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.9. Country of destination ISO Code

I.8. Region of origin 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
[en] accompanying.document.number
Date of issue
Country
Place of issue

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

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.28. Total gross weight

I.27. Total quantity

I.30. Description of consignment

Commodity	Species	Identification Number	Quantity	Nature of commodity

Identification Mark	Package count	Date of collection / production	Plant / Establishment / Centre

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

(1) ○ either [II.1. the in vivo derived embryos(1)/in vivo derived ova(1) described in Part I were collected, processed and stored by an embryo collection team(2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.1. the in vitro produced embryos(1)/micromanipulated embryos(1) described in Part I were produced, processed and stored by an embryo production team(2) approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]

(1) ○ either [II.2. the in vivo derived embryos described in Part I meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the in vivo derived ova described in Part I meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the in vitro produced embryos described in Part I meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the micromanipulated embryos described in Part I meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

☐ [II.3. the consignment consists of embryos of the ovine or caprine species which comply with the following conditions as regards classical scrapie:

(1) ○ either [they were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1) ○ or [they were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1) ○ or [they were collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with the first subparagraph of point 2.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1) ○ or [they were collected from ovine animals and

(1) ○ either [are of the ARR/ARR prion protein genotype;]

(1) ○ or [carry at least one ARR allele and were collected after the date of 1 January 2015;]]

II.4. the ova or embryos described in Part I come from female donors of the ovine(1)/caprine species(1) which meet the requirements of Chapter IV(3) of Annex D to Directive 92/65/EEC;

(1) ○ either [II.5. the embryos described in Part I were conceived as a result of artificial insemination of the donor females with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.5. the embryos described in Part I were conceived as a result of in vitro fertilisation of ova complying with the conditions in Chapter III(II)(2) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.5. the ova have not been in contact with semen of the ovine and caprine species;]

II.6. the ova or embryos described in Part I were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.19.

Part II: Certification

II. Health information

Notes

This animal health certificate shall be completed according to 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 I.11: Place of dispatch shall correspond to the embryo collection team or embryo production team of embryos collection/production.

Box I.12: Place of destination shall correspond to the embryo collection team, embryo production team, germinal product processing establishment, germinal product storage centre or to the establishment of ova/embryos destination.

Box I.19: Identification of container and Seal number shall be indicated.

Box I.30: "Type": specify if: in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.

Identification number shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Part II:

(1) Delete as appropriate.

(2) Only embryo collection or production teams approved by the competent authority and listed in accordance with Article 11(4) of Directive 92/65/EEC.

Certifying Officer/Official veterinarian

Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	

Part I: Description of consignment

I.1. Odosielateľ
Meno/názov
Adresa
Krajina Kód ISO

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Príjemca
Meno/názov
Adresa
Krajina Kód ISO

I.6. Operator conducting assembly operations independently of an establishment
Meno/názov
Adresa
Číslo schválenia
Krajina Kód ISO

I.7. Krajina pôvodu Kód ISO

I.9. Country of destination Kód ISO

I.8. Region of origin Kód

I.10. Región určenia Kód

I.11. Place of dispatch
Meno/názov
Adresa
Číslo schválenia
Krajina Kód ISO

I.12. Miesto určenia
Meno/názov
Adresa
Číslo schválenia
Krajina Kód ISO

I.13. Miesto nakládky
Meno/názov
Adresa
Číslo schválenia
Krajina Kód ISO

I.14. Date and time of departure

I.15. Dopravný prostriedok

Druh	Dokument	Identifikácia

I.16. Transporter
Meno/názov
Adresa
Číslo schválenia
Krajina Kód ISO

I.17. Sprievodné doklady
[sk] accompanying.document.number
Date of issue
Country
Place of issue

I.18. Transport conditions
Teplota okolia ☐ Mrazené ☐ Chladené ☐

I.19. Číslo kontajnera/číslo pečate

I.20. Certified as
Germinal products ☐

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

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

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

I.24. Estimated journey time

I.25. Journey Log

I.26. Celkový počet balení
I.28. Celková hrubá hmotnosť

I.27. Celkové množstvo

I.30. Description of consignment

Tovar	Druh	Identification Number	Množstvo	Nature of commodity

Identification Mark	Počet balení	Dátum zberu	Plant / Establishment / Centre

Part II: Certification

II. Zdravotné informácie

Ja, podpísaný úradný veterinárny lekár, týmto potvrdzujem, že:

(1) ○ buď [II.1. embryá získané in vivo(1)/vajíčka získané in vivo(1), opísané v časti I, odobral, spracoval a skladoval tím na odber embryí(2), ktorý je schválený a pod dohľadom v súlade s kapitolou I oddielom III bodom 1 prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.1. embryá vyprodukované in vitro(1)/embryá podrobené mikromanipulácii(1), opísané v časti I, vyprodukoval, spracoval a skladoval tím na produkciu embryí(2), ktorý je schválený a pod dohľadom v súlade s kapitolou I oddielom III bodmi 1 a 2 prílohy D k smernici 92/65/EHS;]

(1) ○ buď [II.2. embryá získané in vivo opísané v časti I spĺňajú požiadavky kapitoly III oddielu II bodu 1 prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.2. vajíčka získané in vivo opísané v časti I spĺňajú požiadavky kapitoly III oddielu II bodu 2 prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.2. embryá vyprodukované in vitro opísané v časti I spĺňajú požiadavky kapitoly III oddielu II bodu 3 prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.2. embryá podrobené mikromanipulácii opísané v časti I spĺňajú požiadavky kapitoly III oddielu II bodu 4 prílohy D k smernici 92/65/EHS;]

☐ [II.3. Zásielka pozostáva z embryí oviec alebo kôz, ktoré spĺňajú tieto podmienky, pokiaľ ide o klasickú klusavku:

(1) ○ buď [boli odobrané zvieratám, ktoré boli od narodenia nepretržite držané v chove alebo chovoch so zanedbateľným alebo kontrolovaným rizikom klasickej klusavky v súlade s kapitolou A oddielom A bodom 1 prílohy VIII k nariadeniu (ES) č. 999/2001;]

(1) ○ alebo [boli odobrané zvieratám, ktoré boli počas posledných troch rokov pred odberom nepretržite držané v chove alebo chovoch, ktoré počas posledných troch rokov pred odberom spĺňali požiadavky stanovené v kapitole A oddiele A bode 1.3. písm. a) až f) prílohy VIII k nariadeniu (ES) č. 999/2001;]

(1) ○ alebo [boli odobrané zvieratám, ktoré boli od narodenia nepretržite držané v členskom štáte alebo pásme členského štátu so statusom zanedbateľného rizika klasickej klusavky schváleným v súlade s kapitolou A oddielom A bodom 2.2 prílohy VIII k nariadeniu (ES) č. 999/2001;]

(1) ○ alebo [boli odobrané ovciam a

(1) ○ buď [majú genotyp priónového proteínu ARR/ARR;]

(1) ○ alebo [sú nositeľom aspoň jednej alely ARR a boli odobrané po 1. januári 2015;]]

II.4. vajíčka alebo embryá opísané v časti I pochádzajú z darcovských oviec(1)/kôz(1)samičieho pohlavia, ktoré spĺňajú požiadavky kapitoly IV bodu 3 prílohy D k smernici 92/65/EHS;

(1) ○ buď [II.5. embryá opísané v časti I boli počaté ako dôsledok umelej inseminácie darcovských samíc spermou, ktorá bola odobraná, spracovaná, skladovaná a prepravovaná za podmienok, ktoré sú v súlade s požiadavkami kapitoly I oddielu I, kapitoly II oddielu I a kapitoly III oddielu I prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.5. embryá opísané v časti I boli počaté ako dôsledok oplodnenia vajíčok in vitro v súlade s podmienkami v kapitole III oddiele II bode 2 prílohy D k smernici 92/65/EHS spermou, ktorá bola odobraná, spracovaná, skladovaná a prepravovaná za podmienok, ktoré sú v súlade s požiadavkami kapitoly I oddielu I, kapitoly II oddielu I a kapitoly III oddielu I prílohy D k smernici 92/65/EHS;]

(1) ○ alebo [II.5. vajíčka neboli v kontakte so spermou oviec ani kôz;]

II.6. vajíčka alebo embryá opísané v časti I boli odoslané na miesto nakládky v zaplombovanom kontajneri v súlade s kapitolou III oddielom II bodom 6 prílohy D k smernici 92/65/EHS, ktorý bol označený číslom uvedeným v kolónke I.19.

Part II: Certification

II. Zdravotné informácie

Poznámky

Tento certifikát zdravia zvierat treba vyplniť podľa poznámok k vyplneniu certifikátov uvedených v kapitole 2 prílohy I k vykonávaciemu nariadeniu Komisie (EÚ) 2020/2235.

Časť I:

Kolónka I.11:	Miesto odoslania zodpovedá tímu na odber embryí alebo tímu na produkciu embryí, ktorý vykonal odber/produkciu embryí.
Kolónka I.12:	Miesto určenia zodpovedá tímu na odber embryí, tímu na produkciu embryí, zariadeniu na spracovanie zárodočných produktov, inseminačnej stanici na skladovanie zárodočných produktov alebo zariadeniu, do ktorého sú vajíčka/embryá určené.
Kolónka I.19:	Uvádza sa identifikácia kontajnera a číslo plomby.
Kolónka I.30:	„Typ“: Uveďte, či: ide o embryá získané in vivo, oocyty získané in vivo, embryá vyprodukované in vitro alebo embryá podrobené mikromanipulácii.
	Identifikačné číslo zodpovedá úradnej identifikácii zvieraťa.
	Dátum odberu sa uvádza v tomto formáte: dd/mm/rrrr.
	Schvaľovacie číslo tímu zodpovedá tímu na odber embryí alebo tímu na produkciu embryí, ktorý vajíčka/embryá odobral/vyprodukoval.

Časť II:

(1)	Nehodiace sa prečiarknite/vymažte.
(2)	Len tímy na odber alebo produkciu embryí schválené príslušným orgánom a uvedené v súlade s článkom 11 ods. 4 smernice 92/65/EHS.

Certifikujúci úradník/Úradný ve terinárny lekár

Meno (veľkými písmenami)	Kvalifikácia a titul
Dátum podpisu	Podpis
Pečiatka	