



## SUBMISSION FORM PCR SAMPLES EQUINE (not for post-mortem material)

Number of samples:	Authorisation: Receipt sticker:	Submission number: <b>Te be filled out by GD</b>
Abscess content <input type="checkbox"/> Biopt <input type="checkbox"/> EDTA <input type="checkbox"/> Faeces <input type="checkbox"/> Semen <input type="checkbox"/> Lavage <input type="checkbox"/> Swab <input type="checkbox"/> Urine <input type="checkbox"/> Tissue <input type="checkbox"/> Cooled <input type="checkbox"/> Uncooled <input type="checkbox"/>	Date   Initials	Leave this box blank

Please fill out this form as **COMPLETELY** as possible

Practice: _____ Vet: _____ Adress: _____ Postal code + city + country: _____ Phone number: _____	Customer no: <input type="text"/> Submitter is veterinarian Result to veterinarian Invoice to veterinarian
--	---

Reference on result and invoice:

Pilot/Project GD-no.: \_\_\_\_\_

Sampling date:  -  -       Sampling time (hour:min) :  :

Sample number	Name	Chip number
		<input style="width:100%;" type="text"/>
		<input style="width:100%;" type="text"/>
		<input style="width:100%;" type="text"/>
		<input style="width:100%;" type="text"/>
		<input style="width:100%;" type="text"/>

**Explanation / Anamnesis**

**Client**

Name: \_\_\_\_\_

Signature: \_\_\_\_\_

Date: \_\_\_\_\_

<b>Bacteria - Regular</b>	<b>Bacteria - Urgent (note different rate!) And sample material cannot be combined with regular research!</b>	
Blood		
Faeces		
Swab		
Lavage		
Absces- content		
Urine		
Other		
<b>Viruses - Regular</b>	<b>Viruses - Urgent (note different rate!) And sample material cannot be combined with regular research!</b>	
Blood		
Biopt		
Faeces		
Semen		
Swab		
Tissue		<b>Urgent: samples must arrive at the lab by 10:00 AM on a business day for results to be posted that same day</b>
<b>Parasites</b>		
Blood		
Other		
<b>Combination packages (Parasites / Viruses / Bacteria)</b>		

### 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 Address  Postcode Tel.				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							
	I.12. Place of origin  Establishment  Name Address  Postcode				I.13. Place of destination  Establishment  Name Address  Postcode				Other  Approval number														
	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  Postcode				Approval number  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																			
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				Manufacturing plant				Batch number			

COUNTRY

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

<b>Part II: Certification</b>	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.		
	<b>Notes</b>			
	<b>Part I:</b>			
	-	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.:		
	<b>Animal species:</b>	For Category 3 material and products derived therefrom destined for use as feed material. Select from the following: Aves, Ruminants, Non-Ruminants, <i>Mammalia</i> , <i>Pesca</i> , <i>Mollusca</i> , <i>Crustacea</i> , Invertebrates.		
	<b>Nature of commodity:</b>	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'.		
	<b>Category:</b>	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.		
	<b>Treatment type:</b>	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).		
	<b>Batch number:</b>	Enter batch number or ear tag number, if applicable.		
	<b>Part II:</b>			
	<i>The signature must be in a different colour to that of the printing.</i>			
	Signature			
	Done at .....	(place)	on.....	(date)
	.....			
	(signature of the responsible person/consignor)			
	(name, in capital letters)			