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

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

I.19. Container No / Seal No

I.20. Certified as
Transformation ☐ Processing ☐ Treatment ☐

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 net weight	I.28. Total gross weight

I.30. Description of consignment

1. 30 PHARMACEUTICAL PRODUCTS

3002 Human blood; animal blood prepared for therapeutic, prophylactic or diagnostic uses; antisera and other blood fractions and modified immunological products, whether or not obtained by means of biotechnological processes; vaccines, toxins, cultures of micro-organisms (excluding yeasts) and similar products

Antisera, other blood fractions and immunological products, whether or not modified or obtained by means of biotechnological processes

Part I: Description of consignment

300212 Antisera and other blood fractions

30021200 Antisera and other blood fractions

#1.	Commodity	Nature of commodity	Category	Manufacturing plant	Quantity
Species		Batch number	Package count	Net weight	

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

○ (1)either [II.1 The animal by-products described in Part I were obtained from kept animals:

○ (1)either [killed for the purpose of the prevention and control of:

○ (1) either [(introduce the name of the relevant category A disease) following the instructions of the competent authority in accordance with Regulation (EU) 2020/687 and are intended for processing by

○ (1) either [Method 1 to 5.]]]

○ (1) or [incineration.]]]

○ (1) or [co-incineration.]]]

○ (1)or [(introduce the name of the relevant emerging disease) following the instructions of the competent authority in accordance with emergency measures adopted by the Commission pursuant to Article 259 of Regulation (EU) 2016/429 and are intended for processing by

○ (1) either [Method 1 to 5.]]]

○ (1) or [incineration.]]]

○ (1) or [co-incineration.]]]

○ (1)or [not subject to killing by the competent authority for the purpose of prevention and control of category A diseases or emerging diseases, kept in establishments located in restricted zones established for the prevention and control of animal diseases in accordance with

○ (1) either [Delegated Regulation (EU) 2020/687,]

○ (1) or [temporary special disease control measures referred to in Article 71 of Regulation (EU) 2016/429,]

○ (1) or [emergency measures adopted by the Commission in accordance with Article 259 of Regulation (EU) 2016/429,]

and the animal by-products are moved from that restricted zones in compliance with the conditions set out in (2), for the

○ (1) either [processing by methods 1-5 as set out in Chapter II of Annex IV and in case of ensilage of by-products obtained from aquatic animals as set out in point K of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011.]]]

○ (1) or [processing or treatment by methods referred to in Annex X, Annex XI or Annex XIII to Regulation (EU) No 142/2011.]]]

○ (1) or [production of processed petfood, other than raw petfood referred to in Annex XIII to Regulation (EU) No 142/2011.]]]

○ (1) or [transformation into compost or biogas referred to in Section 1 of Chapter III of Annex V to Regulation (EU) No 142/2011.]]]

○ (1) or [II.1 The animal by-products described in Part I were obtained from wild animals of listed species found dead or killed for the purpose of the prevention and control of (introduce the name of the relevant category A disease) following the instructions of the competent authority (3) in accordance with Article 64(2)(c) of Delegated Regulation (EU) 2020/687 and are intended for processing by

○ (1) either [Method 1 to 5.]]

Part II: Certification

II. Health information

○ (1) or [incineration.]]

○ (1) or [co-incineration.]]

(1) either ☐ [II.2 Unprocessed animal by-products (unprocessed animal by-products other than hides and skins, hides and skins, colostrum, milk and dairy products, indicate as appropriate) obtained from bovine animals kept in vaccination zone I in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, points (3.5) and (3.7), to Commission Delegated Regulation (EU) 2023/361.]

(1) or ☐ [II.2 Unprocessed animal by-products (unprocessed animal by-products other than hides and skins, hides and skins, colostrum, milk and dairy products, indicate as appropriate) obtained from bovine animals kept in vaccination zone II in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, points (3.6) and (3.7), to Commission Delegated Regulation (EU) 2023/361.]

Notes

Part I:

- Box reference I.9 and I.11: delete as appropriate.
- Box reference I.12, I.13 and I.17: approval number or registration number.
- Box reference I.14: complete if different from 'I.1. Consignor'.
- Box reference I.25: for 'processing', 'treatment', 'transformation'
- Box reference I.31: Nature of commodity: 'ABPs referred to in Article 22(5) of Delegated Regulation (EU) 2020/687.' Category: 'Category 1','Category 2', 'Category 3'

Part II:

(1) Delete as appropriate.

(2) Insert the number of the relevant article(s), the title and date of publication in the Official Journal of the European Union of the relevant legal act adopted by the Commission providing those conditions or the reference to the legal act or instruction approved and made public by the competent authority providing for those conditions.

(3) See special legislation on the prevention of transmissible diseases.

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