EUROPEAN UNION **Export Health Certificate**

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority

I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

I.10. Region of destination Code

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Ambient ☐ Controlled temperature ☐ Frozen ☐ Chilled ☐

I.17. Accompanying documents
Commercial document reference Date of issue
Country Place of issue

I.19. Container No / Seal No

I.20. Certified as

Transhumance ☐	Pollination ☐	Animal Feedingstuff ☐	Pharmaceutical use ☐
Game Restocking ☐	Fattening ☐	Registered equidae ☐	Quarantine ☐
Artificial reproduction ☐	Training ☐	Further process ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Pets ☐	Other ☐	Production ☐
Technical use ☐	Ornamental bird food ☐	Relaying ☐	Slaughter ☐
Approved Bodies ☐	Human consumption ☐	Pet food ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Circus exhibition ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

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

1. 30 PHARMACEUTICAL PRODUCTS

3002 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

300212 Antisera and other blood fractions

30021200 Antisera and other blood fractions

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

Part II: Certification

II. Health information

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 (d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011 and in particular Chapter IV of Annex XIII thereto, and certify that the blood or blood products of equidae described above:

II.1. consist of blood or blood products from equidae that satisfy the health requirements below;

II.2. consist exclusively of blood or blood products of equidae not intended for human or animal consumption;

II.3. have been obtained from animals that originate from the EU Member States or from a third country, territory or part thereof listed in the column "third countries lists" of row No 3 of Table 2 in Section 1 of Chapter II of Annex XIV to Regulation (EU) No 142/2011 where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders (Burkholderia mallei), equine encephalomyelitis (all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies, anthrax;

II.4. have been derived from blood from equidae which was collected under the supervision of a veterinarian in slaughterhouses approved in accordance with Regulation (EC) No 853/2004 of the European Parliament and of the Council, in slaughterhouses approved and supervised by the competent authority of the country of collection and in facilities approved and supervised by the competent authority of the country of collection for the purpose of collecting blood from equidae for the production of blood products for purposes other than feeding for farmed animals;

II.5. have been derived from blood which was collected from equidae:

II.5.1. which on inspection on the date of blood collection did not show clinical signs of any of the compulsorily notifiable diseases listed in Annex I to Council Directive 2009/156/EC , and of equine influenza, equine piroplasmosis, equine rhinopneumonitis and equine viral arteritislisted in point 4 of Article 1.2.3 of the Terrestrial Animal Health Code of the World Organisation for Animal Health (0IE), 2010 edition;

II.5.2. which have been kept for at least 30 days prior to the date of and during blood collection on holdings under veterinary supervision which were not subject to a prohibition order pursuant to Article 4(5) or restrictions for African horse sickness in accordance with Article 5 of Directive 2009/156/EC;

II.5.3. which had no contact with equidae from a holding which was subject to a prohibition order for animal health reasons pursuant to Article 4(5) of Directive 2009/156/EC;

II.5.4. for which the period for the prohibition order referred to in points II.5.2. and II.5.3 has been determined as follows:

(2) ○ either [not all the animals of species susceptible to the disease located on the holding have been slaughtered , in which case the period of prohibition must be at least:

— six months in the case of glanders (Burkholderia mallei), beginning on the date on which the equidae infected with the disease are slaughtered,

— six months in the case of equine encephalomyelitis of any type, including Venezuelan equine encephalomyelitis, beginning on the date on which the equidae infected with the disease are slaughtered, in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, and the remaining animals have shown a negative reaction to two Coggins tests carried out three months apart,

— six months from the date of the last recorded case of vesicular stomatitis,

— one month from the date of the last recorded case of rabies,

— 15 days from the date of the last recorded case of anthrax;]

(2) ○ or [all the animals of species susceptible to the disease located on the holding have been slaughtered and the premises were disinfected, in which case the period of prohibition must be 30 days beginning on the date on which the animals were slaughtered, and the premises disinfected, except in the case of anthrax, where the period of prohibition shall be 15 days;]

Part II: Certification

II. Health information

II.6. blood products come from an establishment or plant approved or registered by the competent authority of the third country meeting the specific conditions set out in Article 23 or 24 of Regulation (EC) No 1069/2009;

II.7. blood products have been produced from blood which fulfils the conditions referred in II.4 and II.5 and

(2) ○ either [has been collected from equidae which have been kept for a period of at least three months, or since birth if less than three months old, prior to the date of collection on holdings under veterinary supervision in the country of collection which during that period and the period of blood collection has been free of:

(a) African horse sickness for two years;

(b) Venezuelan equine encephalomyelitis for a period of at least two years;

(c) glanders

(2) ○ either [for a period of three years;]

(2) ○ or [for a period of six months where the animals have passed the post-mortem inspection for glanders in the slaughterhouse referred to in II.4, including a careful examination of mucous membranes from the trachea, larynx, nasal cavities and sinuses and their ramifications, after splitting the head in the median plane and excising the nasal septum;]

(d) in the case of blood products other than serum and plasma, vesicular stomatitis for six months;]]

(2) ○ or [has been subjected to at least one of the following treatments, followed by an effectiveness check, for the inactivation of possible causative pathogens for African horse sickness, equine encephalomyelitis of all types including Venezuelan equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and glanders (Burkholderia mallei):

(2) ☐ either [heat treatment at a temperature of 65°C for at least three hours;]

(2) ☐ and/or [irradiation at 25 kGy by gamma rays;]

(2) ☐ and/or [change in pH to pH 5 for two hours;]

(2) ☐ and/or [heat treatment of at least 80°C throughouttheir substance;]]

II.8. all precautions have been taken to avoid contamination of the blood and blood products with pathogenic agents during production, handling and packaging;

II.9. blood and blood products were packed in sealed impermeable containers clearly labelled "NOT FOR HUMAN OR ANIMAL CONSUMPTION" and bearing:

(a) in the case of blood, the approval number of the establishment of collection;

(b) in the case of blood products, the approval number of the establishment of production;

II.10. the products were stored in enclosed storage.

Part II: Certification

II. Health information

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 to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.

— 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 transit commodity. The products in transit can 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 case of unloading and reloading, the consignor must inform the BCP of entry into Great

— 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 heading: 30.02.

— 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 for animal consumption.

— Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

— Box reference I.28:

(a) Manufacturing plant:

(i) in the case of blood, provide the approval number of the registered establishment of collection;

(ii) in the case of blood products, provide the approval number of the establishment of production;

(b) Species: select amongst the following: Equus cabalus, Equus asinus, Equus cabalus*asinus.

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.

Certifying Officer

Part II: Certification

II. Health information		
Name (in capital letters) Date of signature Stamp	Qualification and title Signature	

Část I

I.1. Odesílatel
Název
Adresa
Země Kód ISO

I.2. Referenční číslo IMSOC
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Příjemce
Název
Adresa
Země Kód ISO

I.3. Ústřední příslušný orgán

I.4. Local competent authority

I.7. Země původu Kód ISO

I.9. Country of destination Kód ISO

I.8. Region of origin Kód

I.10. Region určení Kód

I.11. Place of Dispatch
Název
Adresa
Číslo schválení
Země Kód ISO

I.12. Místo určení
Název
Adresa
Číslo schválení
Země Kód ISO

I.13. Místo nakládky
Název
Adresa
Číslo schválení
Země Kód ISO

I.14. Date and time of departure

I.15. Dopravní prostředky

Typ	Doklad	Identifikace

I.16 Entry Point

I.18. Transport conditions
Okolní ☐ Controlled temperature ☐ Zmrazené ☐ Chlazený ☐

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 ☐	Pollination ☐	Krmivo pro zvířata ☐	Použití pro farmaceutické účely ☐
Game Restocking ☐	Výkrmová ☐	Evidovaní koňovití ☐	Karanténa ☐
Umělé rozmnožování ☐	Training ☐	Další postup ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Zvířata v zájmovém chovu ☐	Jiné ☐	Production ☐
Technické použití ☐	Ornamental bird food ☐	Sádkování ☐	Porážka ☐
Schválené orgány ☐	Lidská spotřeba ☐	Krmivo pro domácí zvířata ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Cirkus/výstava ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Země Kód ISO
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Země Kód ISO

I.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	I.25. Celková hrubá hmotnost

Část I

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

300212 Antiséra a jiné krevní složky

30021200 Antiséra a jiné krevní složky

Komodita	Druh	Množství	Číslo šarže	Výrobní zařízení

Chladírenské zařízení	Bourárna	Datum zmrazení	Datum výroby	Datum porážky

Čistá hmotnost	Product Description	Počet balení	Identifikační značka

Part II: Certification

II. Informace týkající se zdraví

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 (d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011 and in particular Chapter IV of Annex XIII thereto, and certify that the blood or blood products of equidae described above:

II.1. consist of blood or blood products from equidae that satisfy the health requirements below;

II.2. consist exclusively of blood or blood products of equidae not intended for human or animal consumption;

II.3. have been obtained from animals that originate from the EU Member States or from a third country, territory or part thereof listed in the column "third countries lists" of row No 3 of Table 2 in Section 1 of Chapter II of Annex XIV to Regulation (EU) No 142/2011 where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders (Burkholderia mallei), equine encephalomyelitis (all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies, anthrax;

II.4. have been derived from blood from equidae which was collected under the supervision of a veterinarian in slaughterhouses approved in accordance with Regulation (EC) No 853/2004 of the European Parliament and of the Council, in slaughterhouses approved and supervised by the competent authority of the country of collection and in facilities approved and supervised by the competent authority of the country of collection for the purpose of collecting blood from equidae for the production of blood products for purposes other than feeding for farmed animals;

II.5. have been derived from blood which was collected from equidae:

II.5.1. which on inspection on the date of blood collection did not show clinical signs of any of the compulsorily notifiable diseases listed in Annex I to Council Directive 2009/156/EC , and of equine influenza, equine piroplasmosis, equine rhinopneumonitis and equine viral arteritislisted in point 4 of Article 1.2.3 of the Terrestrial Animal Health Code of the World Organisation for Animal Health (0IE), 2010 edition;

II.5.2. which have been kept for at least 30 days prior to the date of and during blood collection on holdings under veterinary supervision which were not subject to a prohibition order pursuant to Article 4(5) or restrictions for African horse sickness in accordance with Article 5 of Directive 2009/156/EC;

II.5.3. which had no contact with equidae from a holding which was subject to a prohibition order for animal health reasons pursuant to Article 4(5) of Directive 2009/156/EC;

II.5.4. for which the period for the prohibition order referred to in points II.5.2. and II.5.3 has been determined as follows:

(2) ○ either [not all the animals of species susceptible to the disease located on the holding have been slaughtered , in which case the period of prohibition must be at least:

— six months in the case of glanders (Burkholderia mallei), beginning on the date on which the equidae infected with the disease are slaughtered,

— six months in the case of equine encephalomyelitis of any type, including Venezuelan equine encephalomyelitis, beginning on the date on which the equidae infected with the disease are slaughtered, in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, and the remaining animals have shown a negative reaction to two Coggins tests carried out three months apart,

— six months from the date of the last recorded case of vesicular stomatitis,

— one month from the date of the last recorded case of rabies,

— 15 days from the date of the last recorded case of anthrax;]

(2) ○ or [all the animals of species susceptible to the disease located on the holding have been slaughtered and the premises were disinfected, in which case the period of prohibition must be 30 days beginning on the date on which the animals were slaughtered, and the premises disinfected, except in the case of anthrax, where the period of prohibition shall be 15 days;]

Part II: Certification

II. Informace týkající se zdraví

II.6. blood products come from an establishment or plant approved or registered by the competent authority of the third country meeting the specific conditions set out in Article 23 or 24 of Regulation (EC) No 1069/2009;

II.7. blood products have been produced from blood which fulfils the conditions referred in II.4 and II.5 and

(2) ○ either [has been collected from equidae which have been kept for a period of at least three months, or since birth if less than three months old, prior to the date of collection on holdings under veterinary supervision in the country of collection which during that period and the period of blood collection has been free of:

(a) African horse sickness for two years;

(b) Venezuelan equine encephalomyelitis for a period of at least two years;

(c) glanders

(2) ○ either [for a period of three years;]

(2) ○ or [for a period of six months where the animals have passed the post-mortem inspection for glanders in the slaughterhouse referred to in II.4, including a careful examination of mucous membranes from the trachea, larynx, nasal cavities and sinuses and their ramifications, after splitting the head in the median plane and excising the nasal septum;]

(d) in the case of blood products other than serum and plasma, vesicular stomatitis for six months;]]

(2) ○ or [has been subjected to at least one of the following treatments, followed by an effectiveness check, for the inactivation of possible causative pathogens for African horse sickness, equine encephalomyelitis of all types including Venezuelan equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and glanders (Burkholderia mallei):

(2) ☐ either [heat treatment at a temperature of 65°C for at least three hours;]

(2) ☐ and/or [irradiation at 25 kGy by gamma rays;]

(2) ☐ and/or [change in pH to pH 5 for two hours;]

(2) ☐ and/or [heat treatment of at least 80°C throughouttheir substance;]]

II.8. all precautions have been taken to avoid contamination of the blood and blood products with pathogenic agents during production, handling and packaging;

II.9. blood and blood products were packed in sealed impermeable containers clearly labelled "NOT FOR HUMAN OR ANIMAL CONSUMPTION" and bearing:

(a) in the case of blood, the approval number of the establishment of collection;

(b) in the case of blood products, the approval number of the establishment of production;

II.10. the products were stored in enclosed storage.

Part II: Certification

II. Informace týkající se zdraví

Notes

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

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— 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 for animal consumption.

— Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

— Box reference I.28:

(a) Manufacturing plant:

(i) in the case of blood, provide the approval number of the registered establishment of collection;

(ii) in the case of blood products, provide the approval number of the establishment of production;

(b) Species: select amongst the following: Equus cabalus, Equus asinus, Equus cabalus*asinus.

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.

Certifying Officer

Part II: Certification

II. Informace týkající se zdraví		
Name (in capital letters) Datum podpisu Razítko	Qualification and title Podpis	