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
Activity ID
Country ISO Code

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

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

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.25. Journey Log

I.26. Total number of packages

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Identification Number	Quantity	Nature of commodity

Identification Mark	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 ova/embryos(1) described in Part I:

II.1. were produced/collected(1), processed and stored by an embryo collection/production(1) team(2) approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC;

II.2. meet the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;

II.3. come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to Directive 92/65/EEC;

(1) ○ either [II.4. are in vivo derived embryos which:

II.4.1. were conceived as a result of artificial insemination with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1) ○ either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1) ○ or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and have been washed with trypsin;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and have been washed with trypsin;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;];]

(1) ○ or [II.4. are in vitro produced/micromanipulated(1) embryos which:

II.4.1. were conceived as a result of in vitro fertilisation with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1) ○ either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1) ○ or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]

(1) ○ or [II.4. are in vivo derived ova which originate from a Member State or region thereof:

(1) ○ either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

Part II: Certification

II. Health information

(1) ○ or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]

(1) ○ or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]

(1) ○ or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]

II.5. were sent to the place of loading in a sealed container under conditions complying with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

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

Part I:

Box I.11: Place of dispatch shall correspond to the embryo collection team or embryo production team of oocytes/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 oocytes/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 indicated in Box I.11.

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 Council Directive 92/65/EEC.

Certifying Officer/Official veterinarian

Name (in capital letters) Authority name

Date of signature Signature

Stamp