ANNEXES

ANNEX I

Part 1

Technical specifications of electronic means of identification for equine animals

1. Where applied to equine animals, the electronic means of identification referred to in points (c), e and (f) of Annex III to Delegated Regulation (EU) 2019/2035 (the electronic means of identification) shall display:

   (a) a three-digit ISO-3166 compatible country code;

   (b) a numeric individual animal code of 12 digits.

2. The electronic **means of identification** must be:

   (a) read-only passive transponders applying HDX or FDX-B technology, complying with ISO standards 11784 and 11785;

   (b) readable by reading devices, complying with ISO standard 11785, capable of reading HDX and FDX- B transponders.

3. The electronic means of identification must be readable at the minimum reading distance of:

   (a) 12 centimetres for ear tags when read with a portable reader;

   (b) 15 centimetres for injectable transponders when read with a portable reader.

4. The electronic means of identification must have been tested with favourable results as regards the following:

   (a) conformance with the ISO standards 11784 and 11785, in accordance with the method referred to in point 7 of the ISO standard 24631-1;

   (b) achievement of minimum performance on reading distances referred in point 3 of this Part, in accordance with the procedures referred to in point 7 of the ISO standard 24631-3.

5. Alternative composition of the code referred to in point 1(b):

   (a) a two-digit species code;

   (b) a two-digit manufacturer code;

   (c) a numeric-individual animal code of 8 digits.

Part 2

Technical specifications of means of identification for equine animals

1. The means of identification referred to in points (a) and (b) of Annex III to Delegated Regulation (EU) 2019/2035 for equine animals shall be:

   (a) non-reusable;

   (b) of non-degradable material;

   (c) tamper-proof;

   (d) easy to read throughout the lifetime of the equine animals;

   (e) designed in such way that they can remain securely attached to the equine animals without being harmful to it;

   (f) easily removable from the food chain.

2. The means of identification referred to in point 1 shall carry one of the following non-removable inscriptions:

   (a) a three-digit ISO-3166 compatible country code;

   (b) a numeric individual animal code of at least 12 digits.

3. The means of identification referred to in point 1 may carry other information, if authorised by the competent authority, provided that the inscriptions referred to in point 2 remain visible and legible.

ANNEX II

Part 1
Content of the single lifetime identification document

DOCUMENT D'IDENTIFICATION DES ÉQUIDÉS

Ces instructions sont rédigées en vue d'assister l'utilisateur et n'entravent pas l'application des règles établies par le règlement d'exécution (UE) 2021/… [*Reference to present Regulation*].

I. Le document d'identification doit comporter toutes les instructions nécessaires à son utilisation ainsi que les coordonnées de l'organisme émetteur en français, en anglais et dans une des langues officielles de l'État membre ou du pays tiers dans lequel l'organisme émetteur a son siège.

II. Le document d'identification doit contenir les renseignements suivants :

1. Section I - Identification

L'équidé doit être identifié par l'autorité compétente ou par l'organisme émetteur ou la personne physique visés à l'article 5, paragraphe 1, du règlement d'application (UE) n° 2021/... [*Référence au présent règlement*]. Le numéro unique d'identification valable à vie doit permettre d'identifier clairement l'équidé ainsi que la base de données établie par l'autorité compétente ou l'organisme émetteur qui a délivré le document d'identification et doit être compatible avec le numéro universel d'identification des équidés (UELN).

Dans la description à la partie A de la section I, notamment au point 3, l'utilisation d'abréviations doit être évitée autant que possible. Au point 5 de la partie A de la section I, un champ doit être prévu pour insérer au moins quinze chiffres du code transmis par le transpondeur.

A la partie B de la section I le signalement graphique doit être renseigné en utilisant un stylo à bille à encre rouge pour les marques et un stylo à bille à encre noire pour les épis, ou en conséquence si complété par voie électronique, en tenant compte des lignes directrices fournies par la Fédération Équestre Internationale (FEI) ou par Weatherbys.

La partie C de la section I doit être utilisée pour enregistrer toute rectification aux détails d'identification.

2. Section II – Administration de médicaments vétérinaires

Les parties I et II ou la partie III de cette section doivent être dûment complétées suivant les instructions établies dans cette section.

3. Section III – Marque de validation/Licence

Nécessaire pour les mouvements conformément à l'article 92, paragraphe 2, du règlement délégué (UE) 2020/688.

4. Section IV – Certificat zootechnique

Si l'équidé est inscrit ou enregistré et admissible à l'entrée dans un livre généalogique tenu par une organisme de sélection, le document d'identification doit indiquer le pedigree ainsi que la classe du livre généalogique dans laquelle l'équidé est inscrit conformément aux règles du organisme de sélection qui délivre le certificat zootechnique.

5. Section V – Propriétaire

Le nom du propriétaire ou de son agent ou représentant doit être mentionné si l'autorité compétente, l'organisme émetteur le requiert ou l'organisation qui gère les chevaux enregistrés en vue des compétitions ou courses le requiert.

6. Section VI – Enregistrement des contrôles d'identité

À chaque fois que les lois et règlements l'exigent, l'identité de l'équidé doit faire l'objet d'une vérification enregistrée par l'autorité compétente, au nom de l'organisme émetteur ou de l'organisation qui gère les chevaux enregistrés en vue des compétitions ou courses.

7. Section VII and VIII – Enregistrement des vaccinations

Toutes les vaccinations doivent être enregistrées à la section VI (grippe équine seulement) et à la section VII (toutes les autres vaccinations). Ces informations peuvent être fournies par l'apposition d'un autocollant.

8. Section IX – Examen de laboratoire

Les résultats de tous les examens pratiqués pour déceler une maladie transmissible peuvent être consignés.

9. Section X – Châtaignes *(en option)*

Cette section est nécessaire au respect du modèle de document d'identification de la Fédération Equestre Internationale (FEI).

III. Sauf s'il est détruit sous surveillance officielle à l'abattoir, le document d'identification doit être retourné à l'organisme émetteur après que l'animal est mort, a dû être détruit, a été perdue ou volée ou a été abattu à des fins de contrôle de la maladie.

IDENTIFICATION DOCUMENT FOR EQUIDAE

These instructions are drawn up to assist the user and do not impede on the rules laid down in Implementing Regulation (EU) 2021/… [*Reference to present Regulation*]

I. The identification document must contain all the instructions needed for their use and the details of the issuing body in French, English and one of the official languages of the Member State or third country where the issuing body has its headquarters.

II. The identification document must contain the following information:

1. Sections I – Identification

The equine animal shall be identified by the competent authority or by the issuing body or natural person as referred to in Article 5(1) of Implementing Regulation (EU) 2021/… [*Reference to present Regulation*]. The unique code shall clearly identify the equine animal and the database established by the competent autority or issuing body which issued the identification document and shall be compatible with the universal equine life number (UELN).

In the narrative in Part A of Section I, in particular in point 3 thereof, abbreviations must be avoided, where possible. In point 5 of Part A of Section I, the space must be provided for at least 15 digits of the transponder code.

In Part B of Section I the outline diagram shall be completed using red ball point ink for marks and black ball point ink for whorls, or accordingly if completed electronically, taking into account the guidelines provided for by the International Federation for Equestrian Sports (FEI) or the Weatherbys.

Part C of Section I must be used to record modifications to identification details.

2. Section II – Administration of veterinary medicinal products

Parts I and II or Part III of this Section must be duly completed in accordance with the instructions set out in this Section.

3. Section III – Validation mark/Licence

Required for movements in accordance with Article 92(2) of Delegated Regulation (EU) 2020/688.

4. Sections IV – Zootechnical certificate

If the equine animal is entered or registered and eligible for entry in a breeding book maintained by a breed society, the identification document shall contain the pedigree and the breeding book class in which the equine animal is entered in accordance with the rules of the breed society issuing the zootechnical certificate.

5. Section V – Owner

The name of the owner or its agent or representative must be stated where required by the competent autority, issuing body or the organisation which manages registered horses for competitions or races.

6. Section VI – Recording of identity checks

Whenever laws and regulations so require, checks conducted on the identity of the equine animal must be recorded by the competent authority, on behalf of the issuing body or by the organisation which manages registered horses for competitions or races.

7. Sections VII and VIII – Vaccination record

All vaccinations must be recorded in Section VII (equine influenza only) and in Section VIII (all other vaccinations). The information may take the form of a sticker.

8. Section IX – Laboratory health tests

The results of all tests carried out to detect transmissible diseases may be recorded.

9. Section X – Chestnuts (optional)

This section shall be required for compliance with the model of the identification document of the International Federation for Equestrian Sports (FEI).

III. Except where it is destroyed under the offical supervision at the slaughterhouse, the identification document must be returned to the competent authority or issuing body after the animal has died, had to be destroyed, was lost or stolen or was slaughtered for disease control purposes.

/ official language

SECTION I
Partie A - Détails d'identification
Part A – Identification details
/ *official language*

(1)(a) Espèce:
Species:
/ *official language*

(1)(b) Sexe:
Sex:
/ *official language*

(2)(a) Date de naissance:
Date of birth:
/ *official language*

(2)(b) Lieu et pays de naissance:
Place and country of birth :
/ *official language*

2(c) Nom (optionnel):
Name (optional):
/ *offical language*

(3) Signalement:
Description:
/ *official language*

(3)(a) Robe:
Colour:
/ *official language*

(3)(b) Tête:
Head:
/ *official language*

(3)(c) Ant. G:
Foreleg L:
/ *official language*

(3)(d) Ant. D:
Foreleg R:
/ *official language*

(3)(e) Post G:
Hind leg L:
/ *official language*

(3)(f) Post D:
Hind leg R:
/ *official language*

(3)(g) Corps:
Body:
/ *official language*

(3)(h) Marques:
Markings:
/ *official language*

(9) Date/Date/ *official language:*

(10) Lieu/Place/ *official language*:

(4) Code Unique ou Numéro unique d'identification valable à vie (15 chiffres):
Unique Code or lifer number: (15 digits):
/ *official language*
□□□-□□□-□□□□□□□□□□
Code-barres (optionnel)
Bar-Code (optional)
/ *official language*

(5) Code du transpondeur (si disponible)
Transponder code (where available)
/ *official language*
□□□ □□□ □□□ □□□ □□□
Système de lecture (si différent de ISO 11784)
Reading system (if not ISO 11784)
/ *official language*
Code-barres (optionnel)
Bar-Code (optional)
/ *official language*

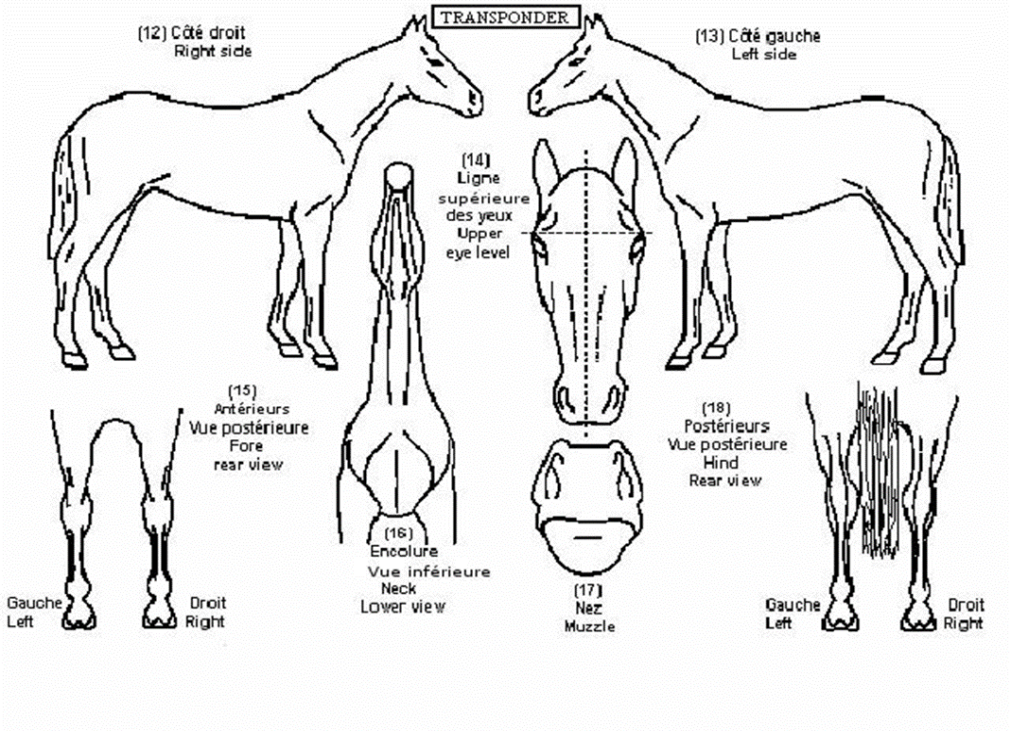
(6) Méthode alternative de vérification d'identité (si applicable)/Alternative method for identity verification (if applicable)/ *official language*:

(7) Information sur toute autre méthode appropriée donnant des garanties pour vérifier l'identité de l'animal (groupe sanguin / code ADN) (optionnel)/ Information on any other appropriate method providing guarantees to verify the identity of the animal (blood group/DNA code) (optional)/ *official language*:

(8) **Nom et adresse du destinataire du document/Name and address of person to whom document is issued/** *official language***:**

(11) Signature de la personne qualifiée (nom en lettres capitales)/Signature of qualified person (name in capital letters)/ *official language*
Cachet de l'organisme émetteur ou de l'autorité compétente/ stamp of issuing body or competent authority/ *official language*

Partie B – Signalement graphique
Part B – Outline Diagram
/ official language



Signature et chachet du vétérinair ou de la personne qualifée or de l'authité compétente (nom en lettres capitales)
Signature and stamp of the veterinarian or the qualified person or the competent authority (name in capital letters)
/ offical language

Note for the competent authority or issuing body [*not to be printed in identification document*]: Slight variations from this model outline diagram are permitted, provided they were in use before the date of application of this Regulation.

Partie C – Castration, vérification de la description, autres modifications
Part C – Castration, verification of the description, other modifications
/ *official language*

<table>
<tr><td>Castration/Castration/ official language</td><td colspan="2">Identification/Identification/ official language</td></tr>
<tr><td rowspan="4">Date et lieu de la castration/Date and place of castration/ official language

Signature et cachet du vétérinaire/Signature and stamp of veterinarian/ official language</td><td>Vérification de la description/Verification of the description / official language
Mentionner/Include/ official language:</td><td rowspan="4">Signature de la personne qualifiée (nom en lettres capitales)/Signature of qualified person (name in capital letters)/ official language
Cachet de l’organisme émetteur ou de l’autorité compétente/ Stamp of issuing body or competent authority/ official language
Date et lieu/Date and place/ official language</td></tr>
<tr><td>1. Rectifications/ Amendments/ official language</td></tr>
<tr><td>2. Adjonctions/ Additions/ official language</td></tr>
<tr><td>3. Enregistrement d'un document d' identification dans la base de données d'un organisme émetteur autre que l' organisme qui a délivré le document/ Registration of an identification document in the database of an issuing body other than the body which issued the original document/ official language</td></tr>
</table>

Note for the competent authority or issuing body [*not to be printed in identification document*]: Slight variations from this model are permitted, provided they were in use before the date of application of this Regulation. Part C of Section I may be completed in hand-writing

SECTION II

Administration de médicaments vétérinaires
Administration of veterinary medicinal products
/ *official language*

Code Unique /Unique Code/*official language*:
□□□-□□□-□□□□□□□□□

Partie/Part/*official language* I

Date et lieu de délivrance de la présente section[1]/Date and place of issue of this Section[1]/ *official language*:.....

Organisme émetteur de la présente section du document d'identification[1]/Issuing body for this Section of the identification document[1]/ *official language*: .....

Partie/Part/official language II

Remarque/ Note/ *official language*	**L'équidé n'est pas destiné à l'abattage pour la consommation humaine, et par conséquent, l'équidé peut recevoir des médicaments vétérinaires autorisés conformément à l'article 8, paragraphe 4, du règlement (UE) 2019/6 ou administrés conformément à l'article 112, paragraphe 4, du ledit règlement. /** **The equine animal is not intended for slaughter for human consumption, and may therefore undergo the administration of veterinary medicinal products authorised in accordance with Article 8(4) of Regulation (EU) 2019/6 or administered in accordance with Article 112(4) of that Regulation./ *official language***

Déclaration / Declaration / *official language*	**L'animal équine décrit dans le présent document d'identification n'est pas destiné à l'abattage pour la consommation humaine./ The equine animal described in this identification document is not intended for slaughter for human consumption/ *official language***	
Date et lieu/ Date and place/ *official language:*	Vétérinaire responsable procédant conformément à l'article 112, paragraphe 4, du règlement (UE) 2019/6/Veterinarian responsible acting in accordance with Article 112(4) of Regulation (EU) 2019/6[2]/ *official language* :	Nom (en lettres capitales) et signature du vétérinaire responsable/ Name (in capital letters) and signature of the veterinarian responsible/ *official language*
	Autorité compétente[2] ou organisme émetteur[2]/ Competent authority[2] or issuing body[2] / *official language*	Nom (en lettres capitales) et signature de la personne responsable[2]/ Name (in capital letters) and signature of the person responsible[2]/ *official language*

Partie/Part/*official language* III

Remarque/Note/ *official language*: **L'équidé est destiné à l'abattage pour la consommation humaine./The equine animal is intended for slaughter for human consumption. / *official language***

Sans préjudice du règlement (CE) nº 470/2009 ni de la directive 96/22/CE, l'équidé peut faire l'objet d'un traitement médicamenteux conformément à l'article 115, paragraphe 1, du règlement (UE) 2019/6 à condition que l'équidé ainsi traité ne soit abattu en vue de la consommation humaine qu'au terme d'un temps d'attente général de six mois suivant la date de la dernière administration de substances listées conformément à l'article 115, paragraphe 5, du ledit règlement./Without prejudice to Regulation (EC) No 470/2009 and Directive 96/22/EC, the equine animal may be subject to medicinal treatment in accordance with Article 115(1) of Regulation (EU) 2019/6 under the condition that the equine animal so treated may only be slaughtered for human consumption after the end of the general withdrawal period of six months following the date of last administration of the substances listed in accordance with Article 115(5) of that Regulation./ *official language*.

ENREGISTREMENT DE LA MÉDICATION/ MEDICATION RECORD/*offical language*			
Date de la dernière administration, telle que prescrite, conformément à l'article 115, paragraphe 1, du règlement (UE) 2019/6[(2)]/Date of last administration, as prescribed, in accordance with Article 115(1) of Regulation (EU) 2019/6[(2)]/ *official language*	Substance(s) fondamentale(s) incorporée(s) dans le médicament vétérinaire administré conformément à l'article 115, du règlement (UE) 2019/6[(2)], comme mentionné dans la première colonne[(2)(3)(4)]/ Essential substance(s) incorporated in the veterinary medicinal product administered in accordance with Article 115 of Regulation (EU) 2019/6 as mentioned in the first column[(2)(3)(4)]/ *official language*	Vétérinaire responsable appliquant et/ou prescrivant le traitement médicamenteux/ Veterinarian responsible applying and/or prescribing administration of veterinary medicinal product/ *official language*	
		Nom/Name/ *official language*:[5] .... ……. Adresse/ Address/ *official language*:[5]..... ..... Code postal/ Postal code/ *official language*:[5] ……... Lieu/Place/ *official lan*guage:[5] ……..…… …… Téléphone/ Telephone/ *official language*:[6]	Signature/ Signature/ *official language*

		Nom/Name/ *official language*:[5] .... ……. Adresse/ Address/ *official language*:[5]..... ..... Code postal/ Postal code/ *official language*:[5] ……... Lieu/Place/ *official lan*guage:[5] ……..…… …… Téléphone/ Telephone/ *official language*:[6] ……..	Signature/ Signature/ *official language*

Partie/Part/*official language* IV[(7)]

Remarque/Note/ *official lang*uage: **Les échanges des équidés enregistrés auxquels ont été administrés des médicaments vétérinaires contenant du trembolone allyle ou des substances beta-agonistes aux fins indiquées à l'article 4 de la Directive 96/22/CE peuvent s'effectuer avant la fin de la période d'attente, conformément à l'article 7, paragraphe 1, de la Directive 96/22/CE/ Trade in registered equidae to which veterinary medicinal products containing allyl trenbolone or beta-agonists have been administered for the purposes referred to in Article 4 of Directive 96/22/EC, may take place before the end of the withdrawal period, in accordance with Article 7(1) of Directive 96/22/EC/** *official language*

Date de la dernière administration conformément à l'article 4 de la Directive 96/22/CE/ Date of last administration in accordance with Article 4 of Directive 96/22/EC/ *offical language*	Substance(s) incorporée(s) dans le médicament vétérinaire administré conformément à l'article 4 Directive 96/22/CE/ Substance(s) incorporated in the veterinary medicinal product administered in accordance with Article 4 of Directive 96/22/EC/ *offical language*	Vétérinaire appliquant et/ou prescrivant le traitement médicamenteux/ Veterinarian applying and/or prescribing administration of veterinary medicinal product/ *official language*

		Nom/Name/ *official language*:[5] .... ……. Adresse/ Address/ *official language*:[5]..... ..... Code postal/ Postal code/ *official language*:[5] ……... Lieu/Place/ *official lan*guage:[5] ……..…… …… Téléphone/ Telephone/ *official language*:[6]	Signature/ Signature/ *official language*

Partie/Part/*official language* V[(8)]

Remarque/Note/ *official lang*uage:	**L'équidé est destiné à l'abattage pour la consommation humaine./The equine animal is intended for slaughter for human consumption. / *official language***

L'abattage de l'équidé est pour des raisons administratives retardé d'au moins six mois conformément à l'article 38, paragraphe 2(b) du règlement d'exécution (UE) 2021/ …… [*Reference to this Regulation*]/ The slaughter of the equine animal is for administrative reasons delayed for at least six months in accordance with Article 38(2)(b) of Implementing Regulation (EU) 2021/…. [*Reference to this Regulation*]/ *official lang*uage

Date de la suspension / Date of suspension/ *official language*	Lieu/ Place/ *official language*	Autorité compétente[2] ou organisme émetteur[2]/ Competent authority[2] or issuing body[2] / *official language*	Nom (en lettres capitales) et signature de la personne responsable[2]/ Name (in capital letters) and signature of the person responsible[2]/ *official language*

1 Information à ne fournir que si la présente section est délivrée à une autre date que la section I./Information only required if this Section is issued at a different date than Section I. / *official language*

2 Biffer les mentions inutiles./ Cross out what is not applicable./ *official language*

3 Il est indispensable de spécifier les substances en se fondant sur la liste de substances établie conformément à l'article 115, paragraphe 5, du règlement (UE) 2019/6./ Specification of substances against list of substances established in accordance with Article 115(5) of Regulation (EU) is compulsory./ *official language*

4 Les informations relatives à d'autres médicaments vétérinaires administrés conformément au règlement (UE) 2019/6 sont facultatives./Information on other veterinary medicinal products administered in accordance with Regulation (EU) 2019/6 is optional./ *official language*

5 Nom, adresse, code postal et lieu (en lettres capitales)./Name, address, postal code and place (in capital letters)./ *official language*

6	Numéro de téléphone selon le modèle [+ code pays (code régional) numéro]./Telephone in format [+country code (regional code) number]./ *official language*
7	La partie IV doit être complétée conformément à l'article 44 du règlement d'application (UE) 2020/,,, [R*eference to present Regulation*]/ Part IV to be completed in accordance with Article 44 of Implementing Regulation (EU) 2020/… [*Reference to this Regulation*]/ *official language*
8	L'impression de cette référence n'est obligatoire que pour les duplicata de document d'identification délivrés conformément à l'article 38, paragraphe (2)(b), du règlement (UE) 2021/ [*reference to present Regulation*]./The print of this reference is only mandatory for duplicate identification documents issued in accordance with Article 38(2)(b). of Regulation (EU) ….. [*reference to present Regulation*]./ *official language*

SECTION III

MARQUE DE VALIDATION OU LICENCE/ VALIDATION MARK OR LICENCE/*official language*

Code Unique /Unique Code/*official language*: □□□-□□□-□□□□□□□□□		
Conformément à l'article 92, paragraphe (2), du règlement délégué (UE) 2021/688/ In accordance with Article 92(2) of Delegated Regulation (EU) 2020/688/ *official language* Marque de validation valable jusque à /Validation mark valid until/*offical language*: …….. or Licence valable jusque à / Licence valid until/ *official language*: …….	Autorité compétente ou organisme émetteur/ Competent authority or issuing body/ *official language*	Date/Date/ *official language* Lieu/Place/ *official language* Nom (en lettres capitales) et signature de la personne qualifiée /Name (in capital letters) and signature of qualified person / *official language* Stamp of competent authority or issuing body/ *official language*
Note: The layout of this Section may vary as well as its order in the single lifetime identification document, provided the order and numbering of Sections I and II are observed. The licence or validation mark may be placed behind a transparent window in the cover for fast checks.		

SECTION IV

Certificat zootechnique pour les échanges de reproducteurs de race pure de l'espèce équine (*Equus caballus* et *Equus asinus*), conformément à l'annexe V, partie 2, chapitre I, du règlement (UE) 2016/1012
Zootechnical certificate for trade in purebred breeding animals of the equine species (*Equus caballus* and *Equus asinus*), in accordance with Chapter I of Part 2 of Annex V to Regulation (EU) 2016/1012 /
official language

<table>
<tr><td colspan="4">PART I</td></tr>
<tr><td colspan="4">1. Name of issuing breed society or competent authority
(provide contact details and, where available, a reference to the website)</td></tr>
<tr><td colspan="2">2. Name of breeding book</td><td colspan="2">3. Name of breed</td></tr>
<tr><td colspan="2">4. Name and commercial name of animal(1) and code of country of birth(2)</td><td colspan="2">5.1. Individual identification number(3)</td></tr>
<tr><td colspan="2">6. Breeding book number(5)</td><td colspan="2">5.2. Unique Life Number(4) ☐☐☐-☐☐☐-☐☐☐ ☐☐☐ ☐☐☐</td></tr>
<tr><td colspan="4">7. Identification of animal(1)(6)
7.1. Transponder code(1) ☐☐☐ ☐☐☐ ☐☐☐ ☐☐☐ ☐☐☐
Reading system (if not ISO 11784)(1)
Bar-Code(1)
7.2. Alternative method for identity verification(1)</td></tr>
<tr><td colspan="2">8. Date of birth of animal
(use format dd/mm/yyyy)</td><td colspan="2">9. Country of birth of animal</td></tr>
<tr><td colspan="4">10. Name, address and e-mail address(1) of breeder</td></tr>
<tr><td colspan="4">11. Pedigree(7)(8)</td></tr>
<tr><td>11.1. Sire</td><td>11.1.1. Paternal Grandsire
Breeding book number and section</td><td>11.1.1.1.(1) Paternal Grand-Grandsire
Breeding book number and section</td><td></td></tr>
</table>

Breeding book number and section		11.1.1.2.[1] Paternal Grand-Granddam Breeding book number and section	
	11.1.2. Paternal Granddam Breeding book number and section	11.1.2.1.[1] Paternal Grand-Grandsire Breeding book number and section	
		11.1.2.2.[1] Paternal Grand-Granddam Breeding book number and section	
11.2. Dam Breeding book number and section	11.2.1. Maternal Grandsire Breeding book number and section	11.2.1.1.[1] Maternal Grand-Grandsire Breeding book number and section	
		11.2.1.2.[1] Maternal Grand-Granddam Breeding book number and section	
	11.2.2. Maternal Granddam Breeding book number and section	11.2.2.1.[1] Maternal Grand-Grandsire Breeding book number and section	
		11.2.2.2.[1] Maternal Grand-Granddam Breeding book number and section	

12.1. Done at *(insert place of issue)* 12.2. Done on *(insert date of issue in format dd/mm/yyyy)* 12.3. Name and capacity of the signatory *(insert in capital letters name and capacity of the individual[9] authorised by the issuing breed society or competent authority to sign this part of the zootechnical certificate)*	12.4. Signature

(1) Keep empty if not applicable.

(2) Enter country code where required by international agreements on the breed.

(3) The individual identification number in accordance with point 3 of Chapter I of Part 1 of Annex II to Regulation (EU) 2016/1012 of the European Parliament and of the Council, referred to as 'unique code' in Article 114(1)(a) of Regulation (EU) 2016/429 of the European Parliament and of the Council, and recorded in accordance with Article 8(2) of Implementing Regulation (EU) 2021/…. [*Reference to current Regulation*].

(4) Unique life number as defined in point (o) of Article 2 of Implementing Regulation (EU) 2015/262, if assigned in accordance with that Implementing Regulation

(5) Required if different from the individual identification number or unique life number assigned in accordance with Implementing Regulation (EU) 2015/262.

(6) Not required if Part I of the zootechnical certificate is an integral part of the single lifetime identification document issued by a breed society. If the single lifetime identification document was issued in accordance with Implementing Regulation (EU) 2015/262, the unique life number as defined in point (o) of Article 2 of that Implementing Regulation shall be stated.

(7) If necessary, include additional generations.

(8) Enter the individual identification number in accordance with point 3 of Chapter I of Part 1 of Annex II to Regulation (EU) 2016/1012, referred to as 'unique code' in Article 114(1)(a) of Regulation (EU) 2016/429. If the individual identification number is either not available or different from the number under which the animal is entered in the breeding book, enter the breeding book number.

(9) That individual shall be representative of the breed society or competent authority referred to in Article 30(2)(b) of Regulation (EU) 2016/1012.

<table>
<tr><td colspan="2">PART II</td></tr>
<tr><td>1.1. Individual identification number[1]

1.2. Unique Life Number[2] □□□-□□□-□□□□□□□□□</td><td rowspan="2">2. Identification of animal[3]
2.1. Transponder code[4] □□□ □□□ □□□ □□□ □□□
Reading system (if not ISO 11784)[4]
Bar-Code[4]

2.2. Alternative method for identity verification[3]</td></tr>
<tr><td>3. Sex</td></tr>
<tr><td>4. Class within the main section of the breeding book[4]
4.1. Name of breeding book[5] 4.2. Class within the main section[6]</td><td>5. Name, address and e-mail address[4] of owner[7]</td></tr>
<tr><td colspan="2">6. Additional information[4][8][9]
6.1. Results of performance testing

6.2. Up-to-date results of the genetic evaluation carried out last on
(insert date in format dd/mm/yyyy)

6.3. Genetic defects and genetic peculiarities of animal in relation to the breeding programme

6.4. System of identity verification and result[4][10][11] 6.5. Results of parentage control[4][10][12]</td></tr>
</table>

7. Insemination/mating[13][14]

7.1. Date *(use format dd/mm/yyyy)*

7.2. No of covering certificate[15]

7.3. Identification of the donor male

7.3.1. Individual identification number[1]

7.3.2. Unique Life Number[2] ☐☐☐-☐☐☐-☐☐☐☐☐☐☐☐☐

7.3.3. System of identity verification and result[4][10][11]

7.3.4. Results of parentage control[4]

8.1. Done at *(insert place of issue)*	8.2. Done on *(insert date of issue in format dd/mm/yyyy)*
8.3. Name and capacity of the signatory *(insert in capital letters name and capacity of the individual[16] authorised by the issuing breed society or competent authority to sign this part of the certificate)*	8.4. Signature

(1) The individual identification number in accordance with point 3 of Chapter I of Part 1 of Annex II to Regulation (EU) 2016/1012, referred to as 'unique code' in Article 114(1)(a) of Regulation (EU) 2016/429 of the European Parliament and of the Council, and recorded in accordance with Article 8(2) of Implementing Regulation (EU) 2021/…. [*Reference to current Regulation*]..

(2) Unique life number as defined point (o) of Article 2 of Implementing Regulation (EU) 2015/262, if assigned in accordance with that Implementing Regulation.

(3) Not required if information corresponds to information in point 7 of Part I and Parts I and II are an integrated whole and indivisible and contained in or attached to the single lifetime identification document. If the single lifetime identification document was issued in accordance with Implementing Regulation (EU) 2015/262, the unique life number as defined in point (o) of Article 2 of that Regulation shall be stated.

(4) Keep empty if not applicable.

(5) Required if different from point 2 of Part I.

(6) Not required where this information is provided in Section V of the identification document issued in accordance with Commission Implementing Regulation (EU) 2015/262.

(7) Not required if information on the owner is available and up-to-date in other parts of the single lifetime identification document.

(8) If necessary use additional paper.

(9) If that genetic information can be accessed on a website, a reference to that website may be provided instead, if authorised by the competent authority in accordance with Article 32(3) of Regulation (EU) 2016/1012.

(10) Based on DNA analysis or analysis of its blood group.

(11) Required in accordance with Article 22(1) of Regulation (EU) 2016/1012 for purebred breeding animals of the equine species used for the collection of semen for artificial insemination. It may be required by breed societies in accordance with Article 22(2) of Regulation (EU) 2016/1012 for purebred breeding animals of the equine species used for the collection of oocytes and embryos. Indicate details or reference to laboratory report where such details can be accessed by the issuing breed society.

(12) If required by the breeding programme.

(13) Required in the case of pregnant females. Information may be indicated in a separate document.

(14) Delete as appropriate.

(15) If not applicable, provide results of parentage control in point 7.3.4.

(16) That individual shall be a representative of the breed society or competent authority referred to in Article 30(2)(b) of Regulation (EU) 2016/1012.

Note for the issuing authority [*not to be printed in identification document*]: Layout variations from this model are permitted, provided that the required minimum information is ensured. Footnotes may not be printied provided a reference is made to the accessible explanation

SECTION V

Coordonnées du propriétaire

1. Pour les compétitions sous compétence de la Fédération équestre internationale (FEI), la nationalité du cheval doit être celle de son propriétaire.
2. En cas de changement de propriétaire, le document d'identification doit être immédiatement déposé auprès de 1'organisation, 1'association ou le service officiel 1'ayant délivré avec le nom et l'adresse du nouveau propriétaire afin de le lui transmettre après ré-enregistrement.
3. S'il y a plus d'un propriétaire ou si le cheval appartient ä une société, le nom de la personne responsable du cheval doit être inscrit dans le document d'identification ainsi que sa nationalité. Si les propriétaires sont de nationalités différentes, ils doivent préciser la nationalité du cheval.
4. Lorsque la FEI approuve la location d'un cheval par une Fédération équestre nationale, les détails de ces transactions doivent être enregistrés par la Fédération équestre nationale intéressée.

Details of ownership

1. For competition purposes under the auspices of the, International Federation for Equestrian Sports (FEI) the nationality of the horse shall be that of its owner.
2. On change of ownership the identification document must immediately be lodged with the issuing body, organisation, association or official service, giving the name and address of the new owner, for re-registration and forwarding to the new owner.
3. If there is more than one owner or the horse is owned by a company, then the name of the individual responsible for the horse must be entered in the identification document together with his nationality. If the owners are of different nationalities, they have to determine the nationality of the horse.
4. When the FEI approves the leasing of a horse by a national equestrian federation, the details of these transactions must be recorded by the national equestrian federation concerned.

official language

Date d'enregistrement par l'organisation, 1'association ou le service officiel **Date of registration by the organisation, association, or official service** / *official language*	**Nom du propriétaire** **Name of owner** / *official language*	**Adresse du propriétaire** **Address of owner** / *official language*	**Nationalité du propriétaire** **Nationality of owner** / *official language*	**Signature du propriétaire** **Signature of owner** / *official language*	**Cachet de 1'organisation, association ou service officiel et signature** **Organisation, association or official service stamp and signature**/ *official language*

SECTION VI

Contrôles d'identité de l'équidé décrit dans ce document d'identification

L'identité de l'équidé doit être contrôlée chaque fois que les lois et règlements 1'exigent et il doit être certifié qu'elle est conforme à la description donnée dans la section I du document d'identification.

Control of identification of the equine animal described in the identification document

The identity of the equine animal must be checked each time this is required by the rules and regulations and certified that it conforms to the description given in Section I of the identification document.

/ *official language*

Date (jj/mm/aaaa) **Date** (dd/mm/yyyy) */ official language*	**Ville et pays** **Place and country** / *official language*	**Motif du contrôle (concours, certificat sanitaire, etc.)** **Purposeof of check (event, health certificate, etc.)** / *official language*	**Nom (en lettres capitales), qualité de la personne ayant vérifié 1'identité et signature** **Name (in capital letters), capacity of official verifying the identity and signature,** / *official language*

SECTION VII

Grippe équine seulement
ou
Grippe équine utilisant des vaccins combinés

Enregistrement des vaccinations

Toute vaccination subie par l'équidé doit être portée dans le cadre ci-dessous de façon lisible et précise et complétée par le nom et la signature du vétérinaire.

Equine influenza only
or
equine influenza using combined vaccines

Vaccination record

Details of every vaccination which the equine animal has undergone must be entered clearly and in detail, and completed with the name and signature of veterinarian.

/ official language

Date **Date** */ official language*	**Lieu** **Place** */ official language*	**Pays** **Country** */ official language*	**Vaccin/Vaccine**			**Nom (en lettres capitales) et signature du vétérinaire** **Name (in capital letters) and signature of veterinarian** */ official language*
			Nom **Name** */ official language*	**Numéro du lot** **Batch number** */ official language*	**Maladie(s)** **Disease(s)** */ official language*	

SECTION VIII

Maladies autres que la grippe équine **Diseases other than equine influenza** */ official language*

Enregistrement des vaccinations *Vaccination record*

Toute vaccination subie par l'équidé doit être portée dans le cadre ci-dessous de façon lisible et précise et complétée par le nom et la signature du vétérinaire.

Details of every vaccination which the equine animal has undergone must be entered clearly and in detail, and completed with the name and signature of veterinarian.

Date **Date** */ official language*	**Lieu** **Place** */ official language*	**Pays** **Country** */ official language*	**Vaccin/Vaccine**			**Nom (en lettre capitales) et signature du vétérinaire** **Name (in capital letters) and signature of veterinarian** */ official language*
			Nom **Name** */ official language*	**Numéro du lot** **Batch number** */ official language*	**Maladie(s)** **Disease(s)** */ official language*	

SECTION IX

Examen de laboratoire

Le résultat de tout examen effectué par un vétérinaire pour une maladie transmissible ou par un laboratoire agréé par le service vétérinaire officiel du pays (‘laboratoire officiel’) doit être noté clairement et en détail par le vétérinaire qui représente l'autorité demandant l’examen.

Laboratory health test

The result of every test carried out for a transmissible disease by a veterinarian or by a laboratory authorised by the official veterinary service of the country (‘official laboratory’) must be entered clearly and in detail by the veterinarian acting on behalf of the authority requesting the test.

/ official language

Date de prélèvement Sampling date */ official language*	**Maladie transmissible concernée Transmissible disease tested for** */ official language*	**Nature de l’examen Type of test** */ official language*	**Résultat de l’examen Result of test** */ official language*	**Laboratoire officiel ayant effectué l’examen Official laboratory which carried out the test** */ official language*	**Nom (en lettres capitales) et signature du vétérinaire Name (in capital letters) and signature of veterinarian** */ official language*

SECTION X

Châtaignes

Dessiner le contour de chaque châtaigne dans la carré correspondant: à ne remplir que pour les chevaux sans marque et avec moins de trois épis

Chestnuts

The outline of each of the four chestnut must be drawn in the appropriate square for all horses without markings and with less than three whorls.

/ *official language*

Antérieur droit/Right Foreleg/ *official language*	**Postérieur droit/Right Hindleg/** *official language***.**
Antérieur gauche/Left Foreleg/ *official language*	**Postérieur gauche/Left Hindleg/** *official language*

Part 2
Additional requirements for the single lifetime identification document for equidae

The single lifetime identification document shall:

(a) be in the format of a printed passport with a paper size between 210 x 148 mm (A5) and 250 x 200 mm;

(b) have a distinct cover (front and back) that provides sufficient protection, which may be embossed with the logo of the issuing body, and may have a pocket at the inside back cover for the insertion of pages containing Sections IV to X, as appropriate;

(c) have at least Sections I, II and III indivisibly machine-rivetted to prevent pages being fraudulently removed or replaced;

(d) where serial numbers are applied, have at least Sections I, II and III printed on pages bearing the serial number of the single lifetime identification document;

(e) have at least each page of Sections I, II and III numbered in the format "page number/total number of pages";

(f) have Part A of Section I sealed with a transparent adhesive laminate after the required information has been entered, unless Section I of the identification document is printed by the issuing body after the required information was entered in a way that prevents alterations;

(g) have the General Instructions provided for in Part 1 printed in the document if it contains Sections I to X. In the case of a single lifetime identification document comprised of only Sections I to III, the printing of the General Instructions provided for in Part 1 is optional.

ANNEX III

Part 1
Information stored on plastic cards or smart cards

The plastic card or smart card shall contain at least the following:

1. Visible information on the plastic card or smart card:
   - competent authority;
   - unique code;
   - species and sex;
   - the last 15 digits of the code transmitted by the transponder;
   - a photograph of the equine animal (optional).
2. Electronic information on the smart card accessible by use of standard software:
   - all compulsory information in Sections I to X of the single lifetime identification document;
   - logging of any modification of previously entered information;
   - a photograph of the equine animal (optional).

Part 2
Physical characteristics of the plastic cards and smart cards

The plastic cards and smart cards shall have the following physical characteristics:

- in accordance with ISO standard 7810 and ISO standard 7816-1;
- the material used shall be made secure against forgery;
- information contained in the front and reverse side of the card shall be legible with the eye, using a minimum character size of 5 points.

ANNEX IV

Model of temporary identification document referred to in Article 24

<table>
<tr><td rowspan="2">Competent authority</td><td>TEMPORARY DOCUMENT
(Article 24 of Commission Implementing Regulation (EU) 2021/..</td><td>Name of Country</td></tr>
<tr><td>Name and Address of keeper/owner:
…</td><td>Unique code
□□□-□□□-□□□□□□□□□□

Barcode of Unique code (where available)</td></tr>
<tr><td colspan="2">Name of animal:</td><td rowspan="4">Transponder code/ear tag

□□□ □□□ □□□ □□□ □□□
Bar-Code (optional)/ear tag</td></tr>
<tr><td colspan="2">Sex:</td></tr>
<tr><td colspan="2">Colour:</td></tr>
<tr><td colspan="2">Date of birth:</td></tr>
<tr><td colspan="3">Alternative method for identity verification (if available):</td></tr>
<tr><td>Date and place of issuing:</td><td>Name (in capital letters) and capacity of signatory</td><td>Signature</td></tr>
</table>

Note for the competent authority or delegated body [*not to be printed in identification document*]: Slight variations from this model are permitted.