



The official passport is in Bulgarian language!

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**SECTION I/Section I**

**Part A – Partie A – Details d'identification/Part A – Identification details**

(D(a))	BMFL:/Especie:/Species:	(4)	<p>Unique code or lifetime number: (15 characters):/Code Unique ou Numero unique d'identification VALABLE a vie (15 chiffres):/Unique Code or Lifer number: (15 digits):</p> <p>Barcode (no M36op)/Code-barres (optionnel)/Bar-Code (optional)</p>
(1)(6)	Gender: /Sex:/Sex:	(5)	<p>Transponder code (if available)/Code du transpondeur (si disponible)/Transponder code (where available)</p> <p>Reading system (if not ISO 11784)/Systeme de lecture (si different de ISO 11784)/Reading system (if not ISO 11784) Barcode (no M36op)/Code-barres (optionnel)/Bar-Code (optional)</p>
(2)(a)	Date of birth:/0a1f de naissance:/Date of birth:		
(2)(6)	Place and country of birth: /Lieu et pays de naissance:/Place and country of birth:		
(2)(B)	Name (no Ma6op):/Nom (optionnel):/Name (optional):		

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### (3) Described:/Signalement:/Description:

(3)(a)	Color:/Who:/So1oyg:
(3)(b)	Chapter:/Tete:/Head:
(3)(c)	Front leg, left:/App1. G:/Foreleg L:
(3)(d)	Front leg, right:/App1. D:/Foreleg R:
(3)(e)	Rear leg, left:/Rov1 G:/Hind leg L:
(3)(f)	Rear leg, fleceH:/Post D:/Hind leg R:
(3)(g)	Body:/Corps:/Body:
(3)(h)	Tags:/Marques:/Markings:

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(6)	Alternative method of authentication (if applicable)/Methode Alternative deverification d'identite (si applicable)/Alternative method for identity verification (if applicable)
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(7)	Information on any other appropriate method enabling the identification of the animal (blood group/DNA code) (optional);/Information surtout autre methode appropriee don- Nant des garanties pour verifier l'identite 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)
(8)	Date/Date/Date:
(9)	Location/Lieu/Place:

(10) Signature of the qualified person (name 6yKBn)/Signature de la personne qualifiee (nom en lettres capitales)/Signature of qualified person (name in capital letters):

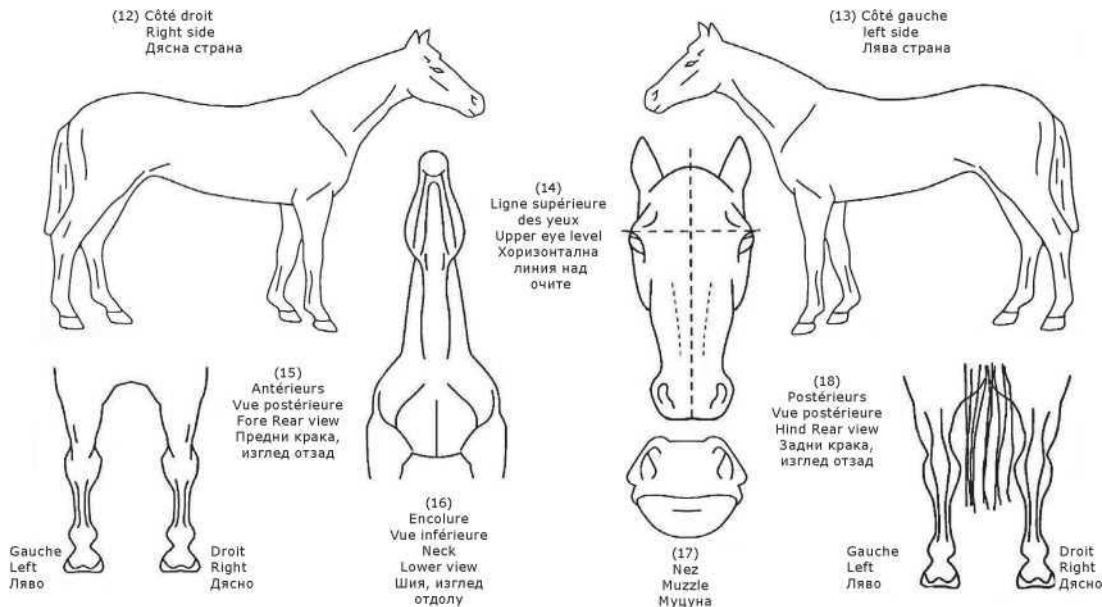
Stamp of the competent authority or delegated body/Cachet de l'Autorite competente ou de l'organisme delegue/stamp of competent authority or delegated body

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## Part B – Contour Scheme/Partie C – Signalement graphique/Part C – Outline Diagram

C  
W  
CH

### TRANSPONDER



Signature de la personne qualifiée (nom en lettres capitales) et chachet de l'autorité competente ou de l'organisme delegue

Signature of the qualified person (name in capital letters) and stamp of the competent authority or delegated body

Signature of the qualified person (name in capital letters) and stamp of the competent authority or delegated body

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Part C – Castration, verification de la description, autrestion/Part C – Castration, verification of the description, other modifications		
Castration/Castration/ Castration/	Identification/Identification/Identification/	
Date and place of castration/Date et lieu de la castration/Date and place of castration       Signature and stamp of veterinarian/Signature et cachet du veterinaire/Signature and stamp of veterinarian	Verification de la description/Verification of the deScription Includes/Mentionner/Include:	Signature of the qualified person (name in capital letters)/Signature de la personne qualifiee (nom en lettres capitales)/Signature of qualified person (name in capital letters):       Printed competent authority or delegated body/Cachet de de l'Autorite competente ou l'organisme delegue/Stamp of COMPETent authority or delegated body
	1. Amendments/rectifications/Amendments	
	2. Additions/Adjonctions/Additions	
	3. Registration of an identification document in the database of a competent authority or delegated authority other than that which issued the original document/Enregistrement d'un document d'identification d'identification dans la base de donnees d'un Autorite competente ou d'un organ- ISME delegue autre que celui qui a delivre le document original/Registration of an identification document in the database of a competent authority or delegated body than that issued the original document	Date and place/Date et lieu/Date and place

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## Administration of medicinal products/Administration of medicinal products

Code Unique/Unique Code:

### Part I/Partie I

Date and place of issue of this Section<sup>1/Date</sup> et lieu delivrance de la presente section<sup>1/Date</sup> and place of issue of this Section<sup>1</sup>: .....

Competent authority or delegated body for this section of identification<sup>document 1/Autorite competente</sup>  
ou organisme delegue de la presente section du document d<sup>identification 1/Competent authority or delegated body for this Section of the</sup>  
identification document<sup>1</sup>: .....

### Part II/Partie II/Part II

**Note/The single ungulate is not intended for slaughter for human consumption and may be Remarkue/ administered veterinary medicinal products authorised in accordance with Article 8(4) of**

**Note: Regulation (EU) 2019/6 or medicinal products administered in accordance with Article 112(4) of that Regulation./**

**L'equide n'est pas DestinE a l'abattage pour la consommation humaine, et par consequent, l'equide peut recevoir des medicaments veterinaires autorises conformement a l'article 8, paragraphe 4, du reglement (UE) 2019/6 ou des medicaments administres conformement a l'article 112, paragraphe 4, du ledit reglement./**

**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 medicinal products administered in accordance with Article 112(4) of that Regulation.**



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<b>Declaration/Declaration/Declaration</b>	<b>The equine animal described in this identification document is not intended for slaughter for human consumption/L'animal equine decrit dans le present document d'identification n'est pas DestinE a l'abattage pour la consommation humaine./The equine animal described in this identification document is not intended for slaughter for human consumption</b>		
Date and place/Date et lieu/Date and place:	Responsible veterinarian acting in accordance with Article 112(4) of Regulation (EU) 2019/62/Veterinaire responsable procedant conformement a l'article 112, paragraphe 4, du reglement (UE) 2019/6/Veterinarian responsible acting in accordance with Article 112(4) of Regulation (EU) 2019/62:	Responsible veterinarian/veterinaire responsable/Veterinarian responsible	
	Competent authority <sup>2</sup> or delegated body <sup>2/Autorite competente<sup>2</sup> ou organisme delegue<sup>2/Competent authority<sup>2</sup> or delegated body<sup>2</sup> </sup></sup>	Name (in capital letters) and signature of the responsible person <sup>2</sup> /Name (in capital letters) and signature of the person responsible <sup>2</sup>	
		Name/Nom/Name <sup>5</sup> :	Signature/Signature/Signature
		Address/Adresse/Address <sup>5</sup> :	
		Postcode/Code postal/Postal code <sup>5</sup> :	
		Location/Lieu/Place <sup>5</sup> :	
		Phone/Telephone/Telephone <sup>6</sup> :	

**Part III/Partie III/Part III**

**Note/ Remarque/ Note:** **The equine animal is intended for slaughter for human consumption./L'équide est DestinE a l'abatt— age pour la consommation humaine./The equine animal is intended for slaughter for human consumption.**

Without prejudice to Regulation (EC) No 470/2009 and Directive 96/22/EC, the equine animal may be subjected to medical treatment in accordance with Article 115(1) of Regulation (EU) 2019/6, provided that the treated equine animal may be slaughtered for human consumption only after the total withdrawal period of 6 months has elapsed from the date of the last application of the substances listed in accordance with Article 115(5) of that Regulation./ Sans prejudice du reglement (CE) No 470/2009 ni de la Directive 96/22/CE, l'équide peut faire l'objet d'un traitement medicamenteux conformement a l'article 115, paragraphe 1, du reglement (UE) 2019/6 a condition que l'équide losi traite ne soit abattu en vue de la consommation humaine qu'au terme d'un temps d'attente general de sixis suivant la date de la la derniere administration de substances conformement a l'article 115, paragraphe 5, du ledit reglement./Without prejudice to Regulation (EC) No 470/2009 and Directive 96/22/EC, the equine animal may be subject to medic Inaltreatment 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.

DOCUMENTATION HA TREATMENT/ENREGISTREMENT DE LA MEDICATION/MEDICATION RECORD		
<p>Date and place of the last prescribed application in accordance with Article 115(1) of Regulation (EU) 2019/6<sup>(2)</sup>/Date et lieu de la derniere administration, telle que prescrite, conformement a l'article 115, paragraphe 1, du reglement (UE) 2019/6<sup>(2)</sup>/Date and place of last administration, as prescribed, in accordance with Article 115<sup>(1)</sup> of Regulation (EU) 2019/6<sup>2</sup>&gt;</p>	<p>Essential substance(s) included in the medicinal product administered in accordance with Article 115 of Regulation (EU) 2019/6 as specified in the first column<sup>(2) &amp;lt;t,h) (-&gt;)/Substance(s) essentielle(s) incorporee(s) dans le medicament administre conformement a l'article 115, du reglement (UE) 2019/6<sup>2</sup>&gt;, comme mentionne dans la premiere colonne<sup>(2)(3)(4)/Essential substance(s) incorporated in the medicinal product administered in accordance with Article 115 of Regulation (EU) 2019/6 as mentioned in the first column<sup>(2)(3)(4)</sup></sup></sup></p>	<p>Responsible veterinarian performing and/or administering a medicinal product/veterinaire responsable administrant et/ou prescrivait l'administration d'un medicament/Veterinarian responsible administering and/or prescribing the administration of the medicinal product</p>

		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature
		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature
		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature

		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature
		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature
		Name/Nom/Name <sup>5</sup> : ----- Address/Adresse/Address <sup>5</sup> : ----- Postcode/Code postal/Postal code <sup>5</sup> : ----- Location/Lieu/Place <sup>5</sup> : ----- Phone/Telephone/Telephone <sup>6</sup> : ----- -----	Signature/Signature ----- Signature

Part IV/Partie IV/Part IV &<sup>7</sup>>

<p><b>Note/ Remarque/ Note:</b></p> <p><b>Trade in registered equidae which have been administered veterinary medicinal products containing allyl trenbolone or beta-agonists 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./Les échanges des équides 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/EC/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.</b></p>			
<p>Date of last Annex in accordance with Article 4 of Directive 96/22/EC/ 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/</p>	<p>Substance(s) included in the veterinary medicinal product administered in accordance with Article 4 of Directive 96/22/EC/Sub-position(s) incorporée(s) dans le médicament vétérinaire administré conformément à l'article 4 Directive 96/22/EC/Substance(s) incorporated in the veterinary medicinal product administered in accordance with Article 4 of Directive 96/22/EC/</p>	<p>Responsible veterinarian performing and/or prescribing the administration of a veterinary medicinal product/ Vétérinaire responsable administrant et/ou prescrivant l'administration d'un médicament vétérinaire/ Veterinarian responsible administering and/or prescribing administration of veterinary medicinal product</p>	
		<p>Name/Nom/Name<sup>5</sup>:</p> <p>Address/Adresse/Address<sup>5</sup>:</p> <p>Postcode/Code postal/Postal code<sup>5</sup>:</p> <p>Location/Lieu/Place<sup>5</sup>:</p> <p>Phone/Telephone/Telephone<sup>6</sup>:</p>	<p>Signature/Signature/Signature</p>

Part IV/Partie IV/Part V & 7>

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**Note/Remarque/Note:** **The equine animal is intended for slaughter for human consumption./L'équide est DestinE a l'abattage pour la consommation humaine.**

The slaughter of the equine animal has been postponed for administrative reasons by at least six months in accordance with Article 38(2)(b) of Implementing Regulation (EU) 2021/963/L'abattage de l'équide est pour des raisons administratives retardé d'au moins six mois conformément à l'article 38, paragraphe 2(b) du règlement d'exécution (UE) 2021/963/  
 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) 2021/963/

Date of suspension/Date de la suspension/Date of suspension	Location/Lieu/Place	Competent authority <sup>2</sup> or delegated body 2/Autorité compétente <sup>2</sup> ou organisme délégué 2/Competent authority <sup>2</sup> or delegated body <sup>2</sup>	Name (in capital letters) and signature of the responsible person/Nom (en lettres capitales) et signature de la personne responsable/Name (in capital letters) and signature of the person responsible

1. Information is required only if this section is issued on a date other than that of Section I./Information a ne fournir que si la presente section est delivree a une autre date que la section I./Information only required if this Section is issued at a different date than Section I.

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

3. Substances must be indicated according to the list of substances established in accordance with Article 115(5) of Regulation (EU) 2019/6./Il est indispensable de specifier les substances en se fondant sur la liste de substances etablie conformement a l'article 115, paragraphe 5, du reglement (UE) 2019/6./Specification of substances against list of substances established in accordance with Article 115(5) of Regulation (EU) is compulsory.

4. Information on other veterinary medicinal products administered in accordance with Regulation (EU) 2019/6 is optional./Les informations relatives a d'autres medicaments veterinaires administres conformement au reglement (UE) 2019/6 sont facultatives.

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

6. Telephone in format [+ country code (location code) number]./Numero de telephone selon le modele [+ code pays (code regional) numero]./Telephone in format [+ country code (regional code) number].

7. Part IV doit etre completee conformement a l'article 44 du reglement d'application (UE) 2021/963//Part IV to be completed in accordance with Article 44 of Implementing Regulation (EU) 2021/963/

8. The printing of this reference is mandatory only for duplicate identification documents issued in accordance with Article 38(2)(b) of Regulation (EU) 2021/963./L'impression de cette reference n'est obligatoire que pour les duplicata de document d'identification delivres conformement a l'article 38, paragraphe (2)(b), du reglement (UE) 2021/963./The print of this reference is only mandatory for duplicate identification documents issued in accordance with Article 38(2)(b) of Regulation (EU) 2021/963.

**SECTION III/SECTION III**

Validating Mark or Licence/Marque de validation ou licence/Validation Mark or Licence/

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<b>Code Unique/Unique Code:</b>		
<p>In accordance with Article 92(2) of Delegated Regulation (EU) 2020/688./Conformement a Particle 92, paragraphe (2), du reglement delegue (UE) 2021/688/In accordance with Article 92(2) of Delegated Regulation (EU) 2020/688/</p> <p>Valid until/Marque de validation valablejusque a/Validation mark valid until:</p>	<p>Competent authority or delegated body/Auto rite competente ou organisme delegue/Competent authority or delegated body/</p>	<p>Date/Date/Date</p> <p>Location/Lieu/Place/</p> <p>Name (in capital letters) and signature of qualified person</p>
<p>or</p> <p>License valid until/Licence VALABLE jusque a/Licence valid until</p>		<p>Stamp of competent authority or delegated body/Cachet de l'au- torite competente ou de l'organisme delegue/Stamp of competent authority or delegated body/</p>



**SECTION III/SECTION IV**

		<b>Code Unique/Unique Code:</b>
<b>Coordonnees du proprietaire</b>	<b>Details of ownership</b>	<b>Ownership data</b>
1. Pour les competitions sous competence de la Federation equestre internationale (FEI), la nationalite du cheval doit etre celle de son proprietaire.	1. For competition purposes under the auspices of the, International Federation for Equestrian Sports (FEI) the nationality of the horse should be that of its owner.	1. For competitions organised by the Federation Equestre Internationale (FEI), the nationality of the horse is that of its owner.
2. En cas de changement de proprietaire, le document d'identification doit etre immediatement depose aupres de l'organisation, l'association ou le service officiel l'ayant delivre avec le nom et l'adresse du nouveau proprietaire afin de le lui transmettre apres re-enregistrement.	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.	2. In the event of a change of owner, the identification document must be transmitted without delay to the issuing authority, organisation, association or official service, with the name and address of the new owner, for re-registration and transmission to the new owner.
3. S'il y a plus d'un proprietaire ou si le cheval appartient a une societe, le nom de la personne responsable du cheval doit etre inscrit dans le document d'identification que sa nationalite. Si les proprietaires sont de nationalites differentes, ils doivent preciser la nationalite du cheval.	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 into the identification document together with his nationality. If the owners are of different nationalities, they have to determine the nationality of the horse.	3. If there is more than one owner or the horse is owned by a company, the name and nationality of the person responsible for the horse shall be entered in the identification document. If the owners are of different nationalities, they are obliged to determine the nationality of the horse.
4. Lorsque la FEI approuve la location d'un cheval par une Federation equestre nationale, les details de ces transactions doivent etre enregistres par la Federation equestre nationale interessee.	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.	4. When the FEI approves the rental of a horse by a national equestrian federation, the information on these operations must be recorded by the relevant national equestrian federation.

<b>Date of registration by the organisation, association or official service/Date d'enregistrement par l'organisation, l'association ou le service officiel/Date of registration by the organisation, association, or official service</b>	<b>Name of owner/Nom du propriétaire/Name of owner</b>	<b>Address of owner/Adresse du propriétaire/Address of owner</b>	<b>Nationality of owner/Nationalité du propriétaire/Nationality of owner</b>	<b>Signature of owner/Signature du propriétaire/Signature of owner</b>	<b>Stamp and signature of the organisation, association or official service stamp, association or official service signature/Organisation, association or official service stamp and signature</b>

<b>Date of registration by the organisation, association or official service/Date d'enregistrement par l'organisation, l'association ou le service officiel/Date of registration by the organisation, association, or official service</b>	<b>Name of owner/Nom du propriétaire/Name of owner</b>	<b>Address of owner/Adresse du propriétaire/Address of owner</b>	<b>Nationality of owner/Nationalité du propriétaire/Nationality of owner</b>	<b>Signature of owner/Signature du propriétaire/Signature of owner</b>	<b>Stamp and signature of the organisation, association or official service stamp, association ou service officiel et signature/Organisation, association or official service stamp and signature</b>

### SECTION V/SECTION V

Zootechnical certificate for trade in purebred breeding animals of the equine species (*Equus caballus* and *Equus asinus*) in accordance with by Chapter I of Part 2 of Annex V to Regulation (EU) 2016/1012

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

PART I	
1. Name of the issuing breed society or competent authority (insert contact information and, if available, a reference to it)	
2. Name of the breeding book	3. Breed name
4. Name and trade name of the animal <sup>(1)</sup> and country of birth code <sup>(2)</sup>	5.1. Individual identification number <sup>(3)</sup> 5.2. Unique Life Number <sup>(4)</sup>
6. Herd-book number <sup>(5)</sup>	
7. Animal identification <sup>(1)(6)</sup> 7.1. Transponder's code for the transponder Reading system (if not ISO 11784) <sup>(1)</sup> Barcode – Barcode 7.2. Alternative Identity Check Method	
8. Date of birth of the animal  <i>(in format dd/mm/yyyy)</i>	9. Country of birth of the animal
10. Name, address and e-mail of the breeder	

11. Pedigree <sup>(7)(8)</sup>				
11.1. Father Breeding book number and section	11.1.1. Father's Father Breeding book number and section	11.1.1.1. <sup>(1)</sup> Father of father Breeding book number and section		
		11.1.1.2. <sup>(1)</sup> Father's father's mother Breeding book number and section		
	11.2.1. Father's Mother Breeding book number and section	11.1.2.1. <sup>(1)</sup> Father of the father's mother Breeding book number and section		
		11.1.2.2. <sup>(1)</sup> Mother of father's mother Breeding book number and section		
11.2. Mother Breeding book number and section	11.2.1. Mother's Father Breeding book number and section	11.2.1.1. <sup>(1)</sup> Father of mother's father Breeding book number and section		
		11.2.1.2. <sup>(1)</sup> Mother of mother's father Breeding book number and section		
	11.2.2. Mother of mother Breeding book number and section	11.2.2.1. <sup>(1)</sup> Father of mother of mother Breeding book number and section		
		11.2.2.2. <sup>(1)</sup> Mother of mother's mother Number by herd book and section		
12.1. Done at		12.2. Done on		
<i>(place of issue)</i>		<i>(date of issue in dd/mm/yyyy format)</i>		
12.3. Name and function of the signatory			12.4. Signature	

(name and function of the <sup>person 89)</sup> to whom the issuing breed society or competent authority has conferred powers to sign this part of the zootechnical certificate, enter capital letters)

(1) If not applicable, not completed.

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

(3) The individual identification number in accordance with Chapter 3 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 listed in accordance with Article 8(2) of Implementing Regulation (EU) 2021/963.

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

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

(6) Not required if part! of the zootechnical certificate forms an integral part of the unique lifetime identification document issued by a breed society. In case the single lifetime identification document has been issued in accordance with Implementing Regulation (EU) 2015/262, the unique life number set out in point (o) of Article 2 of that Implementing Regulation shall be provided.

(7) If necessary, more generations shall be indicated.

(8) Indicate the individual identification number in accordance with point 3 of Chapter!Part 1 of Annex II to Regulation (EU) 2016/1012 referred to as 'unique code' in point (a) of Article 114(1) of Regulation (EU) 2016/429. If there is no individual identification number available or it is different from the number under which the animal is entered in the breeding book, enter the breeding book number.

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

## PART II

1.1. Individual identification number <sup>(1)</sup>		2. Animal identification <sup>(3)</sup>	
1.2. Unique Life Number <sup>(2)</sup>		2.1. Transponder Code <sup>(4)</sup>	
		Reading system (if not ISO 11784) <sup>(4)</sup> Barcode <sup>(4)</sup>	
3. Paul		2.2. Alternative method of identity verification <sup>(3)</sup>	
4. Class in the main section of the breeding book <sup>(4)</sup>		5. Name, address and e-mail <sup>(4)</sup> of the owner <sup>(7)</sup>	
4.1. Name of the breeding book <sup>(5)</sup>	4.2. Class in the main section <sup>(6)</sup>		
6. Additional information <sup>(4)(8)(9)</sup>			
6.1. Results of Productivity Research			
6.2. Up-to-date results of the genetic evaluation last carried out on			
..... <i>(date in format dd/mm/yyyy)</i>			
6.3. Genetic defects and genetic features of the animal in relation to the breeding programme			
6.4. Identity and result verification system <sup>(4)(10)(11)</sup>		6.5. Origin verification results & <sup>4</sup> > & <sup>10</sup> > & <sup>12</sup> >	
7. Insemination/coating <sup>(4)(5)</sup>			
7.1. Date (in format dd/mm/yyyy)			
7.2. Cover certificate number <sup>(15)</sup>			
7.3. Identification of the donor male			
7.3.1. Individual identification number <sup>(1)</sup>			
7.3.2. Unique Life Number <sup>(2)</sup> <span style="float: right;">□</span>			

7.3.3. Identity and result verification system <sup>(4)</sup> (10)(11)		7.3.4. Results of verification of origin <sup>(4)</sup>
8.1. Done at	8.2. Done on	
<i>(place of issue)</i>		<i>(date of issue in dd/mm/yyyy format)</i>
8.3. Name and function of the signatory		8.4. Signature
<i>(name and function of the person <sup>(16)</sup>to whom the issuing breed society or competent authority has conferred powers to sign this part of the zootechnical certificate, enter capital letters)</i>		

(<sup>1</sup>) The individual identification number in accordance with point 3 of Chapter I Part 1 of Annex II to Regulation (EU) 2016/1012 referred to as 'unique code' in Article 114(1)(a) of Regulation (EU) 2016/1012

(EU) 2016/429 of the European Parliament and of the Council and listed in accordance with Article 6(2) of Implementing Regulation (EU) 2021/963.

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

(<sup>3</sup>) Not required if the information corresponds to Part 7 and Parts I and II are one inseparable whole and are included or attached to the single lifetime identification document. In case the single lifetime identification document has been issued in accordance with Implementing Regulation (EU) 2015/262, the unique life number set out in point (o) of Article 2 of that Regulation shall be provided.

(<sup>4</sup>) If not applicable, not to be filled in.

(<sup>5</sup>) Required if different from point 2 of Part I.

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

(<sup>7</sup>) Not required if owner information is available and updated in other parts of the unique lifetime identification document.

(<sup>8</sup>) If necessary, add additional sheets.

(<sup>9</sup>) If the genetic information is available on a website, a reference to it may be provided instead if authorised by the competent authority in accordance with Article 32(3) of Regulation (EU) 2016/1012.

(<sup>10</sup>) Based on DNA analysis or blood type analysis.

(<sup>P</sup>) 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. Breed societies may be required 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 the data or file number referring to the database where these data are available.

(<sup>12</sup>) If required by the breeding programme.

(<sup>13</sup>) It is required in pregnant females. The information may be presented in a separate document.

(<sup>14</sup>) Delete as appropriate.

(<sup>15</sup>) If not applicable, the results of the verification of origin referred to in point 7.3.4. shall be provided.

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







**SECTION VII/SECTION VII**

<b>Only equine influenza or equine influenza using combined vaccines</b>			<b>Grippe equine seulement OU Grippe equine utilisant des VACCINS combines</b>			<b>Equine influenza only Or equine influenza using combined vaccines</b>
Registration of vaccinations			Enregistrement des vaccinations			The Vaccination Record
The details of all vaccinations of the equine animal shall be entered in a detailed and detailed manner, supplemented by the name and signature of the veterinarian.			Toute vaccination subie par l'equide doit etre portee dans le cadre ci-dessous de Fagon lisible et precise et completee par le nom et la signature du veterinaire.			Details of any 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	Location of Lieu Place	Country Pays Country	Vaccine/Vaccin/Vaccine			Name (in capital letters) and signature of veterinarian Nom (en lettres capitales) et signature du veterinaire Name (in capital letters) and signature of veterinarian
			Name Nom Name	Batch number Numero du lot Batch number	Disease(s) Maladie(s) Disease(s)	









**SECTION IX/SECTION IX**

<b>Laboratory Health Research</b>		<b>Examen de laboratoire</b>		<b>Laboratory health test</b>	
<p>The results of any test carried out for a communicable disease by a veterinarian or a laboratory authorised by the official veterinary authority of the country ('the official laboratory') must be entered in an examination and in detail by the veterinarian acting on behalf of the authority at whose request the examination is carried out.</p>		<p>Le resultat de tout examen effectue par un veterinaire pour une maladie transmissible ou par un laboratoire agree par le service veterinaire officiel du pays (laboratoire officiel) doit etre note clairement et en detail par le veterinaire qui represente l'Autorite demandant l'examen.</p>		<p>The result of any 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.</p>	
Date of sampling Date de prelevement Sampling date	Contagious disease for which Maladie transmissible-concernee Transmissible disease tested for	Type of study Nature de l'examen Type of test	Result of the Resultat de l'examen Result of Test study	Official laboratory that conducted the study Laboratoire officiel ayant effectue l'examen Official laboratory which carried out the test	Name (in capital letters) and signature of veterinarian Nom (en lettres capitales) et signaTURE du veterinaire Name (in capital letters) and signaTURE of veterinarian



Date of sampling Date de prelevement Sampling date	Contagious disease for which Maladie transmissible- concernee Transmissible dis- ease tested for	Type of study Nature de l'examen Type of test	Result of the Resultat de l'examen Result of Test study	Official laboratory that conducted the study Laboratoire officiel ayant effectue l'examen Official laboratory which carried out the test	Name (in capital letters) and signature of veterinarian Norn (en lettres capitales) et signature du veterinaire Name (in capital letters) and signature of veterinarian

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**SECTION X/SECTION X**

**Chestnuts**

The contours of the four chestnuts must be drawn in the appropriate square for all horses without marking and with less than three spins.

**Chataignes**

Dessiner le contour de chaque chataigne dans la carre correspondant: a ne remplir que pour les chevaux sans marque et avec moins de Trois EPIS

**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.

<b>Foreleg/ Anterieur droit/ Right Foreleg</b>	<b>Back leg, right/ Posterieur droit/ Right Hindleg</b>
<b>Front leg, left/ Anterieur gauche/ Left Foreleg</b>	<b>Back leg, left/ Posterieur gauche/ Left Hindleg</b>

The passport is a lifetime identification document and must accompany each movement of the equine animal.

The passport was issued on the basis of the data in the integrated information system VetIS.

The passport must be protected from damage to destruction or loss. If any of these events occur, the official veterinarian must be notified.

The passport shall be considered invalid in the case of:

- broken integrity of the self-destructive sticker or hologram
- any revision of the unique code of the transponder number or of the date of birth
- lack of signature or stamp of an official from the RFSD
- lack of a page or failure to read the information on a particular page
- death of the animal

The completion of Part II of Section II shall permanently exclude equidae from the food chain.

In the event of the death of the equine passport, the passport shall be handed over for annulment to the relevant official veterinarian.

Le passport est un document viager pour l'identification et il doit accompagner les equides chez leur mouvement/transportation.

Le passport est Edite sur la base des donnees dans la systeme informatique d'integration Vetis.

Le passport doit etre proteger d'alteration, destruction ou la perte. S'il s'agit d'un cas pareil on doit informer le veterinaire officiel.

Le passport n'est pas valide chez:

- en cas de destruction de transpondeur ou de la hologramme;
- chaque Redaction de code unique, de numero de transpondeur ou a la date de naissance;
- Le manqué de la signature ou le cachet d'employer de l'Etat du Regionale direction de securite des aliments;
- Le manqué de la page ou la destruction qui ne permete pas de lire l'information sur une page determinee;
- La Mort d'animal.

Remplissement de la partie II dans la section II exclure l'animal de la Chaine alimen-tAire.

Chez la mort des equides, le passport doit etre annuler par le veterinaire officiel.

This passport is an identification document valid for life and should accompany the equine animal during every movement/transportation.

This passport is issued on the databases in the integrated information system – VetIS.

This passport should be guarded against damage, destruction or loss. In case of the abovementioned events, it is compulsory that the official veterinarian should be notified.

This passport should be considered invalid in case of:

- distorted wholeness of the self-destroying sticker or of the hologram;
- any change of the unique code, of the transponder's number or of the birth date;
- absence of signature or stamp of the official of the Regional food safety directorate;
- absence of a page or any damage which makes the reading of the information on the current page impossible;
- death of the equine animal.

The completion of Part II of Section II excludes the equine animal from the food chain Entirely for good.

On the death of the equine animal, the passport should be forwarded for annulment to the relevant official veterinarian.