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SUBMISSION FORM CYTOLOGY/PATHOLOGY COMPANION ANIMALS

Number of samples:	Authorization	Receive sticker:	Submission number: **To be filled out by GD**
biopsy ☐ dissection ☐	Date Initials		

Please fill out this form as completely as possible.

Veterinary surgeon/practice:	Country:
Vet:	Email:
Address:	Fax:
Postcode:	Phone:
City:	Rel. nr.: ☐☐☐☐☐☐☐

Please note, the report and invoice will be sent to the veterinary practice.

Animal species ☐ Dog ☐ Cat ☐ Bird ☐ Other _____

Owner / holder:	Animal name:	
Address:	Breed:	
Postcode + city:	Sex: ☐ Male ☐ Female	Neutered: ☐ Yes ☐ No
Country:	Date of birth: ☐☐ - ☐☐ - ☐☐	
Your reference:	Chip number: ☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐	

Date biopsy/ death: ☐☐ - ☐☐ - ☐☐ Date sent: ☐☐ - ☐☐ - ☐☐

Reference on result and invoice: ☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐☐

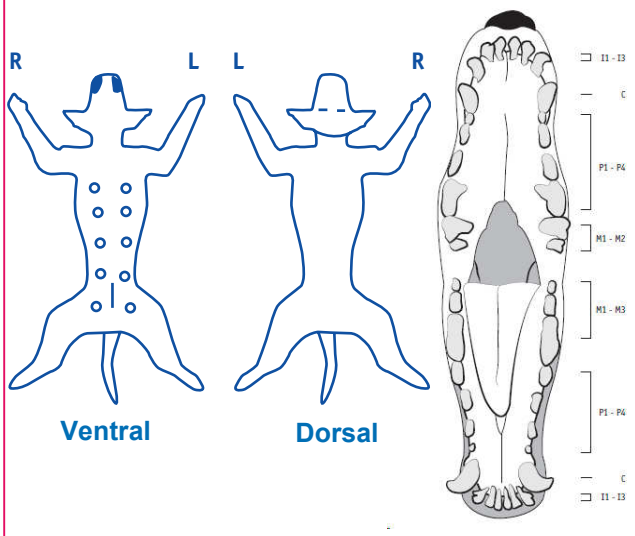
History/clinical signs/treatment (For biopsies please clearly state the following information: location, size, completely or partially removed, attached to the surrounding tissue, etc.)

Please turn over for additional writing space

Producer:	**Treatment declaration**
Name: _____ **Signature:** _____ **Date:** _____	**As producer of the microscopic preparations of animal tissues and/or animal tissue blocks and/or paraffin animal tissue sections I hereby declare that the material I submit with this form has been fixed in formalin, irradiated or treated otherwise ensuring a proven effect to inactivate any possible present pathogens. When producer's signature is missing on the submission form, GD is not allowed to receive the material and has to destroy the material upon arrival.**

Origin	Number of smears	Aspiration biopsy
		☐ Smeared and dried on air
		Smeared body fluid
		☐ Direct smear ☐ Smear after centrifugation
		Additional cytological test
		☐ Immunocytological subtyping lymphoma

Localization of the process



Histological biopsy

☐ Incision ☐ Excision

Tumour completely enclosed

☐ Yes ☐ No

☐ Fixated in 10% buffered formalin

Autopsy	Euthanasia
☒ Not cosmetic	☐ Yes ☐ No

Removal after autopsy

☐ Regular removal

☐ Cremation

To prevent the spread of pathogenic microorganisms it is not allowed to return the pet to its owner after performing autopsy.

Additional tests

In case of mast cell tumour: AgNOR and KI 67	☐ Yes	☐ No
In case of mast cell tumour: AgNOR	☐ Yes	☐ No
In case of mast cell tumour: KI 67	☐ Yes	☐ No
In case of lymphoma: immunohistochem. subtyping lymphoma	☐ Yes	☐ No
If possible further investigation lymphona (PARR) only dogs	☐ Yes	☐ No
If possible further investigation lymphona (PARR) only cats	☐ Yes	☐ No

History/clinical signs/treatment (For biopsies please clearly state the following information: location, size, completely or partially removed, attached to the surrounding tissue, etc.)

5769567601

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

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

I.4. Local competent authority

I.5. Consignee
Name GD Animal Health
Address Attn: CMD
Rigastraat 25
Postcode 7418 EW Deventer, The Netherlands
Tel. +31 (0) 570 63 33 91

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
				Deventer, The Netherlands			

I.12. Place of origin
Establishment ☒
Name
Address
Approval number
Postcode

I.13. Place of destination Deventer, The Netherlands
Establishment ☒ Other ☐
Name GD Animal Health
Address Rigastraat 25
Approval number
Postcode 7418 EW Deventer

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
Address
Approval number
Postcode
Member State

I.18. Description of commodity

I.19. Commodity code (CN code)

I.20. Quantity

I.21. Temperature of products
☐ Ambient ☐ Chilled ☐ Frozen ☐ Controlled temperature

I.22. Number of packages

I.23. Seal/Container No

I.24. Type of packaging

I.25. Commodities certified for:
☐ Animal feedingstuff ☐ Technical use ☒ For research / diagnosis only

I.26.

I.27. Transit through Member States ☐
Member State ISO code
Member State ISO code
Member State ISO code

I.28. Export ☐
Third country ISO code
Exit point Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Nature of commodity	Category	Treatment type	Approval number of establishments: Manufacturing plant	Batch number

COUNTRY **Animal by-products/derived products not intended for human consumption**

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.

Signature

Done at …… on.....

(place) (date)

.....

(signature of the responsible person/consignor)
(name, in capital letters)