

# Horses to the European Union (OMAR)

EUHOR13

Effective from 30 August 2024

**Te Kāwanatanga o Aotearoa** New Zealand Government

# TITLE

Animal Products Notice: Horses to the European Union (OMAR)

# COMMENCEMENT

This Animal Products Notice comes into force on 30 August 2024

# REVOCATION

This Animal Products Notice revokes and replaces:

• Animal Products Notice: Horses to the European Union (OMAR) dated 15 September 2022

## **ISSUING AUTHORITY**

This Animal Products Notice is issued under sections 167(1) and 60(1) of the Animal Products Act 1999.

Dated at Wellington, 29 August 2024

Trish Mead Manager Animal Health & Exports Ministry for Primary Industries (acting under delegated authority of the Director-General)

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30 August 2024

# Introduction

This introduction is not part of the Animal Products Notice, but is intended to indicate its general effect.

## Purpose

The purpose of this document is to set out the zoosanitary requirements necessary to export compliant horses from New Zealand to or via the European Union.

## Background

The Animal Products Act 1999 provides the controls and mechanisms needed to give and to safeguard official assurances or zoosanitary certificates to facilitate the entry of animal material including live animals, hatching eggs, semen and embryos, and products into overseas markets.

Notices issued as Overseas Market Access Requirements (OMARs) under section 60(1)(a) and (b) of the Animal Products Act specify the requirements that are necessary or desirable for the purpose of facilitating access to overseas markets or are in accordance with the requirements of the relevant authority of the importing country.

OMARs may also determine the form and content of the official assurances that can be issued for animal material or product, including live animals, hatching eggs, semen or embryos, which meet the specified requirements.

Where the OMAR determines the form and content of the official assurances, a separate export certificate template is available to authorised persons, recognised persons and registered exporters who have applied for access to the certificate templates, to facilitate the completion and issuing of the relevant official assurance. That template will be an amendable version of the form set in the OMAR.

Notices issued under section 60(1)(c) of the Animal Products Act to safeguard the assurances provided by New Zealand, and guidance in the form of Operational Codes, should be read in conjunction with this Notice.

This OMAR specifies the requirements that must be met by exporters of horses to be exported from New Zealand to or via the European Union and determines the form and content of the official assurance that must accompany the horses to be exported. It is based on:

- Regulation (EU) 2016/429 of the European Parliament and of the Council
- Commission Delegated Regulation (EU) 2020/692
- Commission Implementing Regulation (EU) 2021/403

#### Who should read this Animal Products Notice?

- Exporters of horses to the European Union.
- Exporters of horses to any destination when the horses will be transiting the European Union.

### Why is this important?

This Notice is important because it sets out the requirements that need to be met so that the Director-General of the New Zealand Ministry for Primary Industries (MPI) can certify that the horse meets the requirements for export to, or transit through, the European Union which New Zealand, in consultation with the authorities in the European Union, has determined will apply. It should be noted that although the Horse may comply with

these requirements and be given an official assurance (by way of a certificate), the importing country ultimately retains control over what horses it clears for entry.

# **Document History**

Version Date	Section Changed	Change(s) Description
1 November 2021	All sections (EUPER13 & EUTRA13)	New General Animal Health Law as described in EU Regulations 2016/429, with an accompanying model certificate as written in EU Regulations 2021/403.
15 September 2022	All sections (EUHOR13)	Merging of Horses to the European Union (EUPER13) and Horses Transiting the European Union (EUTRA13) into a single OMAR.
	Sections 1.2 and 1.3	Refinement of the definition of a registered horse and adding of measures to satisfy the health assurances required for such animals.
	Part 2	Amendment of the certificate template for the export of horses to the European Union according to updates to the model certificate published by the European Union. Amendments are limited to the notes section of the certificate.
	Part 3	Amendment of the certificate template for the export of horses transiting the European Union according to updates to the model certificate published by the European Union. Amendments are limited to the notes section of the certificate.
15 August 2024	Section 1.3	Amend the text to account for the removal of Part 3, and the changes to certification of horses transiting the European Union.
	Part 2	Amendment of the certificate template for the export of horses to the European Union according to updates to the model certificate published by the European Union. Included amendments are those provided for in Commission Implementing Regulation (EU) 2024/351 of 17 January 2024, and Commission Implementing Regulation (EU) 2024/399 of 29 January 2024. The change also means that the same model certificate now applies to horses with a destination in the European Union, and those that are transiting.
	Part 3	Deleted Part 3. Horses that are transiting the European Union must now be certified on the certificate in Part 2.

### Other information

#### Export non-conformances

Exporters should note that, under section 51 of the Animal Products Act 1999, where they have exported animal material or products, including live animals, hatching eggs, semen and embryos, that are refused entry by the foreign government they have a statutory duty to notify the Director-General of MPI not later than 24 hours after they have first knowledge of the event.

#### Liability

Section 61A of the Animal Products Act 1999 states that:

The Crown is not liable, and nor is the Director-General or any employee of the Ministry liable, for any loss arising through the refusal or failure of the relevant authority of an overseas market to admit export animal material or animal product to that market.

#### Related documents

OMAR documents can be downloaded from <u>https://www.mpi.govt.nz/export-requirements/omars-for-live-animals-semen-and-embryos/</u>

When you click on the + symbol on the right-hand side of any OMAR document, you can view the related information and documents (guidance document and export certificate template).

Export certificates for this OMAR, including translated versions where applicable, are provided for in documents made available according to the following naming convention: *Horses to the European Union* (*Export Certificate*) {*Ianguage*}. The export certificates are password-protected through a RealMe ® account.

# Part 1: Requirements

## 1.1 Application

- (1) This Notice applies to the export of live equine animals from New Zealand to the European Union.
- (2) This Notice also applies to the export of live equine animals from New Zealand to any destination where those live equine animals will be transported through the European Union.
- (3) This Notice applies to the following countries:
  - Austria, Belgium, Bulgaria, Croatia, Cyprus, the Czech Republic, Denmark, Estonia, Finland, France (including Reunion Island which is a Department of France), Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, the Netherlands, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, (EU member states).
  - b) Northern Ireland and Norway (non-EU member states).
    - i) Northern Ireland is not part of Great Britain and under the Brexit Northern Ireland Protocol will remain aligned with EU requirements.
    - ii) Horses cannot be exported directly into Switzerland. Horses can only be exported to Switzerland through another European Union country.

### 1.2 Definitions

(1) In this Notice, unless the context otherwise requires:

Act means the Animal Products Act 1999.

**Assembly operation** means the assembling of kept terrestrial animals from more than one establishment for a period shorter than the required residency period.

**Container** means any crate, box, receptacle or other rigid structure used for the transport of animals which is not the means of transport.

**Disease has not been reported** means that no animal or group of animals of relevant species kept on the establishment has been classified as a confirmed case of that disease and any suspect case of that disease has been ruled out.

**Equine animal** means an animal of species of solipeds belonging to the genus Equus (including horses, asses, and zebras) and the offspring of crossings of those species.

Listed third country, territory or zone thereof means a third country, territory or zone thereof included in a list of third countries, territories or zones thereof from which the entry into the Union of a particular species and category of animals.

Means of transport means road or rail vehicle, vessels and aircraft.

**Registered equine animal** means a purebred breeding animal of the species *Equus caballus* and *Equus asinus* entered or eligible for entry in the main section of a breeding book established by a breed society or breeding body recognised in accordance with Articles 4 or 34 of Regulation (EU) 2016/1012.

**Registered horse** means a kept animal of the species *Equus caballus* registered to compete in races or cultural equestrian events with an international association or organisation, either directly or through its national federation or branches, which manages horses for competition or racing. A registered horse may be exempted from certain requirements if they comply with additional guarantees. The exemption is based on the expectation that such horses will have a high level of health. Refer to *Horses to the European Union (Guidance)* for information on how to meet those guarantees.

**Sanitary group** means a group of listed third countries in which common animal health risks as regards diseases listed for equine animals prevail that require specific risk-mitigating measures and health guarantees when equine animals enter into the Union, New Zealand has been assigned by the European Union to Sanitary group A.

- (2) A term used in this Notice that is defined in the Act or the following Notices (or their successors) has the meaning given to it in the Act or that Notice:
  - a) Animal Products Notice: Official Assurance Requirements for Live Animals and Germplasm.
  - b) Animal Products Notice: Recognised Laboratories.

#### **1.3 Requirements for export**

- (1) Each horse exported from New Zealand must be accompanied by an official assurance in the form of a zoosanitary certificate, a sample version of which is included in Part 2, when it is:
  - a) exported to the European Union; or
  - b) exported to any other destination and is scheduled to be transported through the European Union to reach that destination.
- (2) A zoosanitary certificate must be completed and issued by an authorised person.
- (3) In order to issue a zoosanitary certificate, the authorised person must be satisfied that:
  - a) From the time of loading at the pre-export isolation facility until the time of export to the European Union, the horse has not been in contact with other terrestrial animals of:
    - i) the same species, not intended for entry into, or transit through, the European Union;
    - ii) other species susceptible to the same diseases, not intended for entry into, or transit through, the European Union; or
    - iii) a lower health status.
  - b) The means of transport used for the transport of the horse is:
    - i) constructed in such a way that:
      - 1) the horse cannot escape or fall out;
      - 2) visual inspection of the space where the horse is kept is possible; and
      - 3) the escape of excrements, litter or feed is prevented or minimised.
    - ii) cleaned and disinfected, with a disinfectant authorised by MPI, and dried or allowed to dry immediately before every loading of animals intended for entry into the European Union.
  - c) The containers in which the horse is to be transported to the European Union in the means of transport:
    - i) comply with the requirements in clause (3) b)i) of this Notice;
    - ii) contain only animals of the same species and category coming from the same establishment; and
    - iii) are cleaned and disinfected and dried or allowed to dry before loading of animals intended for entry into the European Union.
  - d) The horse was individually identified prior to being exported at least by one of the following methods:
    - i) an injectable transponder, or ear tag with a visible, legible and indelible display of:
      - 1) the identification code of the horse which establishes an unequivocal link between the horse and the accompanying animal health certificate; and
      - 2) the ISO-3166 two-digit alpha or three-digit numeric country code of the exporting country.

- ii) an identification document, issued at the latest at the time of certification for entry into the European Union, which:
  - describes and depicts the horse, including the alternative methods of identification, so as to establish an unequivocal link between the horse and the accompanying identification document; and
  - contains information on the individual code emitted by an implanted injectable transponder in the case where this code does not comply with the specifications in clause (3)d)i).
- e) The horse has not been vaccinated for Venezuelan equine encephalomyelitis for at least 60 days prior to scheduled date of export.
- f) The horse has not been vaccinated for African Horse Sickness.
- g) The proposed shipment otherwise meets the requirements of this Notice.

## 1.4 Specific requirements for the zoosanitary certificate

(1) Exporters intending to make use of the derogation for Registered horses in section II.3.2 of the certificate must apply for authorisation from MPI. See Horses to the European Union (Guidance) for details.

#### 1.5 Laboratories

(1) Where this Notice requires laboratory testing to be undertaken for official purposes the testing, unless otherwise stated must be done in laboratories recognised by MPI for this testing.

# Part 2: Zoosanitary Certificate



#### NEW ZEALAND MINISTRY FOR PRIMARY INDUSTRIES

# ANIMAL HEALTH/OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION OF EQUINE ANIMALS (MODEL "EQUI-X")

Name       I.3       Central competent authority         Address       I.3       Central competent authority         Ministry for Primary Industries       OR CODE         New Zealand       NZ         I.5       Consignee/Importer         Name       Name         Address       I.6         Operator responsible for the consignment         Name       Name         Address       Address         Country       ISO country code         Name       Address         Country       ISO country code         Name       ISO country code         I.7       Country of origin         New Zealand       NZ         I.7       Country of origin         New Zealand       NZ         I.8       Region of origin         NZ       I.9         Country of destination       ISO country code         I.8       Region of origin       Code         NZ-0       I.10       Region of destination         L11       Place of dispatch       I.12	COL	JNTRY	: NEW ZEALAND				Anima	l health/of	ficial certificate to the EU
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I.18     Transport conditions       I.19     Container number/Seal number			Identification			Coun	try	ISO co	untry code
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		I.18	Transport conditions						
Container No Seal No		I.19	Container number/Sea	l number					
			Container No		Seal	No			
I.20 Certified as or for		I.20	Certified as or for						
□ Further keeping □ Registered equine animal □ Registered horse			□ Further keeping	□ Registered equin	e anima	al	□ Registered horse		
I.21  For transit I.22  For internal market		I.21	For transit				I.22	et	
Third country ISO country code I.23			Third country	ISO country code			I.23		
I.25 Total quantity One (1)				I.25 Total quantit	ty One	(1)			

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# NEW ZEALAND Certificate model EQUI-X I.27 Description of consignment CN code Species Subspecies/Category Sex Identification System Identification number Age

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Certificate model EQUI-X

	II. He	alth inform	ation	II.a Certifi	icate reference	II.b IMSOC reference			
	П.	Animal h	health att	estation					
		I, the und	dersigned	official vete	rinarian, hereby certify that:				
	II.1.	I, the undersigned official veterinarian, hereby certify that: The equine animal described in Part I:							
		II.1.1.		tended for sl tion of infect	ghter in the framework of the mals, and:				
		<sup>(1)</sup> either	on Delegated Regulation (EU)						
		<sup>(1)</sup> <i>or</i>	[is a reg	gistered hors	e as defined in Article 2, point (12), of Delegated Regula	ation (EU) 2020/692;]			
		<sup>(1)</sup> <i>or</i>	[is an e	quine animal	other than a registered equine animal or a registered ho	rse;]			
		II.1.2.	Regulati <i>dd/mm/y</i> the last 4	on (EU) 201 <i>yyy</i> ) <sup>(2)</sup> , this c	or symptoms of diseases listed for equine animals in C 8/1882 during the clinical examination carried out on date being within the last 24 hours or, in the case of a regi on the last working day prior to the date of dispatch of t shment;	stered equine animal, within			
		II.1.3.		e requiremen fficial certifi	nts attested in points II.2. to II.5., and where applicable cate;	in point II.6., of this animal			
		II.1.4.			written declaration, signed by the operator responsible al health/official certificate.	for the animal, which is			
	II.2.	Attestation on third country, territory or zone thereof and in establishment of dispatch							
uc	II.2.1. The equine animal described in Part I is dispatched from New Zealand (insert name of territory, or zone thereof), a third country or territory, or zone thereof, which on the dat animal health/official certificate has the Code: <sup>(3)</sup> and is assigned to Sanitary Group								
Part II: Certification		II.2.2.	there ha African and ther	s been no cli horse sickne e have been r	escribed in Part I comes from a third country or territor inical, serological (in unvaccinated equine animals) or e ss during the last 24 months prior to the date of dispatc no systematic vaccinations against African horse sicknes lispatch of the animal to the Union.	pidemiological evidence of h of the animal to the Union			
Part I		II.2.3.		ine animal de thereof in w	escribed in Part I comes from an establishment situated in hich:	a third country or territory,			
		<sup>(1)</sup> either			<i>rolderia mallei</i> (glanders) has not been reported during the animal to the Union.]	e last 36 months prior to the			
		(1) <sub>or</sub>			ramme for infection with <i>Burkholderia mallei</i> (glanders) during the last 36 months prior to the date of dispatch of t				
			<sup>(1)</sup> either	L.	n with <i>Burkholderia mallei</i> (glanders) has not been repo during the last 36 months prior to the date of dispatch of				
			<sup>(1)</sup> or	the last 3	n with <i>Burkholderia mallei</i> (glanders) has been reported 66 months prior to the date of dispatch of the animal to ast outbreak, the establishment has remained under more	the Union and following the			
				<sup>(1)</sup> either	[until the date on which the remaining equine animal been subjected to a complement fixation test for infe <i>mallei</i> (glanders) <sup>(4)</sup> , carried out, with negative results a on samples taken at least 6 months after the date on have been killed and destroyed.]]]	ction with <i>Burkholderia</i> at a serum dilution of 1 in 5,			
				(1) <sub>0</sub> r	[for at least 30 days after the date on which the last an establishment was killed and destroyed, and the estal disinfected.]]]				
		II.2.4.		ine animal de thereof in w	escribed in Part I comes from an establishment situated in hich:	a third country or territory,			
		<sup>(1)</sup> either	[surra h Union.]		reported during the last 24 months prior to the date of d	ispatch of the animal to the			
		<sup>(1)</sup> <i>or</i>	[a surve	illance progr	amme for surra recognised by the Union <sup>(2)</sup> has been ca	rried out during the last 24			

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Certificate model EQUI-X

II. Health informa	tion	II.a Certific	ate reference	II.b IMSOC reference
	months p	prior to the d	ate of dispatch of the animal to the Union, and:	
	<sup>(1)</sup> either[		t been reported in the establishment during the last 24 m of the animal to the Union.]]	onths prior to the date of
	<sup>(1)</sup> or	of the ani	been reported in the establishment during the last 24 month mal to the Union, and following the date of the last out under movement restrictions:	
		<sup>(1)</sup> either	[until the date on which the remaining animals in the subjected to an enzyme-linked immunosorbent assay(E card agglutination test for trypanosomosis (CATT) at a carried out, with negative results, on samples taken at leas which the last infected animal has been removed from	LISA) for trypanosomosis a serum dilution of 1 in 4 ast 6 months after the date
		<sup>(1)</sup> <i>or</i>	[for at least 30 days after the date on which the last an establishment was either killed and destroyed or slaugh was cleaned and disinfected.]]]	
II.2.5.		ne animal de reof in which	scribed in Part I comes from an establishment situated in a n:	third country or territory,
<sup>(1)</sup> either	[dourine Union.]	has not been	n reported during the last 24 months prior to the date of o	dispatch of the animal to t
(1) <sub>0</sub> r			amme for dourine recognised by the Union <sup>(2)</sup> has been c late of dispatch of the animal to the Union, and:	arried out during the last
	<sup>(1)</sup> either		has not been reported in the establishment during the last 2 sh of the animal to the Union.]]	24 months prior to the dat
	<sup>(1)</sup> or	dispatch	has been reported in the establishment during the last 24 of the animal to the Union, and following the date of nent has remained under movement restrictions:	
		<sup>(1)</sup> either	[until the date on which the remaining equine animals in castrated male equine animals, have been subjected to for dourine, carried out with negative results at a seru samples taken at least 6 months after date on which the killed and destroyed or slaughtered, or the date on whi equine animals have been castrated.]]]	a complement fixation test im dilution of 1 in $5^{(4)}$ o infected animals have been
		<sup>(1)</sup> or	[for at least 30 days after the date on which the last anii establishment was either killed and destroyed or slaught was cleaned and disinfected.]]]	
II.2.6.	The equ	ine animal de	escribed in Part I comes from an establishment in which:	
<sup>(1)</sup> either		nfectious and al to the Uni	aemia has not been reported during the last 12 months pric on.]	or to the date of dispatch of
(1) <sub>0</sub> r	animal to		aemia has been reported during the last 12 months prior to and following the date of the last outbreak the establishm is:	
	<sup>(1)</sup> either	to an agar anaemia c interval o	date on which the remaining equine animals in the establis gel immuno-diffusion test (AGID or Coggins test) or EL arried out, with negative results, on samples taken on two f 90 days following the date on which the infected ani l or slaughtered, and the establishment was cleaned and	ISA <sup>(4)</sup> for equine infectiou occasions with a minimum nals have been killed an
	<sup>(1)</sup> or		st 30 days after the date on which the last animal of listed s r killed and destroyed or slaughtered, and the establ d.]]	•
II.2.7.	The equ	ine animal de	escribed in Part I comes from an establishment in which:	
	II.2.7.1.		with rabies virus in kept terrestrial animals has not been r to the date of dispatch of the animal to the Union;	reported during the last 3
	II.2.7.2.		ungulates has not been reported during the last 15 days p mal to the Union.	prior to the date of dispate

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Certificate model EQUI-X

II. Health inform	ation II.a	Certificate reference	II.b IMSOC referen
II.2.8.	been in cont in points II.2	ct with kept animals of listed species .2. to II.2.7.1 during the last 30 days requirement referred to in point II.2.	he operator, the equine animal described in Part I has s which did not comply with the requirements referred prior to the date of dispatch of the animal to the Un 7.2. during the last 15 days prior to the date of dispatc
II.3. Attestation	n of residence	and isolation prior to dispatch to the	e Union
<sup>(1)</sup> either [II.3.1.	of age, the e or zone ther	uine animal described in Part I has be	patch to the Union, or since birth if it is less than 40 of een continuously resident in the third country or terri country or territory, or zone thereof of dispatch fro {.]
<sup>(1)</sup> or [II.3.1.		st 40 days prior to the date of its disp gistered horse described in Part I:	patch to the Union, or since birth if it is less than 40 o
<sup>(1)</sup> either	[has bee	continuously resident in the third of	country or territory, or zone thereof of dispatch.]
(1) <sub>0</sub> r	[entered	he third country or territory, or zon	e thereof of dispatch on one or more occasions fro
	<sup>(1)</sup> either	[a Member State of the Europ	pean Union or Norway;]]]
	<sup>(1)</sup> and/or	registered horses, and from w or zone thereof of dispatch u accordance with Union legisla	or zone thereof authorised for entry into the Union which it was introduced into the third country or terri- under conditions at least as strict as those required ation for the entry of registered horses from that the thereof directly to the Union, and which is:
			ame Sanitary Group <sup>(3)</sup> as the third countr thereof of dispatch;]]]]
		<sup>(1)</sup> and/or [assigned to Sanita	ary Group A, B or C;]]]]
		<sup>(1)</sup> and/or [the United Arab ] Korea, Macao or S	Emirates, Bahrain, China <sup>(5)(6)</sup> , Hong Kong, Japan, So Singapore.]]]]
<sup>(1)</sup> <i>either</i> [II.3.2.		nimal described in Part I is dispatched roup A, B, C, D or G, and :	d from a third country or territory, or zone thereof assi
<sup>(1)</sup> either		st 30 days prior to the date of its dis ce entry from a Member State of the	patch to the Union, or since birth if it is less than 30 c Union or Norway,
	1		equine animals, except in case of a foal at foot of n a third country or territory, or zone thereof assigned
	t		ion from other equine animals, except in case of a fo ent situated in a third country or territory, or zone the r G.]]]
(1) <sub>0</sub> r	the last 30 d or since entr	ys prior to the date of its dispatch to y in accordance with point II.3.1 fro	tablishments under official veterinary supervision du the Union, or since birth if it is less than 30 days of om a Member State of the European Union, Norway is assigned to Sanitary Group A, B, C, D, E or G.]]
<sup>(1)(7)</sup> or [II.3.2.		nimal described in Part I is dispatched roup E, and:	d from a third country or territory, or zone thereof assi
<sup>(1)</sup> either	of age, or si Union, Norv	ce the date of entry in accordance	patch to the Union, or since birth if it is less than 40 o with point II.3.1 from a Member State of the Europ zone thereof which is assigned to Sanitary Group A, F
	<sup>(1)</sup> either	n isolation in a vector-protected est	tablishment.]]]
	t		terinary supervision, and the country or territory, or : the World Organisation for Animal Health (WOAH ess.]]]
<sup>(1)</sup> or	birth if it is le State of the	ss than 30 days of age, or since the da European Union, Norway or a third	the last 30 days prior to the date of its dispatch, or s ate of entry in accordance with point II.3.1 from a Mer country or territory, or zone thereof which is assigne ments under official veterinary supervision, and the

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NEW ZEALAN	EW ZI	EALAN
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II. Health inform	ation	II.a Cert	ificate reference	II.b IMSOC referen
			y, or zone thereof of dispatch to the Union is recogni ickness.]]	sed by the WOAH as officially
<sup>(1)(7)</sup> or [II.3.2.			rse described in Part I is dispatched from a third corry Group F, and:	ountry or territory, or zone the
<sup>(1)</sup> either		the last 4( hment.]]	) days prior to the date of dispatch it has been kep	t in isolation in a vector-prote
(1) <sub>0</sub> r	protecte	d establish	days prior to the date of dispatch to the Union it ha nment and constant monitoring of the vector protection vector-protected establishment.]]	
II.4. Attestation	of vaccind	ation and h	health tests	
<sup>(1)</sup> <i>either</i> [II.4.1.			described in Part I was not vaccinated against Africa he thereof of dispatch and there is no information s	
<sup>(1)</sup> or [II.4.1.			I described in Part I was vaccinated against African h f its dispatch to the Union.]	orse sickness more than 12 mo
<sup>(1)(7)</sup> or [II.4.1.	months in a thir of a cor period c a registe	and at leas d country on plete prino of validity of red vaccin	se described in Part I was vaccinated against Africa t 40 days prior to the date of introduction into the vec or territory, or zone thereof assigned to Sanitary Gro nary course of vaccination against African horse sic of the previous vaccination, by administration accord e which is protective against the circulating serotypes nation was applied on (insert date).]	ctor-protected establishment situ up F, and this vaccination consi kness, or a revaccination withir ing to manufacturer's instruction
II.4.2.			described in Part I has not been vaccinated against V days prior to the date of its dispatch to the Union, a	
<sup>(1)</sup> either		lomyelitis	n establishment situated in a third country or terr has not been reported during the last 24 months pr	
(1) <sub>0</sub> r	duringt date of	he last 6 m dispatch of	establishment in which Venezuelan equine encept onths prior to the date of its dispatch to the Union an f the animal described in Part I to the Union, all equin y healthy, and:	d during the last 21 days prior to
	<sup>(1)</sup> either	vector-p temperat	ine animal described in Part I has been kept protected rotected establishment, in which any equine animal th ure has been subjected with negative result to a virus lomyelitis <sup>(4)</sup> ; and the equine animal described in Pa	hat showed a rise in daily taken b isolation test for Venezuelan eq
		<sup>(1)</sup> either	[was vaccinated against Venezuelan equine er primary course and revaccinated according to m less than 60 days and not more than 12 months animal to the Union.]]]	anufacturer's recommendations
		<sup>(1)</sup> or	[was subjected to a haemagglutination inhibi encephalomyelitis <sup>(4)</sup> , carried out, with negative re- 14 days after the date of commencement of establishment.]]]	sult, on a sample taken not less
	<sup>(1)</sup> or	a rise o	y temperature of the equine animal described in Part I r the animal has been subjected to a virus isol lomyelitis with negative result, and the equine and d to:	ation test for Venezuelan eq
			<ul> <li>a haemagglutination inhibition test for Venez without an increase in antibody titre, carried o occasions with an interval of 21 days, the secon 10 days prior to the date of its dispatch to the</li> </ul>	ut on paired samples taken on d of which was taken during the
			<ul> <li>a reverse transcription-polymerase chain react Venezuelan equine encephalomyelitis virus gen out on a sample taken within the last 48 hours p</li> </ul>	ome <sup>(4)</sup> , with negative result, can
			<ul> <li>protection from vector attacks during the period loading for dispatch to the Union, by combine</li> </ul>	

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II. Health inform	II.a Certificate reference	II.b IMSOC refere
	and insecticides on the animal and disinsectization of	of the stable and the mea
(1)(3)	which it is transported.]]	
<sup>(1)(7)</sup> <i>either</i> [II.4.3.	The equine animal described in Part I is dispatched to the Union from Iceland free from equine infectious anaemia, where it was continuously resident sinc contact with equine animals which have entered Iceland from other third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the third contact with equine animals which have entered Iceland from the the third contact with equine animals which have entered Iceland from the	e birth, and did not come
<sup>(1)</sup> or [II.4.3.	The equine animal described in Part I was subjected with negative result to test (AGID or Coggins test) or to an ELISA for equine infectious anaemia <sup>(4)</sup> taken on	
<sup>(1)</sup> either	[the last 30 days prior to the date of its dispatch to the Union.]]	
(1)(7) <sub>0</sub> r	[the last 90 days prior to the date of its dispatch to the Union from a third thereof assigned to Sanitary Group A.]]	country or territory, or
<sup>(1)</sup> [II.4.4.	The equine animal described in Part I is dispatched from a third country or ter to Sanitary Group B, D or E or from China, or from a third country or ter <i>Burkholderia mallei</i> (glanders) has been reported during the last 36 months p to the Union, and was subjected to a complement fixation test for infecti (glanders) <sup>(4)</sup> carried out with negative result at a serum dilution of 1 in 5 	itory in which infection prior to the date of its disp on with <i>Burkholderia n</i> on a blood sample take
<sup>(1)</sup> [II.4.5.	The equine animal described in Part I is an uncastrated male or female equir dispatched from a third country or territory, or zone thereof assigned to Sa from China, or from a third country in which dourine has been reported duri the date of its dispatch to the Union, and was subjected to a complement fixe out with negative result at a serum dilution of 1 in 5 on a blood sample taken of <i>date</i> ), within the last 30 days prior to the date of its dispatch, to the Union, ar in Part I has not been used for breeding during 30 days prior to and after the	nitary Group B, D, E or ng the last 24 months pr tion test for dourine <sup>(4)</sup> cr on
<sup>(1)</sup> [II.4.6.	The equine animal described in Part I is dispatched to the Union from a thir thereof assigned to Sanitary Group E, from Bolivia, Brazil, Malaysia (Penins country or territory in which surra was reported during the last 24 months pri the Union, and was subjected to a card agglutination test for trypanosomos negative result at a serum dilution of 1 in 4 on a blood sample taken on within the last 30 days prior to the date of its dispatch to the Union.]	ula), Uruguay, or from a or to the date of its dispat is (CATT) <sup>(4)</sup> , carried out
<sup>(1)(7)</sup> [II.4.7.	The equine animal described in Part I is dispatched to the Union from a thir thereof which is assigned to Sanitary Group E, and:	d country or territory, or
<sup>(3)</sup> either	[was subjected to an indirect ELISA or a blocking ELISA for African horse out by the same laboratory on the same day on blood samples taken on two between 21 and 30 days, on	occasions with an interv (insert date,
	<sup>(3)</sup> <i>either</i> [with negative results in each case.]]]	
	<sup>(3)</sup> or [with a positive result in the first sample, and:	
	<sup>(3)</sup> <i>either</i> [the second sample was subsequently tested with neg PCR <sup>(8)</sup> .]]]	ative result in a real time
	<sup>(3)</sup> or [the two samples were tested without more than a two in a virus neutralisation test as described in the Terrestrial Manual for Diagnostic Tests and Vaccine	atest edition of the W
(1) <i>or</i>	[was subjected to an indirect ELISA or a blocking ELISA for African horses on a blood sample taken on ( <i>insert date</i> ), within the last 2 dispatch to the Union, and the third country or territory of dispatch is recogni- free of African horse sickness.]]	days prior to the date
(1) <sub>0</sub> r	[it is a registered horse not vaccinated against African horse sickness and d third country or territory, or zone thereof which is recognised by the WOAI horse sickness.]]	
<sup>(1)(7)</sup> [II.4.8.	The equine animal described in Part I is dispatched to the Union from a thir thereof assigned to Sanitary Group F, and:	d country or territory, or
<sup>(1)</sup> either	[was subjected to an indirect ELISA or a blocking ELISA for African horses same laboratory on the same day on blood samples taken on two occasions wand 30 days, on	vith an interval of betwee

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Cumunan	mouti	EQUI-A

n. neath morn	ation	II.a Certific	cate reference	II.b IMSOC refere
			is after the date of introduction into the vector-protected the last 10 days prior to the date of its dispatch to the Un	
	<sup>(1)</sup> either	[with neg	ative results in each case.]]]	
	<sup>(1)</sup> <i>or</i>	[with a po	ositive result in the first sample, and:	
		<sup>(1)</sup> either	[the second sample was subsequently tested with negative PCR <sup>(8)</sup> .]]]]	e result in a real-time
		<sup>(1)</sup> <i>or</i>	[the two samples were tested without more than a two-fo in a virus neutralisation test as described in the late Terrestrial Manual for Diagnostic Tests and Vaccines.]	st edition of the We
(1) <sub>0</sub> r	sickness date) no	s <sup>(8)</sup> carried ou t less than 28	indirect ELISA or a blocking ELISA and a real-time t with negative result in each case on a blood sample taken days after the date of introduction into the vector-protect t to the date of its dispatch to the Union.]]	on
(1) <sub>0</sub> r	sample	taken on	al-time RT-PCR for African horse sickness <sup>(8)</sup> , carried out w 	date of introduction in
II.5. Attestation	of the tra	nsport condit	tions	
<sup>(1)(7)</sup> either[II.5.1.	thereof a directly contact	assigned to S to the Union with other equ	escribed in Part I is dispatched to the Union from a third co anitary Group A, B, C, D, E or G and arrangements have , without subjecting the animal to any assembly operation uine animals not complying with at least the same health re ficial certificate.]	been made to transp n and without coming
<sup>(1)(7)</sup> or [II.5.1.	Sanitary establisl	Group F an nment withou	hed to the Union from a third country or territory, or zone t d arrangements have been made to transport it directly t coming into contact with other equine animals not compl as described in this animal health/official certificate:	from the vector prote
<sup>(1)</sup> either		l and disinfec	vector-protected conditions and arrangements have been ted in advance with a disinfectant officially recognised in the	
(1) <sub>0</sub> r	have bee into a po of equin	en made to tra ort situated in ie animals, in	pountry or territory, or zone thereof under vector-protected consport it on a vessel which is scheduled directly to a port in a third country or territory, or zone thereof not approved f stalls which were cleansed and disinfected in advance w d country or territory of dispatch.]]	the Union without ca or the entry into the U
II.5.2.	with at	least the same	een made and verified to prevent any contact with other eq e health requirements as described in this animal health/c of certification until the date of dispatch of the animal to	official certificate duri
II.5.3.	before le	oading of the or territory o	is or containers in which the animal is going to be loaded w animals for dispatch to the Union with a disinfectant offic f dispatch and are so constructed that facces, urine, litter or	cially recognised in the
<sup>(1)(9)</sup> [II.6. <b>Public</b>	health a	ttestation [D	elete when the Union is not the final destination of the an	imals]
			narian, hereby certify, that the equine animal described in	
II.6.1.	in the th	ird country c	or territory of dispatch to the Union has not received:	
		any stilbene o pestrogenic, a	bstances listed in Table 2 of the Annex to Commission Re or thyrostatic substances; indrogenic, gestagenic or beta-agonist substances for purp al treatment (as defined in Council Directive 96/22/EC);	
II.6.2.	accorda dispatch	nce with Art ied from a t	s covering equine animals provided by the control plan icle 6(2) of Commission Delegated Regulation (EU) 2 hird country or territory listed for equine animals in tion (EU) 2021/405.	022/2292 and it has

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#### Certificate model EQUI-X

II. Hea	alth information	II.a Certificate reference	II.b IMSOC reference		
	antimicrobial me an antimicrobial down in Comm Regulation (EU)	d official veterinarian, hereby certify that the animals described dicinal products for growth promotion or yield increase or antim that is included in the list of antimicrobials reserved for the treatm ssion Implementing Regulation (EU) 2022/1255, as set out in 2023/905 and originate from a third country or region thereof lis ation (EU) 2023/905.]]	nicrobial medicinal products contain nent of certain infections in humans Article 3 of Commission Delega		
Notes:					
	imal health/official l destination of the	certificate is intended for the entry into the Union of equine ani animals.	mals, including when the Union is		
Europe Ireland	an Union and the E in conjunction wi	greement on the withdrawal of the United Kingdom of Great B uropean Atomic Energy Community, and in particular Article 5( th Annex 2 to that Protocol, references to Union in this animal et of Northern Ireland.	4) of the Protocol on Ireland/North		
		certificate shall be completed in accordance with the notes for t I to Commission Implementing Regulation (EU) 2020/2235.	he completion of certificates provi		
Part I:					
Box ref	Terence I.6: P	rovide the information on the operator responsible for the anim	nal.		
Box ref		rovide the code of the third country or territory, or zone thereof column 2 of the table in Part 1 of Annex IV to Commission Imp			
Box re	ic ic th th	dentification system ": The animal shall be individually ide entification laid down in Article 21(2), point (a), of Delegate entified by an alternative method provided it is recorded in the e animal as referred to in Article 21(2), point (b)(i), of Delegate e identification system and the anatomic place used on the anima s number shall be stated and the name of the competent author	ed Regulation (EU) 2020/692, or e identification document (passport) d Regulation (EU) 2020/692. Spec I. If a passport accompanies the anir		
Part II:					
(1)	Delete if not appl	icable.			
(2)	at the border co	/official certificate shall be issued within the last 10 days prior to ntrol post; in the case of transport by sea, the period may be the duration of the journey by sea.			
	entry into the Uni period where res from that third co	e Union shall not be allowed when the animal was loaded either on from the respective third country or territory, or zone thereof rictive measures have been adopted by the Union against the er untry or territory, or zone thereof. Check against columns 8 and 9 gulation (EU) 2021/404.	f referred to in point II.2.1, or durin ntry into the Union of equine anin		
(3)		ry, territory or zone thereof and the Sanitary Group as appearing Annex IV to Implementing Regulation (EU) 2021/404.	respectively in columns 2 and 3 of		
(4)	European Unio	e, surra, dourine, equine infectious anaemia and Venezuelan equi n Reference Laboratory for Equine Diseases other ses.fr/en/minisite/equine-diseases/sop.			
(5)		country or territory authorised for entry into the Union as appear of Annex IV to Implementing Regulation (EU) 2021/404.	ring respectively in columns 2 and 5		
(6)	Only authorised	f the third country or territory of dispatch is assigned to Sanita	ary Group G.		
(7)					
(8)		horse sickness described by the European Union Reference La.gob.es/en/ganaderia/temas/laboratorios/referencia-union-euro			
(9)		oint, the equine animal, if intended for free circulation in accord	ance with the customs procedures l uncil (OJL 269, 10.10.2013, p.1),		

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II. Health information	on II.a Certificate reference	
		II.b IMSOC reference
be excluded	from slaughter for human consumption in the iden	tification document issued in accordance with Un
animal health	rules.	
(10) Applicable to	consignments entering the Union as from 3 Septe	mber 2026.
Official veterinar		
Official veterinar Name (in capital le		
Name (in capital le	tters)	
	tters)	tion and title
Name (in capital le Date	(Qualifica	
Name (in capital le	tters)	

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Certificate model EQUI-X

Declaration by the operator responsible for entry into the Union of the consignment of equine animal								
	ication of the animal <sup>(1)</sup>	¥.410 .1	X1		2			
Specie	s (Scientific name)	Identification system	Identification number	Age	Sex			
I, the	undersigned operator of	of the equine animal described	above, hereby declare, that:					
- the equine animal								
©either	either [has remained in (insert name of third country or territory, or zone thereof of dispatch to the Union) during a at least 40 days prior to the date of dispatch to the Union, or since birth, or since entry from the European Union or Norway;]							
<sup>(2)</sup> OF	[entered(insert name of third country or territory, or zone thereof of dispatch to the Union) during the required residence period of at least 40 days prior to the date of dispatch to the Union:							
	(a) on(insert date) from(insert name of third country or territory from where the horse entered the third country or territory, or zone thereof of dispatch to the Union)							
	(b) on(insert date) from(insert name of third country or territory from where the horse entered the third country or territory, or zone thereof of dispatch to the Union)							
(c) on(insert date) from(insert name of third country or territory from where the horse entered the third country or territory, or zone thereof of dispatch to the Union);]								
- during the last 15 days prior to the date of dispatch to the Union the equine animal has not been in contact with animals suffering from infectious or contagious diseases transmissible to equine animals;								
- the conditions for residence and isolation prior to dispatch to the Union as applicable in accordance with point II.3. of the accompanying animal health/official certificate for the third country or territory, or zone thereof of dispatch to the Union are fulfilled;								
- the conditions for the transport as applicable in accordance with point II.5. of the accompanying animal health/official certificate for the third country or territory, or zone thereof of dispatch to the Union are fulfilled;								
- I am aware of the animal health and veterinary certification requirements for the movement of equine animals from one Member State of the Euopean Union to another laid down in Commission Delegated Regulation (EU) 2020/688;								
- the equine animal is scheduled to leave the European Union on								
Name and address of the operator:								
Date:								
(Signature)								
(**********								
<sup>(1)</sup> Identification system: The animal shall be individually identified with one of the methods of identification laid down in Article 21(2), point (a), of Delegated Regulation (EU) 2020/692, or be identified by an alternative method provided it is recorded in identification document (passport) of the animal as referred to in Article 21(2), point (b)(i), of Delegated Regulation (EU) 2020/692. Specify the identification system (such as ear tag, transponder) and the anatomic place used on the animal.								
If a passport accompanies the animal, its number shall be stated and the name of the competent authority which validated it.								
Age: Date of birth (dd/mm/yyyy).								
Sex (M = male, $F = female$ , $C = castrated$ ).								
<sup>(2)</sup> Delete if not applicable.								

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