Country **Official certificate to the EU**

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. Certificate reference
I.3. Central competent authority
I.4. Local competent authority

I.2.a. IMSOC reference
Specimen not to be used for imports into the EU

I.5. Consignee
Name
Address
Country ISO Code

I.6. Responsible for the consignment in EU
Name
Address
Country ISO Code

I.7. Country of origin ISO Code	I.8. Region of origin Code	I.9. Country of destination ISO 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 BCP
Authority
Country

I.18. Transport conditions
Ambient ☐ **Chilled** ☐ **Frozen** ☐

I.17. Accompanying documents
Type
Number

I.19. Container No / Seal No

I.20. Certified as
Fattening ☐ **Breeding/production** ☐ **Slaughter** ☐ **Technical Use** ☐ **Pharmaceutical use** ☐ **Game restocking** ☐ **Registered equidae** ☐ **Relaying** ☐ **Quarantine** ☐ **Circus/exhibition** ☐ **Artificial reproduction** ☐ **Further process** ☐ **Approved Bodies** ☐ **Trade samples** ☐ **Canning industry** ☐ **Petfood** ☐ **Production of petfood** ☐ **Manufacture of petfood** ☐ **Animal Feedingstuff** ☐ **Human Consumption** ☐ **Pets** ☐ **Other** ☐

I.21. For transit ☐
Non-EU ISO Code

I.22. For internal market ☐
I.23. For re-entry ☐

I.24. Total number of packages	I.25. Total quantity	I.26. Total net weight	I.26. Total gross weight

I.27. Description of consignment

1. 01 LIVE ANIMALS
0106 Other live animals
Mammals:
010613 Camels and other camelids (Camelidae)
01061300 Camels and other camelids (Camelidae)

Commodity	Species	Identification system	Age	Gender

Part II : Certification

II. Health information

Part II : Certification

II. Health information

II.1. Public health attestation [*to delete when the Union is not the final destination of the animals]

I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:

II.1.1. have not received:

- any stilbene or thyrostatic substances,
- oestrogenic, androgenic, gestagenic or beta-agonist substances for purposes other than therapeutic or zootechnical treatment (as defined in Council Directive 96/22/EC).

II.1.2. fulfil the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Article 29 of Council Directive 96/23/EC and the concerned animals are listed in Commision Decision 2011/163/EU for the concerned country of origin.

II.2. Animal health attestation

I, the undersigned official veterinarian, hereby certify, that the animals described in Part I:

II.2.1. come from the zone with code: _____- _____ (2) which, at the date of issuing this certificate is authorised for entry into the Union of camelid and cervid animals and listed in Part 1 of Annex II to Commission Implementing Regulation (EU) 2021/404.

II.2.2. have remained continuously:

(i) in the zone referred to in point II.2.1. since birth or for a period of time of at least 6 months prior to the date of their dispatch to the Union, and

(ii) in the establishment of origin since birth or for a period of time of at least 40 days prior to the date of their dispatch to the Union, in which no animals have been introduced during that period of time.

II.2.3. had no contact with animals of a lower health status since birth or at least for 6 months prior to the date of their dispatch to the Union.

II.2.4. are not to be killed under a national programme for the eradication of diseases, including the relevant listed diseases referred to in Annex I to Commission Delegated Regulation (EU) 2020/692 and emerging diseases.

II.2.5. have been dispatched directly from the establishment of origin to the Union without passing through any other establishment.

II.2.6. have not been unloaded in any place that does not comply with the requirements laid down in point II.11 since they were dispatched from their establishment of origin until their dispatch to the Union and during that period they have not been in contact with animals of a lower health status.

II.2.7. have been loaded for dispatch to the Union on _____ (dd/mm/yyyy)(3) in a means of transport which was cleaned and disinfected prior to loading with a disinfectant authorised by the competent authority in the third country or territory and constructed in such a way that:

(i) animals cannot escape or fall out;

(ii) visual inspection of the space where animals are kept is possible;

(iii) the escape of animal excrements, litter or feed is prevented or minimized.

II.2.8. have been subjected to a clinical inspection within the 24 hour period prior to loading for dispatch to the Union, carried out by an official veterinarian in the third country or territory of origin, who did not detect signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 and emerging diseases.

II.2.9. have not been vaccinated against:

(i) foot and mouth disease, infection with rinderpest virus, infection with Rift Valley fever virus, infection with peste des petits ruminants virus, Mycobacterium tuberculosis complex (M.bovis, M.caprae and M.tuberculosis), infection with Brucella abortus, B. melitensis and B. suis, and

Part II : Certification

II. Health information

(ii) infection with bluetongue virus (serotypes 1-24) with a live vaccine during the 60 days prior to their dispatch to the Union.

II.2.10. come from a zone:

II.2.10.1. in which:

(i) foot and mouth disease has not been reported for:

either ○ [at least 24 months prior to the date of dispatch to the Union](1)

or ○ [since _____ (dd/mm/yyyy)](1)(4)

(ii) vaccination against foot and mouth disease has not been carried out for at least 12 months prior to the date of dispatch to the Union, and no animals vaccinated against foot and mouth disease have been introduced during that period.

II.2.10.2. infection with rinderpest virus, infection with Rift Valley fever virus, infection with peste des petits ruminants virus has not been reported for the 12 month period prior to dispatch to the Union and during that period:

(i) vaccination against these diseases has not been carried out, and

(ii) animals vaccinated against these diseases have not been introduced.

either ○ [II.2.10.3. which is free from infection with bluetongue virus (serotypes 1-24)](1)(5)

or ○ [II.2.10.3. which is seasonally free from infection with bluetongue virus (serotypes 1-24):

either ○ [II.2.10.3.1. for at least 60 days prior to the date of dispatch of the animals to the Union.](1)(6)

or ○ [II.2.10.3.1. for at least 28 days prior to the date of dispatch of the animals to the Union and the animals have been subjected to a serological test in accordance with Article 9(b) of Delegated Regulation (EU) 2020/692, with negative results, carried out on samples collected at least 28 days following the date of entry of the animal into the seasonally free zone.](1)(6)

or ○ [II.2.10.3.1. for at least 14 days prior to the date of dispatch of the animals to the Union and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of entry of the animal in the seasonally free zone.](1)(6)

or ○ [II.2.10.3. is not free from infection with bluetongue virus (serotypes 1-24) and the animals have been vaccinated against all the serotypes (1 to 24) of bluetongue virus reported during the past 2 years in that zone and are still within the immunity period of time guaranteed in the specifications of the vaccine and

either ○ [II.2.10.3.1. have been vaccinated more than 60 days prior to the date of dispatch to the Union.]](1)

or ○ [II.2.10.3.1. have been vaccinated with an inactivated vaccine and were subjected to a PCR test, with negative results on samples collected at least 14 days after the onset of the immunity protection set in the specifications of the vaccine.]] (1)

or ○ [II.2.10.3. is not free from infection with bluetongue virus (serotypes 1-24) and the animals have been subjected with positive results to a serological test able to detect specific antibodies against all serotypes (1 to 24) of bluetongue virus reported during the past 2 years in that zone and:

either ○ [II.2.10.3.1. the serological test has been carried out on samples collected at least 60 days prior to the date of dispatch of the animals to the Union.]](1)

Part II : Certification

II. Health information

or ○ [II.2.10.3.1.] the serological test has been carried out on samples collected at least 30 days prior to the date of dispatch of the animals to the Union and the animals were subjected to a PCR test, with negative results, carried out on samples collected not earlier than 14 days prior to the date of dispatch of the animals to the Union.]](1)

II.2.11. come from an establishment:

II.2.11.1. which is registered by and under the control of the competent authority of the third country or territory of origin and has a system in place to maintain for at least 3 years up-to-date records containing information regarding:

(i) the species, categories, number and identification of animals on the establishment;

(ii) movements of animals into and out of the establishment;

(iii) mortality in the establishment.

II.2.11.2. which receives regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 and emerging diseases, at a frequency that is proportional to the risk posed by the establishment.

II.2.11.3. which was not subject to national restriction measures for animal health reasons, including listed diseases and emerging diseases, at the time of dispatch to the Union.

II.2.11.4. in and around which, in an area with a 10 km radius, including where appropriate the territory of a neighbouring country, none of the following listed diseases has been reported for at least 30 days prior to dispatch to the Union: foot and mouth disease, infection with Rift Valley fever virus and infection with peste des petits ruminants virus.

either ○ [II.2.11.5. in and around which, in an area with a 150 km radius, including where appropriate the territory of a neighbouring country, epizootic haemorrhagic disease has not been reported for at least 2 years before the date of dispatch to the Union.](1)

or ○ [II.2.11.5. which is located in a zone seasonally free of epizootic haemorrhagic disease.](1)(7)

II.2.11.6. subjected to surveillance to detect infection with Mycobacterium tuberculosis complex (M.bovis, M.caprae and M.tuberculosis) in animals of the same species of animals as the animals of the consignment in accordance with the procedures in points (1) and (2) of part 2 of Annex II to Commission Delegated Regulation (EU) 2020/688 during at least the 12 month period prior to dispatch to the Union and during this period:

(i) only animals from establishments applying the measures provided in the paragraph above have been introduced in the establishment;

(ii) ☐ [infection with Mycobacterium tuberculosis complex (M.bovis, M.caprae and M.tuberculosis) has been reported in animals of the same species of animals as the animals of the consignment kept on the establishment and measures were taken in accordance with point (3) of part 2 of Annex II to Delegated Regulation (EU) 2020/688]].(1)

II.2.11.7. in which infection with Brucella abortus, B. melitensis and B. suis in animals of the same species of animals as the animals of the consignment has not been reported during the last 42 days prior to dispatch to the Union, and the animals of the consignment have been subjected to a test for the detection of infection with Brucella abortus, B. melitensis and B. suis with one of the diagnostic methods provided for in Part 1 of Annex I to Delegated Regulation (EU) 2020/688, with negative results, carried out on a sample taken during the 30 day period prior to dispatch to the Union, and in the case of post-parturient females, taken at least 30 days after parturition.

Part II : Certification

II. Health information

II.2.11.8. in which rabies has not been reported for at least the 30 days prior to dispatch to the Union.

II.2.11.9. in which anthrax has not been reported for at least the 15 days prior to dispatch to the Union.

II.2.11.10. in which surra (Trypanosoma evansi) has not been reported for at least the 30 days prior to dispatch to the Union and if the disease was reported in the establishment of origin during the last 2 years prior to dispatch to the Union, the affected establishment remained under restriction until the infected animals were removed from the establishment and the remaining animals on the establishment were subjected with negative result to a test for surra (Trypanosoma evansi) as described in Part 3 of Annex I to Delegated Regulation (EU) 2020/688 carried out on samples taken at least 6 months after the infected animals were removed from the establishment.

☐ [II.2.11.11. in which, if an infection with Burkholderia mallei (glanders) has been reported during the period of 3 years prior to the date of dispatch to the Union, following the last outbreak the establishment remained under movement restrictions by the competent authority until:

(i) the infected animals have been killed and destroyed; and

(ii) the remaining animals were subjected to a test carried out as described in point 3.1 of Chapter 3.5.11 of the OIE Terrestrial Manual (Version adopted 2015) with negative results on samples taken at least 6 months after the date on which the infected animals were killed and destroyed and the establishment cleaned and disinfected](1)(8)

☐ [II.2.12. originate from an establishment in which infectious bovine rhinotracheitis/infectious pustular vulvovaginitis has not been reported on camelid animals in the 30 day period prior to dispatch to the Union.](1)(9)

Part II : Certification

II. Health information

Notes:

This certificate is intended for entry into the Union of camelid and cervid animals, including when the Union is not the final destination of those animals.

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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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: "Identification system and identification number": Specify the identification system (such as ear tag, tattoo, transponder etc., from the list in Annex III to Commission Delegated Regulation (EU) 2019/2035) and the individual identification codes of the animals in accordance with Article 21(1) of Delegated Regulation (EU) 2020/692.

Part II:

(1) Keep as appropriate.

(2) Code of the zone as it appears in Column 2 of Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

(3) Date of loading: entries of these animals shall not be permitted when the animals were loaded for dispatch to the Union either prior to the date of authorisation for entry into the Union of the third country, territory or zone thereof referred to in point II.2.1., or during a period where restriction measures have been adopted by the Union against entries of these animals from this third country, territory or zone thereof.

(4) Only for countries with the opening date in column 8 in Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

(5) For countries with entry BTV in Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

(6) For countries with entry SF-BTV in Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

(7) For countries with entry SF-EHD in Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

(8) Only applicable for ungulates of the family Camelidae.

(9) Only applicable when the Member State of destination or Switzerland, in accordance with the Agreement between the European Community and the Swiss Confederation on trade in agricultural products (OJ L 114, 30.4.2002), either have disease-free status for infectious bovine rhinotracheitis /infectious pustular vulvovaginitis in bovine animals or an approved eradication programme.

Official veterinarian or Official inspector

Name (in capital letters) | Qualification and title

Date of signature | Signature

Stamp