**Commercial document**

For the transport of animal by-products and derived products not intended for human consumption in accordance with Regulation (EC) No 1069/2009 within the European Union

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| **EUROPEAN UNION** **Commercial document** | | | | | | | | | | | | | | |
| **Part I: Details of dispatched consignment** | I.1. Consignor  Name  Address  Postcode | | | | | | I.2. Document reference No | | | | I.2.a. Local reference No  …………………………………………….. | | | |
| I.3. Central competent authority *n.v.t.* | | | | | | | |
| I.4. Local competent authority *n.v.t.* | | | | | | | |
| I.5. Consignee *Merefelt Livestock Diagnostics*  Name  Address *Pannenweg 200*  *6031 RK*  Postcode *Nederweert ( the Netherlands)*  Tel. *0031 495 461236* | | | | | | I.6. | | | | | | | |
| I.7. | | | | | | | |
| I.8. Country of origin | ISO code | | I.9. Region of origin | | Code | I.10. Country of destination | | ISO code | | | I.11. Region of destination | | Code |
|  |  | |  | |  | *The Netherlands* | | | *NL* | |  | |  |
| I.12. Place of origin  Establishment *X*  Name Approval number  Address    Postcode | | | | | | I.13. Place of destination  Establishment *X* Other 🖵  Name *Merefelt Livestock Diagnostic*  Approval number *209049*  Address *Pannenweg 200*  Postcode *6031 RK Nederweert* | | | | | | | |
| I.14. Place of loading  ………………………………………. | | | | | | I.15. Date of departure | | | | | | | |
|  | I.16. Means of transport  Aeroplane 🖵 Ship 🖵 Railway wagon 🖵  Road vehicle 🖵 Other 🖵    Identification | | | | | | I.17. Transporter  Name Approval number  Address  Postcode Member State | | | | | | | |
|  | I.18. Description of commodity  *Blood/swabs of animal origin* | | | | | |  | I.19. Commodity code (CN code) *051199* | | | | | | |
|  | | | | | | I.20. Quantity  *………………….Kg* | |
|  | I.21. Temperature of products  Ambient 🖵 Chilled *X* Frozen 🖵 Controlled temperature 🖵 | | | | | | | | | | | | I.22. Number of packages  ………………………… | |
|  | I.23. Seal/Container No *n.v.t.* | | | | | | | | | | | | I.24. Type of packaging  *Box / Bag* | |
|  | I.25. Commodities certified for:  Animal feedingstuff 🖵 Technical use *X, diagnostic research* | | | | | | | | | | | | | |
|  | I.26. | | | | | | I.27. Transit through Member States 🖵  Member State ISO code  Member State ISO code  Member State ISO code | | | | | | | |
|  | I.28. Export *n.v.t.* 🖵  Third country ISO code  Exit point Code | | | | | | I.29. | | | | | | | |
|  | I.30. | | | | | | | | | | | | | |
|  | I.31. Identification of the commodities | | | | Approval number of establishments | | | | | | | | | |
| Species  (Scientific name)  *Mammal*  *Porcin* | | Nature of commodity Category Treatment type Manufacturing plant Batch number    *Blood / swab 1 n.v.t. 209049 n.v.t.* | | | | | | | | | | | |

| **COUNTRY** | | **Animal by-products/derived products not intended for human consumption** | | |
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| **Part II: Certification** | II. Health information | | II.a. Certificate reference number | II.b. |
| II.1. Declaration by the consignor  I, the undersigned, declare that:  II.1.1. the information in Part I is factually correct;  II.1.2. all precautions have been taken to avoid contamination of the animal by-products or derived products with pathogenic agents and cross-contamination between various Categories.  **Notes**  **Part I:**   * Box reference I.9. and I.11.: if appropriate. * Box reference I.12., I.13. and I.17.: approval number of registration number. In the case of processed manure indicate in Box I.13 the approval or registration number of plant or holding of destination. * Box reference I.14.: complete if different from “I.1. Consignor”. * Box reference I.25.: technical use: any use other than for animal consumption. * Box reference I.31.:   **Animal species:** For Category 3 material and products derived therefrom destined for use as feed material. Select from the following: Aves, Ruminants, Non-Ruminants, *Mammalia, Pesca, Mollusca,* Crustacea, Invertebrates*.*  **Nature of commodity**: Enter a commodity chosen from the following list: ‘apiculture by-products’, ‘blood products’, ‘blood’, ‘bloodmeal’,, ‘digestion residues’, ‘digestive tract content’, ‘dog-chews’, ‘fishmeal’, ‘flavouring innards’, ‘gelatine’, ‘greaves’, ‘hides and skins’, ‘hydrolysed proteins’, ‘organic fertilisers’, ‘pet food’, ‘processed animal protein’, ‘processed pet food’, ‘raw pet food’, ‘rendered fats’, ‘compost’, ‘processed manure’, ‘fish oil’, ‘milk products’, ‘centrifuge or separator sludge from milk processing’; ‘dicalciumphosphate’, ‘tricalciumphosphate’, ‘collagen’, ‘egg products’, ‘serum of equidae’, ‘game trophies’, ‘wool’, ‘hair’, pig bristles’, ‘feathers, ‘animal by-products for processing’, ’derived products’.  **Category**: Specify Categories 1, 2 or 3 materials.  In case of Category 3 material, indicate the point of Article 10 of regulation (EC) No 1069/2009 that refers to the animal by-product concerned (e.g. Article 10(a), Article 10(b), etc.).  In the case of Category 3 material for use in raw petfood indicate ‘3a’, ‘3b(i)’ or 3b(ii)’ depending on whether the animal by-products are referred to in Article 10(a) or in Article 10(b)(i) or (ii) of Regulation (EC) No 1069/2009.  In the case of hides and skins and products derived therefrom, indicate ‘3b(iii)’ or ‘3(n)’ depending on whether the animal by-products or derived products are referred to in Article 10 (b)(iii) or Article 10(n) of Regulation (EC) No 1069/2009.  Where the consignment is made of more than one category, indicate the quantity and if applicable the number of containers per category of materials.  **Treatment type**: For treated hides and skins indicate the treatment:  ‘(a)’ for dried;  ‘(b)’ for dry-salted or wet-salted for at least 14 days prior to dispatch;  ‘(c)’ for salted for seven days in sea salt with the addition of 2% sodium carbonate.  For Categories 1 and 2 materials describe the method of processing or transformation. Indicate the relevant processing method (choose a method form 1 to 5 referred to in Chapter III of Annex IV to regulation (EU) No 142/2011).  For Category 3 materials and derived products from Category 3 material destined for use in feed: if appropriate describe the nature and the methods of the treatment. Indicate the relevant processing method (choose a method from 1 to 7 referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011).  **Batch number**: Enter batch number or ear tag number, if applicable.  **Part II:**  *The signature must be in a different colour to that of the printing.* | | | |
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|  | Signature  Done at……………………………………………..……………………………………….…. on .  (place) (date)    (signature of the responsible person/consignor) (name, in capital letters) | | | |