legislation;]</p> <p>II.4. the blood that these products were manufactured from has been collected in slaughterhouses approved in accordance with retained EU law, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection.</p> <p>(2) <input type="checkbox"/> II.5. In the case of blood products derived from Artiodactyla, Perissodactyla and Proboscidea including their crossbreeds, other than Suidae and Tayassuidae, the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue:</p> <p>(2) <input type="checkbox"/> either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [change in pH to pH 5 for two hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check]]</p> <p>(2) <input type="checkbox"/> II.6. In the case of blood products derived from Suidae, Tayassuidae, poultry and other avian species, the products have undergone one of the following treatments guaranteeing the absence of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease and highly pathogenic avian influenza, as appropriate to the species:</p> <p>(2) <input type="checkbox"/> either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]</p>		

Part II: Certification	II. Health information			
	(2)	<input type="checkbox"/> and/or	[heat treatment of at least 80°C for Suidae/Tayassuidae (2) and at least 70°C for poultry and other avian species (2) throughout the substance of the product, followed by an effectiveness check]].	
	(2)	<input type="checkbox"/> [II.7.	In the case of blood products derived from species other than those listed in point II.5 or II.6, the products have undergone the following treatment (please specify): _____.]	
	II.8.	The products were:		
	(2)	<input type="radio"/> either	[packed in new or sterilised bags or bottles,]	
	(2)	<input type="radio"/> or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;] and the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';	
	II.9.	the products were stored in enclosed storage;		
	II.10.	all precautions were taken to avoid the contamination of the products with pathogenic agents after treatment;		
	(2)	<input type="checkbox"/>	The treated blood products described above	
	[II.11.			
(2)	<input type="radio"/> either	[is derived from other ruminants than bovine, ovine or caprine animals.]]		
(2)	<input type="radio"/> or	[is derived from bovine, ovine or caprine animals and does not contain and is not derived from:		
(2)	<input type="radio"/> either	[bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]		
(2)	<input type="radio"/> or	(a) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council;		
		(b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC, which there has been no indigenous BSE case,		
		(c) animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]		

Part II: Certification	II. Health information		
	<p>Notes</p> <p>(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.</p> <p>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).</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man.</p> <p>Part I:</p> <ul style="list-style-type: none"> <li>- Box reference I.6: Person responsible for the consignment 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 Great Britain, Channel Islands or Isle of Man.</li> <li>- Box reference I.11 and I.12: Approval number; the registration number of the establishment or plant, which has been issued by the competent authority.</li> <li>- Box reference I.12: Place of destination; this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.</li> <li>- 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 in Great Britain, Channel Islands or Isle of Man, the consignor must inform the BCP of entry into Great Britain, Channel Islands or Isle of Man.</li> <li>- Box reference I.16: Do not use this box until the end of the transitional staging period.</li> <li>- Box reference I.19: use the appropriate Harmonized System (HS) code under the following headings: 05.11, 30.02, 35.02 or 35.04.</li> <li>- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.</li> <li>- 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.</li> <li>- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.</li> <li>- Box reference I.28 in case of Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilian.</li> </ul> <p>Part II:</p> <ul style="list-style-type: none"> <li>(2) Delete as appropriate.</li> <li>- The signature and the stamp must be in a different colour to that of the printing.</li> <li>- 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 of Great Britain, Channel Islands or Isle of Man.</li> </ul>		
Certifying Officer			

<b>Part II: Certification</b>	II. Health information			
	Name (in capital letters)	Qualification and title		
	Date of signature	Signature		
	Stamp			
<b>SPECIMEN</b>				

Část I	I.1. Odesílatel Název Adresa Země <span style="float: right;">Kód ISO</span>		I.2. Referenční číslo IMSOC <b>Specimen not to be used for exports from EU</b> I.2.a. Local Reference																
	I.5. Příjemce Název Adresa Země <span style="float: right;">Kód ISO</span>		I.3. Ústřední příslušný orgán I.4. Local competent authority																
	I.7. Země původu <span style="float: right;">Kód ISO</span>		I.9. Country of destination <span style="float: right;">Kód ISO</span>																
	I.8. Region of origin <span style="float: right;">Kód</span>		I.10. Region určení <span style="float: right;">Kód</span>																
	I.11. Place of Dispatch Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>		I.12. Místo určení Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>																
	I.13. Místo nakládky Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>		I.14. Date and time of departure																
	I.15. Dopravní prostředky		I.16 Entry Point																
	<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 20%;">Typ</th> <th style="width: 20%;">Doklad</th> <th style="width: 60%;">Identifikace</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </tbody> </table>		Typ	Doklad	Identifikace														
	Typ	Doklad	Identifikace																
I.18. Transport conditions Okolní <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Zmrazené <input type="checkbox"/> Chlazený <input type="checkbox"/>		I.17. Průvodní doklady Referenční číslo obchodního do k ladu Datum vydání Země Místo vydání																	
I.19. Č. kontejneru / č. plomby																			
I.20. Certified as Transhumance <input type="checkbox"/> Pollination <input type="checkbox"/> Krmivo pro zvířata <input type="checkbox"/> Použití pro farmaceutické účely <input type="checkbox"/> Game Restocking <input type="checkbox"/> Výkrmová <input type="checkbox"/> Evidování koňovité <input type="checkbox"/> Karanténa <input type="checkbox"/> Umělé rozmnožování <input type="checkbox"/> Training <input type="checkbox"/> Další postup <input type="checkbox"/> Racing <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Competition <input type="checkbox"/> Storage <input type="checkbox"/> Breeding <input type="checkbox"/> Ornamental use/research <input type="checkbox"/> Laboratory <input type="checkbox"/> Rodent food <input type="checkbox"/> Zvířata v zájmovém chovu <input type="checkbox"/> Jiné <input type="checkbox"/> Production <input type="checkbox"/> Technické použití <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Sádkování <input type="checkbox"/> Porážka <input type="checkbox"/> Schválené orgány <input type="checkbox"/> Lidská spotřeba <input type="checkbox"/> Krmivo pro domácí zvířata <input type="checkbox"/> Sales <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Cirkus/výstava <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>																			
I.21. For transit through a third country <input type="checkbox"/> Země <span style="float: right;">Kód ISO</span> EU Exit Authority <span style="float: right;">BCP code</span> EU Entry Authority <span style="float: right;">BCP code</span>		I.22. For transit through Member State(s) <input type="checkbox"/> Země <span style="float: right;">Kód ISO</span>																	
I.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	I.25. Celková hrubá hmotnost																

I.28. Description of consignment

**1. 30 FARMACEUTICKÉ VÝROBKY**

**3002** Lidská krev; zvířecí krev připravená k terapeutickým, profylaktickým nebo diagnostickým účelům; antiséra a ostatní krevní složky a modifikované imunologické výrobky, též získané biotechnologickými procesy; očkovací látky, toxiny, kultury mikroorganismů (kromě kvasinek) a podobné výrobky

Antiséra, ostatní krevní složky a imunologické výrobky, též modifikované nebo získané biotechnologickými procesy

**300219** Ostatní

**30021900** Ostatní

Komodita	Druh	Množství	Číslo šarže	Výrobní zařízení
Chladírenské zařízení	Bourárna	Datum zmrazení	Datum výroby	Datum porážky
Čistá hmotnost	Product Description	Počet balení	Identifikační značka	

Část I

SPECIMEN

Part II: Certification	II. Informace týkající se zdraví		
	<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 8 (c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto, and certify that:</p> <p>II.1. the blood products described above consist of blood products that satisfy the requirements below;</p> <p>II.2. they consist exclusively of blood products not intended for human or animal consumption;</p> <p>II.3. they have been prepared and stored in a plant supervised by the competent authority, exclusively with the following animal by-products:</p> <p>(2) <input type="checkbox"/> either [- blood of slaughtered animals, which is fit for human consumption in accordance with retained EU law, but is not intended for human consumption for commercial reasons;]</p> <p>(2) <input type="checkbox"/> and/or [- blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with retained EU law, but which did not show any signs of diseases communicable to humans or animals, derived from carcasses that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]</p> <p>(2) <input type="checkbox"/> and/or [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]</p> <p>(2) <input type="checkbox"/> and/or [- blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals;]</p> <p>(2) <input type="checkbox"/> and/or [- blood and blood products derived from the production of products intended for human consumption;]</p> <p>(2) <input type="checkbox"/> and/or [- animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1 (2)(d) of Council Directive 96/22/EC or Article 2(b) of Council Directive 96/23/EC;]</p> <p>(2) <input type="checkbox"/> and/or [- animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by retained EU law or, in the absence thereof, in national legislation;]</p> <p>II.4. the blood that these products were manufactured from has been collected in slaughterhouses approved in accordance with retained EU law, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection.</p> <p>(2) <input type="checkbox"/> II.5. In the case of blood products derived from Artiodactyla, Perissodactyla and Proboscidea including their crossbreeds, other than Suidae and Tayassuidae, the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue:</p> <p>(2) <input type="checkbox"/> either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [change in pH to pH 5 for two hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check]]</p> <p>(2) <input type="checkbox"/> II.6. In the case of blood products derived from Suidae, Tayassuidae, poultry and other avian species, the products have undergone one of the following treatments guaranteeing the absence of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease and highly pathogenic avian influenza, as appropriate to the species:</p> <p>(2) <input type="checkbox"/> either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]</p> <p>(2) <input type="checkbox"/> and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]</p>		

Part II: Certification	II. Informace týkající se zdraví			
	(2)	<input type="checkbox"/> and/or	[heat treatment of at least 80°C for Suidae/Tayassuidae (2) and at least 70°C for poultry and other avian species (2) throughout the substance of the product, followed by an effectiveness check]].	
	(2)	<input type="checkbox"/> [II.7.	In the case of blood products derived from species other than those listed in point II.5 or II.6, the products have undergone the following treatment (please specify): _____.]	
	II.8.	The products were:		
	(2)	<input type="radio"/> either	[packed in new or sterilised bags or bottles,]	
	(2)	<input type="radio"/> or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;]	and the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';
	II.9.	the products were stored in enclosed storage;		
	II.10.	all precautions were taken to avoid the contamination of the products with pathogenic agents after treatment;		
	(2)	<input type="checkbox"/>	The treated blood products described above	
	[II.11.			
	(2)	<input type="radio"/> either	[is derived from other ruminants than bovine, ovine or caprine animals.]]	
	(2)	<input type="radio"/> or	[is derived from bovine, ovine or caprine animals and does not contain and is not derived from:	
(2)	<input type="radio"/> either	[bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]		
(2)	<input type="radio"/> or	[(a) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council;	(b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC, which there has been no indigenous BSE case,	
	(c)	animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]		

II. Informace týkající se zdraví

## 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 consignment 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 Great Britain, Channel Islands or Isle of Man.
- Box reference I.11 and I.12: Approval number; the registration number of the establishment or plant, which has been issued by the competent authority.
- Box reference I.12: Place of destination; this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- 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 in Great Britain, Channel Islands or Isle of Man, the consignor must inform the BCP 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 Harmonized System (HS) code under the following headings: 05.11, 30.02, 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 in case of Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilian.

## Part II:

- (2) Delete as appropriate.
- 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 of Great Britain, Channel Islands or Isle of Man.

Certifying Officer

Part II: Certification

<b>Part II: Certification</b>	II. Informace týkající se zdraví			
	Name (in capital letters) Datum podpisu Razítko	Qualification and title Podpis		
	<b>SPECIMEN</b>			