Commission Implementing Regulation (EU) No 780/2013 of 14 August 2013 amending Commission Regulation (EU) No 206/2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (Text with EEA relevance)

ANNEX

ANNEX VI

PART 1

Table 1

"RUM-A" : Model of veterinary certificate for animals of the species listed below that are originating from and intended for an approved body, institute or centre.

Order	Family	Genera/species
Artiodactyla	Antilocapridae	*Antilocapra* ssp.
	Bovidae	*Addax* ssp., *Aepyceros* ssp., *Alcelaphus* ssp., *Ammodorcas* ssp., *Ammotragus* ssp., *Antidorcas* ssp., *Antilope* ssp., *Bison* ssp., *Bos* ssp. *(including Bibos, Novibos, Poephagus), Boselaphus* ssp., *Bubalus* ssp. *(including anoa), Budorcas* ssp., *Capra* ssp., *Cephalophus* ssp., *Connochaetes* ssp., *Damaliscus* ssp. *(including Beatragus), Dorcatragus* ssp., *Gazella* ssp., *Hemitragus* ssp., *Hippotragus* ssp., *Kobus* ssp., *Litocranius* ssp., *Madoqua* ssp., *Naemorhedus* ssp. *(including Nemorhaedus and Capricornis), Neotragus* ssp., *Oreamnos* ssp., *Oreotragus* ssp., *Oryx* ssp., *Ourebia* ssp., *Ovibos* ssp., *Ovis* ssp., *Patholops* ssp., *Pelea* ssp., *Procapra* ssp., *Pseudois* ssp., *Pseudoryx* ssp., *Raphicerus* ssp., *Redunca* ssp., *Rupicapra* ssp., *Saiga* ssp., *Sigmoceros-Alecelaphus* ssp., *Sylvicapra* ssp., *Syncerus* ssp., *Taurotragus* ssp., *Tetracerus* ssp., *Tragelaphus* ssp. (*including Boocerus*).
	Camelidae	*Camelus* ssp., *Lama* ssp., *Vicugna* ssp.

	Cervidae	*Alces* ssp., *Axis-Hyelaphus* ssp., *Blastocerus* ssp., *Capreolus* ssp., *Cervus-Rucervus* ssp., *Dama* ssp., *Elaphurus* ssp., *Hippocamelus* ssp., *Hydropotes* ssp., *Mazama* ssp., *Megamuntiacus* ssp., *Muntiacus* ssp., *Odocoileus* ssp., *Ozotoceros* ssp., *Pudu* ssp., *Rangifer* ssp.
	Giraffidae	*Giraffa* ssp., *Okapia* ssp.
	Moschidae	*Moschus* ssp.
	Tragulidae	*Hyemoschus* ssp., *Tragulus-Moschiola* ssp.

Table 2

"SUI-A" : Model of veterinary certificate for animals of the species listed below that are originating from and intended for an approved body, institute or centre.

Order	Family	Genera/species
Artiodactyla	Suidae	*Babyrousa* ssp., *Hylochoerus* ssp., *Phacochoerus* ssp., *Potamochoerus* ssp., *Sus* ssp.
	Tayassuidae	*Catagonus* ssp., *Pecari-Tayassu* ssp.
	Hippopotamidae	*Hexaprotodon-Choeropsis* ssp., *Hippopotamus* ssp.

Table 3

"TRE-A" : Model of veterinary certificate for animals of the species listed below that are originating from and intended for an approved body, institute or centre.

Order	Family	Genera/species
Perissodactyla	Tapiridae	*Tapirus* ssp.
	Rhinocerotidae	*Ceratotherium* ssp., *Dicerorhinus* ssp., *Diceros* ssp., *Rhinoceros* ssp.
Proboscidea	Elephantidae	*Elephas* ssp., *Loxodonta* ssp.

PART 2

Model RUM-A

COUNTRY **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code Tel.		I.6.	
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 origin Name Approval number Address		I.12.	
I.13. Place of loading Address Approval number		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references		I.16. Entry BIP in EU	
		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code)	
		I.20. Quantity	
I.21.		I.22. Number of packages	
I.23. Seal/Container No		I.24.	
I.25. Commodities certified for: Approved body ☐			
I.26.		I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (scientific name)	Identification system	Identification number	Age	Sex

COUNTRY **Model RUM-A**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

II.1. **Animal health attestation**

I, the undersigned official veterinarian responsible for the approved body, institute or centre/holding (1) of origin certify that the animals described in Part I meet the following requirements:

II.1.1. They come from the country, territory or part thereof described in Box I.7.:

(a) where the diseases referred to in this certificate are notifiable,

►(1) (b) which at the date of issuing this certificate has been free for 12 months from rinderpest.◄

II.1.2. They come from the body, institute or centre/holding (1) described in Box I.11;

(a) which is approved according to the requirements and conditions set out in Part 3 and 4 of Annex VI to Regulation (EU) No 206/2010;

(b) which is not subjected to any restrictions relating to a national programme for the control of infectious diseases to which the animals referred to in Box I.28. are susceptible;

(c) where there have been no clinical cases of the following diseases to which the animals referred to in Box I.28. are susceptible:

— anthrax for the last 30 days;

— foot-and-mouth disease, bluetongue, Rift valley fever, vesicular stomatitis, rabies, contagious bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox, goat pox, contagious caprine pleuropneumonia for the past 6 months;

(d) where there have been no clinical or non-clinical cases of tuberculosis and brucellosis for the past 6 months;

(e) around which in an area of 10 km radius for the last 30 days, there has been no case of the following diseases to which the animals referred to in Box I.28. are susceptible: foot-and-mouth disease, vesicular stomatitis, contagious bovine pleuropneumonia, peste des petits ruminants, sheep pox, goat pox, contagious caprine pleuropneumonia;

(f) around which in an area of 150 km radius for the last 30 days, there has been no case of the following diseases to which the animals referred to in Box I.28. are susceptible: bluetongue, epizootic haemorrhagic disease, Rift valley fever, lumpy skin disease;

(g) in which they have remained since birth or for the past 6 months before dispatch to the Union.

II.1.3. They:

(a) have not come into contact with other animals not complying with at least the same health requirements as described in this certificate for the last 30 days and during their transportation from the approved body, institute or centre/holding (1) to the place of shipment;

(b) were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease and are fit for the intended transport;

(c) are not animals to be killed under a national programme for the eradication of diseases.

II.1.4. **Foot-and-Mouth Disease**

either (1) [(a) They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from foot-and-mouth disease with or without vaccination, and]

or (1) [(a) They have been subjected to the following tests:

— a serological test for evidence of foot-and-mouth disease virus infection carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals (OIE Terrestrial Manual), with negative results, taken within 10 days prior to dispatch to the Union,

— (1)(2)[a probang test for evidence of foot-and-mouth disease virus infection carried out in accordance with the procedures described in the OIE Terrestrial Manual with negative results, (1)(3)[taken 10 days prior to dispatch to the Union] (1)(4)[taken on two occasions 15 days apart, the second of which must have been taken 10 days prior to dispatch to the Union, and]

►(2)(1) (b) they have not been vaccinated against foot-and-mouth disease.◄

COUNTRY **Model RUM-A**

II.	Health information	II.a. Certificate reference number	II.b.

II.1.5. Bluetongue and Epizootic haemorrhagic disease (EHD)

either (1) [They come from the country, territory or part thereof described in Box I.7 which has been free for 24 months from bluetongue/EHD in accordance with the OIE Terrestrial Animal Health Code (OIE Terrestrial Code).]

or (1) [They were held in a vector-protected facility in the approved body, institute or centre/holding (1) for at least 30 days prior to shipment and were subjected to a serology test according to the OIE Terrestrial Manual, with negative results, carried out at least 28 days after introduction into the approved body, institute or centre.]

or (1) [They were held in a vector-protected facility in the approved body, institute or centre/holding (1) for at least 30 days prior to shipment and were subjected to a PCR test according to the OIE Terrestrial Manual, with negative results, carried out at least 14 days after introduction into the approved body, institute or centre.]

or (1) [They come from a seasonally free area and were subjected during that period to an serology test according to the OIE Terrestrial Manual, with negative results, carried out at least 28 days after introduction into the approved body, institute or centre/holding (1).]

or (1) [They come from a seasonally free area and were subjected during that period to a PCR test according to the OIE Terrestrial Manual, with negative results, carried out at least 14 days after introduction into the approved body, institute or centre/holding (1).]

II.1.6. Rift valley fever

either (1) [They come from the country, territory or part thereof described in Box I.7. which has been free for 48 months from Rift valley fever and have not been vaccinated against that disease.]

or (1) [They were held in a vector-protected facility in the approved body, institute or centre/holding (1) for at least 30 days prior to shipment during which the animals showed no clinical signs of Rift valley fever and were protected from vectors between the vector-protected facility and the place of shipment to the Union as well as at the place of shipment.]

or (1) [They have been subjected to a virus neutralisation test (9) with negative results for evidence of Rift valley fever, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken at the beginning of the isolation/quarantine period and at least 42 days later on, the second of which must have been taken within 10 days of dispatch to the Union.]

II.1.7. Brucellosis

either (1) [They come from a country, territory or part thereof described in Box I.7 which has been free for the past 12 months from brucellosis and which have not been vaccinated against that disease;]

or (1) [They have been subjected to a test as laid down and prescribed for international trade by the OIE Terrestrial Manual, in the 30 days prior to dispatch to the Union;]

or (1) [They are castrated males of any age].

II.1.8. Other vaccinations

(a) They have not been vaccinated against vesicular stomatitis,

(5) (b) They have been vaccinated against:

(1) [anthrax on the ..... (dd/mm/yyyy)(date(s)) with the following vaccine(s) ..... (name of vaccine(s) used)],

(1) [rabies on the ..... (dd/mm/yyyy)(date(s)) with the following vaccine(s) ..... (name of vaccine(s) used) and a blood test performed on ..... (dd/mm/yyyy)(date(s)) shows a protective immune response.].

II.1.9. Parasite treatment

They have been treated at least twice during the 40 days prior to dispatch to the Union against internal and external parasites with the following product(s) ..... Specify the active ingredients and the doses of the products used .....

II.1.10. Loading on the means of transport

They have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) (6) in the means of transport described in Box I.15. that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.

COUNTRY **Model RUM-A**

II. Health information	II.a. Certificate reference number	II.b.

Notes

This certificate is to be used for live animals listed in the note for Box I.28. coming from an approved body, institute or centre in a third country, territory ot part thereof, and destined to an approved body, institute or centre situated within a Member State. Use one certificate per species.

Part I:

— 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 shall inform the BIP of entry into the EU.

— Box reference I.19.: Use appropriate HS code: 010613 or 010619.

— Box reference I.28.: *Identification system*: Specify the identification system (tag, tattoos, brand, chip, transponder). The identifier shall include the ISO code of the exporting country and permit tracing of their premises of origin.

Age: months.

Sex (M = male, F = female, C = castrated).

Species: Select the species amongst those listed below:

Order	**Family**	**Genera/species**
Artiodactyla	Antilocapridae	*Antilocapra*
	Bovidae	*Addax* ssp., *Aepyceros* ssp., *Alcelaphus* ssp., *Ammodorcas* ssp., *Ammotragus* ssp., *Antidorcas* ssp., *Antilope* ssp., *Bison* ssp., *Bos* ssp. *(including Bibos, Novibos, Poephagus), Boselaphus* ssp., *Bubalus* ssp. *(including anoa), Budorcas* ssp., *Capra* ssp., *Cephalophus* ssp., *Connochaetes* ssp., *Damaliscus* ssp. *(including Beatragus), Dorcatragus* ssp., *Gazella* ssp., *Hemitragus* ssp., *Hippotragus* ssp., *Kobus* ssp., *Litocranius* ssp., *Madoqua* ssp., *Naemorhedus* ssp. *(including Nemorhaedus and Capricornis), Neotragus* ssp., *Oreamnos* ssp., *Oreotragus* ssp., *Oryx* ssp., *Ourebia* ssp., *Ovibos* ssp., *Ovis* ssp., *Patholops* ssp., *Pelea* ssp., *Procapra* ssp., *Pseudois* ssp., Pseudoryx ssp., *Raphicerus* ssp., *Redunca* ssp., *Rupicapra* ssp., *Saiga* ssp., *Sigmoceros-Alecelaphus* ssp., *Sylvicapra* ssp., *Syncerus* ssp., *Tauro-tragus* ssp., *Tetracerus* ssp., *Tragelaphus* ssp. *(including Boocerus).*
	Camelidae	*Camelus* ssp., *Lama* ssp., *Vicugna* ssp.
	Cervidae	*Alces* ssp., *Axis-Hyelaphus* ssp., *Blastocerus* ssp., *Capreolus* ssp., *Cervus-Rucervus* ssp., *Dama* ssp., *Elaphurus* ssp., *Hippocamelus* ssp., *Hydropotes* ssp., *Mazama* ssp., *Megamuntiacus* ssp., *Muntiacus* ssp., *Odocoileus* ssp., *Ozotoceros* ssp., *Pudu* ssp., *Rangifer* ssp.
	Giraffidae	*Giraffa* ssp., *Okapia* ssp.
	Moschidae	*Moschus* ssp.
	Tragulidae	*Hyemoschus* ssp., *Tragulus-Moschiola* ssp.

Part II:

([1]) Keep as appropriate.

([2]) This attestation is only applicable to *Bovidae* and *Cervidae.*

([3]) This attestation is only applicable to *Bovidae* and *Cervidae* other than African buffalo (*Syncerus caffer*).

([4]) This attestation is only applicable to African buffalo (*Syncerus caffer).*

([5]) Vaccination is not compulsory, but if the animals have been vaccinated, information on the vaccine(s) used and the time of vaccination shall be filled in.

([6]) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country,territory or part thereof described in Boxes I.7. and I.8., or during a period where restrictive measures have been adopted by the Union against imports of these animals from that country,territory or part thereof.

COUNTRY **Model RUM-A**

II. Health information	II.a. Certificate reference number	II.b.
Official veterinarian Name (in capital letters): Date: Stamp:		Qualification and title: Signature:

Model SUI-A

COUNTRY **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code Tel.		I.6.	
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 origin Name Approval number Address		I.12.	
I.13. Place of loading Address Approval number		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references		I.16. Entry BIP in EU	
		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **01.06.19**	
		I.20. Quantity	
I.21.		I.22. Number of packages	
I.23. Seal/Container No		I.24.	
I.25. Commodities certified for: Approved body ☐			
I.26.		I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (scientific name)	Identification system	Identification number	Age	Sex

COUNTRY **Model SUI-A**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

II.1. **Animal health attestation**

I, the undersigned official veterinarian responsible for the approved body, institute or centre/holding (1) of origin certify that the animals described in Part I meet the following requirements:

II.1.1. They come from the country, territory or part thereof described in Box I.7.

(a) where the diseases referred to in this certificate are notifiable,

(b) which at the date of issuing this certificate has been free for the past 12 months from rinderpest.

II.1.2. They come from the body, institute or centre/holding (1) described in Box I.11.

(a) which is approved according to the requirements and conditions set out in Part 3 and 4 of Annex VI to Regulation (EU) No 206/2010;

(b) which is not subjected to any restrictions relating to a national programme for the control of infectious diseases to which the animals referred to in Box I.28. are susceptible;

(c) where there have been no clinical cases of the following diseases to which the animals referred to in Box I.28. are susceptible:

— anthrax for the last 30 days;

— foot-and-mouth disease, vesicular stomatitis, rabies, African swine fever, classical swine fever and swine vesicular disease for the past 6 months;

(d) where there have been no clinical or non-clinical cases of tuberculosis and brucellosis for the past 6 months;

(e) around which in an area of radius of 10 km for the last 12 months, there has been no case/outbreak of African swine fever, classical swine fever and swine vesicular disease;

(f) around which in an area of 10 km radius for the past 30 days, there has been no case/outbreak of foot-and-mouth disease or vesicular stomatitis,

(g) in which they have remained since birth or for the past 6 months before dispatch to the Union.

II.1.3. They:

(a) have not come into contact with other animals not complying with at least the same health requirements as described in this certificate since birth or for the last 30 days and during their transportation from the approved body, institute or centre/holding (1) to the place of shipment;

(b) were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease and are fit for the intended transport;

(c) are not animals to be killed under a national programme for the eradication of diseases.

II.1.4. **Foot-and-Mouth Disease**

either (1) [(a) They come from the country, territory or part thereof described in Box I.7. which at the date of issuing this certificate has been free for the past 12 months from foot-and-mouth disease and;]

or (1) [(a) They have been subjected to a virological and serological test for evidence of foot-and-mouth disease virus infection carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals (OIE Terrestrial Manual), with negative results, taken in the 10 days prior to dispatch to the Union; and]

(b) they have not been vaccinated against foot-and-mouth disease.

II.1.5. **Brucellosis**

(1) *either* [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from brucellosis and have not been vaccinated against that disease]

(1)(3) *or* [They have been subjected, with negative results, to a buffered *Brucella* antigen test for porcine brucellosis taken in the 30 days prior to dispatch to the Union.]

COUNTRY **Model SUI-A**

II. Health information	II.a. Certificate reference number	II.b.

II.1.6. Swine vesicular disease

(1) *either* [They come from the country, territory or part thereof described in box 1.7 which has been free for the past 12 months from swine vesicular disease.]

(1) *or* [They have been subjected, with negative results, to a virology and serology test for evidence of swine vesicular disease, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to the Union.]

II.1.7. Vesicular Stomatitis

(1) *either* [They come from the country, territory or part thereof described in Box I.7 which has been free for the last 6 months from vesicular stomatitis.]

(1) *or* [They have been subjected, with negative results, to a virology and serology test for evidence of vesicular stomatitis, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to the Union.]

II.1.8. Classical swine fever

(1) *either* [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from classical swine fever.]

(1) *or* [They have been subjected to a virological and serological test for classical swine fever carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Terrestrial Manual, with negative results, taken in the 30 days prior to dispatch to the Union.]

II.1.9. African swine fever

(1) *either* [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from African swine fever.]

(1) *or* [They have been subjected, with negative results, to a virus and serology test for African swine fever, as laid down and prescribed for international trade in the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to the Union.]

II.1.10. Aujeszky's disease

According to official information, no clinical, pathological or serological evidence of Aujeszky's disease has been recorded for the last 12 months in the approved body, institute or centre/holding (1) and in an area with a 5 km radius around the approved body, centre or institute, and

They have been subjected, with negative results, to a virology and serology test for evidence of Aujeszky's disease, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to the Union, and

They have not been vaccinated against Aujeszky's disease and have not been in contact with vaccinated animals.

II.1.11. Other vaccinations

(a) They have not been vaccinated against rinderpest, vesicular stomatitis, classical swine fever or swine vesicular disease,

(2)(b) They have been vaccinated against:

(1) [anthrax on the ..... (dd/mm/yyyy) with the following vaccine(s) ..... (name of vaccine (s) used)],

(1) [rabies on the ..... (dd/mm/yyyy) with the following vaccine(s) ..... (name of vaccine (s) used)].

II.1.12. Parasite treatment

They have been treated at least twice in the 40 days prior to dispatch to the Union against internal and external parasites with the following product(s) ..... . Specify the active ingredients and the doses of the products used .....

COUNTRY **Model SUI-A**

II. Health information	II.a. Certificate reference number	II.b.

II.1.13. Loading on the means of transport

They have been loaded for dispatch to the Union on .....(dd/mm/yyyy) ([4]) in the means of transport described in Box I.15. that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.

Notes

This certificate is meant for animals of species listed in the note for Box I. 28. coming from an approved body, institute or centre in a third country, territory ot part thereof, and destined to an approved body, institute or centre located within a Member State.

Part I:

— 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 shall inform the BIP of entry into the EU.

— Box reference I.28.: *Identification system*: Specify the identification system (tag, tattoos, brand, chip, transponder). The identifier shall include the ISO code of the exporting country and permit tracing of their premises of origin.

Age: months.

Sex (M = male, F = female, C = castrated).

Species Select the species amongst those listed below:

Order	**Family**	**Genera/species**
Artiodactyla	Suidae	*Babyrousa* ssp., *Hylochoerus* ssp., *Phacochoerus* ssp., *Potamochoerus* ssp., *Sus* ssp.
	Tayassuidae	*Catagonus* ssp., *Pecari-Tayassu* ssp.
	Hippopotamidae	*Hexaprotodon-Choeropsis, Hippopotamus* ssp.

Part II:

([1]) Keep as appropriate.

([2]) Vaccination is not compulsory, but if the animals have been vaccinated, information on the vaccine(s) used and the time of vaccination must be filled in.

([3]) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex I to Regulation (EU) No 206/2010.

([4]) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the country, territory or part thereof decribed in Boxes I.7. and I.8., or during a period where restrictive measures have been adopted by the Union against imports of these animals from that country,territory or part thereof.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:

Model TRE-A

COUNTRY **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.

I.2. Certificate reference No | I.2.a.

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Postal code
Tel.

I.6.

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 origin
Name Approval number
Address

I.12.

I.13. Place of loading
Address Approval number

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification
Documentary references

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code) **01.06.19**

I.20. Quantity

I.21.

I.22. Number of packages

I.23. Seal/Container No

I.24.

I.25. Commodities certified for:
Approved body ☐

I.26.

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (scientific name)	Identification system	Identification number	Age	Sex

COUNTRY **Model TRE-A**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

II.1. **Animal health attestation**

I, the undersigned official veterinarian responsible for the approved body, institute or centre/holding (1) of origin certify that the animals described in Part I meet the following requirements:

II.1.1. They come from the third country, territory or part thereof described in Box I.7.

(a) where the diseases referred to in this certificate are notifiable,

(b) which at the date of issuing this certificate has been free for the past 12 months from rinderpest.

II.1.2. They come from the body, institute or centre/holding (1) described in Box I.11.,

(a) which is approved according to the requirements and conditions set out in Part 3 and 4 of Annex VI to Regulation (EU) No 206/2010;

(b) which is not subjected to any restrictions relating to a national programme for the control of infectious diseases to which the animals referred to in Box I.28. are susceptible;

(c) where there have been no clinical cases of the following diseases to which the animals referred to in Box I.28. are susceptible:

— anthrax for the last 30 days;

— foot–and–mouth disease, rabies, (1)(2) [African horse sickness] for the past 6 months,

(d) where there have been no clinical or non-clinical cases of tuberculosis for the past 6 months;

(e) around which in an area of 10 km radius for the last 30 days, there has been no case/outbreak of foot–and–mouth disease,

(f) in which they have remained since birth or for the past 6 months before dispatch to the Union,

(1)(2) [(g) around which in an area of radius of 150 km for the last 60 days, there has been no case/outbreak of African horse sickness].

II.1.3. They:

(a) have not come into contact with other animals not complying with at least the same health requirements as described in this certificate since birth or for the past 30 days and during their transportation from the approved body, institute or centre/holding (1) to the place of shipment;

(b) were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease and are fit for the intended transport;

(c) are not animals to be killed under a national programme for the eradication of diseases.

(1)(3) **[II.1.4. Foot–and-Mouth Disease**

either (1) [(a) They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from foot-and-mouth disease with or without vaccination, and]

or (1) [(a) They have been subjected to the following tests:

— a serological test for evidence of foot-and-mouth disease virus infection carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals (OIE Terrestrial Manual), with negative results, taken in the 10 days prior to dispatch to the Union, and

— [a probang test for evidence of foot-and-mouth disease virus infection carried out in accordance with the procedures described in the OIE Terrestrial Manual with negative results, taken 10 days prior to dispatch to the Union and]

(b) have not been vaccinated against foot-and-mouth disease.

II.1.5. Other vaccinations

(a) They have not been vaccinated against rinderpest,

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(4) (b) They have been vaccinated against:

(1) [anthrax on the ..... (dd/mm/yyyy)(date(s)) with the following vaccine(s) ..... (name of vaccine(s) used)],

(1) [rabies on the ..... (dd/mm/yyyy)(date(s)) with the following vaccine(s) ..... (name of vaccine (s) used)].

II.1.6. Parasite treatment

They have been treated at least twice in the 40 days prior to dispatch to the Union against internal and external parasites with the following product(s) ..... Specify the active ingredients and the doses of the products used .....

II.1.7. Loading on the means of transport

They have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) (5) in the means of transport described in Box I.15 that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.

Notes

This certificate is meant for live animals as listed in the note for Box I.28. coming from an approved body, institute or centre in a third country, territory or part thereof, and destined for an approved body, institute or centre located within a Member State. Use one certificate per species.

Part I:

— 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 shall inform the BIP of entry into the EU.

— Box reference I.28.: *Identification system:* Specify the identification system (tag, tattoos, brand, chip, transponder). The identifier shall include the ISO code of the exporting country and permit tracing of their premises of origin.

Age: months.

Sex (M = male, F = female, C = castrated).

Species: Select the species amongst those listed below:

Order	**Family**	**Genera/species**
Perissodactyla	Tapiridae	*Tapirus* ssp.
	Rhinocerotidae	*Ceratotherium* ssp., *Dicerorhinus* ssp., *Diceros* ssp., *Rhinoceros* ssp
Proboscidea	Elephantidae	*Elephas* ssp., *Loxodonta* ssp.

Part II:

(1) Keep as appropriate.

(2) This attestation is only applicable to *Rhinocerotidae*.

(3) This attestation is only applicable to *Elephas*. ssp.

(4) Vaccination is not compulsory, but if the animals have been vaccinated, information on the vaccine(s) used and the time of vaccination must be filled in.

(5) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country,territory or part thereof described in Boxes I.7. and I.8., or during a period where restrictive measures have been adopted by the Union against imports of these animals from that third country, territory or part thereof.

COUNTRY		Model TRE-A
II. Health information	II.a. Certificate reference number	II.b.
Official veterinarian Name (in capital letters): Qualification and title: Date: Signature: Stamp:		

PART 3

Requirements concerning bodies, institutes or centres in third countries

The body, institute or centre in a third country must:

(a) be clearly demarcated and separated from its surroundings;

(b) have adequate means for catching, confining and isolating animals, and have available adequate quarantine facilities and approved standard operating procedures for animals coming from unknown origin;

(c) have a vector-protected structure complying with the following requirements:

  (i) it has appropriate physical barriers at entry and exit points;

  (ii) the openings of the vector-protected structure are vector-screened with mesh of appropriate gauge impregnated regularly with an approved insecticide according to the instructions of the manufacturer;

  (iii) vector surveillance and control are carried out within and around the vector-protected structure;

  (iv) measures are taken to limit or eliminate breeding sites for vectors in the vicinity of the vector-protected structure;

  (v) standard operating procedures are in place, including descriptions of back-up and alarm systems, for the operation of the vector-protected structure and for the transport of the animals from that structure to the place of loading;

(d) keep, for a minimum period of ten years, up-to-date records indicating:

  (i) the number and identity (age, sex, species and individual identification, where appropriate) of the animals of each species present on their premises;

  (ii) the number and identity (age, sex, species and individual identification where appropriate) of animals arriving in or leaving their premises, together with information on their origin or destination, the means of transport, and the health status of those animals;

  (iii) the results of blood tests or any other diagnostic procedures carried out on the animals on their premises;

  (iv) cases of disease and, where appropriate, the treatment administered;

(v) the results of the post-mortem examinations on animals that have died on their premises, including still-born animals;

(vi) observations made during any isolation or quarantine period;

(e) be free from the diseases listed in Annex A to Directive 92/65/EEC or mentioned in the veterinary certificates for the relevant species set out in Part 2 of Annex VI to this Regulation, for at least the previous three years, as evidenced by the records kept pursuant to point (d) and the results of the clinical and laboratory tests carried out on the animals on their premises;

(f) either have an arrangement with a laboratory approved *by the competent authority* to perform post-mortem examinations, or have one or more appropriate premises where these examinations may be performed under the authority of the approved veterinarian;

(g) ensure disposal of the carcasses of animals which die of a disease or are euthanised;

(h) secure, by contract or legal instrument, the services of a veterinarian approved by and acting under the control of the competent authority, who must perform at least the following tasks:

(i) ensure that appropriate disease surveillance and control measures are applied in that body, institute or centre. Such measures must be approved by the competent authority of the third country, territory or part thereof where the body, institute or centre is situated, taking into account the disease situation and must include at least the following elements:

— an annual disease surveillance plan including appropriate control measures concerning zoonoses in the animals present on the premises,

— clinical, laboratory and post-mortem testing of animals suspected to be affected by transmissible diseases and zoonoses,

— vaccination of susceptible animals against infectious diseases and zoonoses;

(ii) ensure that any suspect deaths or the presence of any other symptom suggesting that animals have contracted one or more of the diseases listed in Annex A to Directive 92/65/EEC or mentioned in the veterinary certificates for the relevant species set out in Part 2 of Annex VI to this Regulation are notified without delay to the competent authority, where that particular disease is notifiable in the third country, territory or part thereof concerned;

(iii) ensure that incoming animals have been quarantined as necessary, in accordance with the instructions given by the competent authority;

(iv) ensure compliance with the animal health requirements which the animals must fulfil in order to be introduced into the Union.

PART 4

Conditions concerning the approval of bodies, institutes or centres in third countries

1. Approval must be granted only to those bodies, institutes or centres which comply with the requirements set out in Part 3.

2. Where vector protection is required, the approval of a structure as vector-protected must be granted only if the criteria in point (c) of Part 3 are met. In order to grant the approval, the competent authority must verify at least three times during the required protection period (at the beginning, during and at the end of the period) the effectiveness of the vector protection measures, by means of a vector trap inside the vector protected structure.

3. Each approved body, institute and centre must be assigned an approval number.

4. Approval must be maintained only as long as the following conditions continue to be met:

the premises are under the control of an official veterinarian, who must perform at least the following tasks:

(i) inspect the premises of the body, institute or centre at least once per year;

(ii) audit the activity of the veterinarian referred to in point (h) of Part 3 and the implementation of the annual disease surveillance plan referred to in the first indent of point (h)(i);

(iii) ensure that the provisions laid down in Parts 3 and 4 are met;

(iv) verify that:

— compliance with the animal health requirements which the animals must fulfil in order to be introduced into the Union;

— the results of the clinical, post-mortem and laboratory tests on the animals have revealed no occurrence of the diseases listed in Annex A to Directive 92/65/EEC or mentioned in the veterinary certificates for the relevant species set out in Part 2 of Annex VI to this Regulation.

5. The approval must be withdrawn where the competent authority finds that the requirements of Part 3 are no longer being fulfilled.

6. Where notification is given of the suspicion of the occurrence of one of the diseases listed in Annex A to Directive 92/65/EEC or mentioned in the veterinary certificates for the relevant species laid down in Part 2 of Annex VI to this Regulation, the competent authority must suspend the approval of the body, institute or centre, until the suspicion has been officially ruled out. Depending on the disease involved and the risk of disease transmission, the suspension may relate to the the body, institute or centre as a whole or only to certain categories of animals susceptible to the disease in question. The competent authority must ensure that the measures necessary to confirm or rule out the suspicion and to avoid any spread of disease are taken.

7. Where the suspected disease referred to in point 6 is confirmed, the approval of the body, institute or centre must be withdrawn.

8. Where the approval of a body, institute or centre has been withdrawn, it must be restored only where the following conditions are complied with:

(a) the disease and the source of infection were eradicated on the premises of the body, institute or centre concerned;

(b) the premises of the body, institute or centre concerned were appropriately cleaned and desinfected;

(c) the body, institute or centre concerned complies with the requirements set out in points (a) to (d) and (f) to (h) of Part 3.

9. The competent authority which approved the body, institute or centre must inform the Member States that included the body, institute or centre on their lists of approved bodies, institutes and centres of the suspension, withdrawal or restoration of that approval.

Changes to legislation:
There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 780/2013, ANNEX.