Datum van winning van de (gebruikte) oöcyten of embryo's (in vivo) voor 21 april 2021

EUROPEAN UNION

INTRA

Part I: Description of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Operator conducting assembly operations independently of an establishment
Name
Address
Approval Number
Country ISO Code

I.7. Country of origin ISO Code

I.8. Region of origin Code

I.9. Country of destination ISO Code

I.10. Region of destination Code

I.11. Place of dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16. Transporter
Name
Address
Approval Number
Country ISO Code

I.17. Accompanying documents
Document Type
Accompanying document reference
Date of Issue
Country
Place of issue

In vitro geproduceerde embryo's: unieke referentienummer van het document dat het gebruikte sperma naar de inrichting heeft begeleid.

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

I.19. Container No / Seal No

I.20. Certified as
Germinal products ☐

I.21. For transit through a third country ☐
Third country ISO Code
Exit point BCP code
Entry point BCP code

I.22. For transit through Member State(s) ☐
Member State ISO Code

I.23. For export ☐
Third country ISO Code
Exit point BCP code

I.24. Estimated journey time

I.25. Journey Log

I.26. Total number of packages

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED

0511 Animal products not elsewhere specified or included; dead animals of Chapter | 1 | or 3, unfit for human consumption

051199 Other

05119985 Other

Part I: Description of consignment

#1.	Commodity	Identification Number	Quantity	Nature of commodity	Identification Mark
Species		Package count	Date of collection / production	Plant / Establishment / Centre	Type

Part II: Certification

II. Health information

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

II.1. were collected, processed and stored in compliance with Annex A to Council Directive 89/556/EEC;

II.2. were sent to the place of loading in sealed containers in compliance with Annex A to Directive 89/556/EEC;

II.3. come from donors of the bovine species which comply with Annex B to Directive 89/556/EEC;

II.4. were conceived either by ☐ [artificial insemination] (1) ☐ [in vitro fertilisation] (1) using semen coming from semen collection or storage centres approved in accordance with Council Directive 88/407/EEC and loaded in a Member State or in a third country listed in Annex I to Commission Decision 2004/639/EC (1) (2).

In vivo verkregen

In vitro geproduceerd

II.5. is dispatched:

(1) either ✓ [by an ☐ [embryo collection team] (1) ☐ [embryo production team] (1) * or from a zone not subject to movement restrictions affecting bovine animals and established for reasons of listed diseases relevant for those species or diseases subject to emergency measures relevant for those species, or those restrictions do not apply to these embryos because they were collected before the restrictions were established, and the embryos have not been in contact with other embryos of a lower health status for an adequate period;]

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(1) or ○ [by an ☐ [embryo collection team] (1) ☐ [embryo production team] (1) * or from a zone subject to movement restrictions affecting bovine animals and established for (3), but derogations from movement restrictions have been granted, and:

(1) ☐ [they comply with the requirements set out in (4);]]

(1) ☐ [and in particular, they are (5).]]

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.17: In the case of imported embryos, insert the number of the import certificate.

Box reference I.19: Seal number shall be indicated.

Box reference I.26: Total number of packages shall correspond to the number of containers.

Box reference I.30: “Type”: Specify whether there is (a) penetration or (b) non-penetration of zona pellucida. “Identification mark”: shall correspond to the details identifying the donor cows and the date of collection on the straw.

Part II:

(1) Delete if not applicable.

(2) OJ L 292, 15.9.2004, p.21.

(3) Insert the name of the disease(s).

(4) Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.

(5) Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the

Part II: Certification

II. Health information		

Commission, as referred to in Article 159(2), points (a), (b) and (c), of Regulation (EU) 2016/429 of the European Parliament and of the Council.

Certifying Officer/Official veterinarian	
Name (in capital letters)	Qualification and title
Date of declaration	Signature
Stamp	

*

EWT = embryowinningsteam: winning van oöcyten en/of embryo's (in vivo)
EPT = embryoproductieteam: winning van oöcyten en/of productie van embryo's (in vitro) uit door het team zelf gewonnen oöcyten