

## Veterinary Microbiology Diagnostic Laboratory Submission Form

Department of Veterinary Anatomy, Physiology and Pathology, IVES, University of Liverpool, Leahurst Campus Chester High Road, Neston, CH64 7TE Cheshire
Email: vetbact@liverpool.ac.uk Tel: Bacteriology 0151-794-6118

Animal details/Patient sticker:	Please indicate sample type submitted:			
Owner Animal name	□ Swab*	-	□ CSF	
	☐ Synovial fl	uid	☐ Tracheal wash	
Species Age	☐ Faeces		☐ BAL/Tracheal wash	
Breed Sex	☐ Guttural Po		☐ Abdominal fluid	
Hospital no	☐ Tissue* (fro	<i>'</i>	□ Blood □ Milk	
Vet. Surgeon	☐ Urine:	ed nams	☐ Other *	
	□Catch □Catheter □Cysto			
Practice name	*State/Site			
PhoneE-mail	Date of sampling:			
Is there any evidence of infection? Please specify.				
<b>General Bacteriology and Mycology:</b>	<u>Skin</u>			
☐ Routine BACTERIOLOGY examination of various specimens including bacterial culture, MALDI-TOF ID and AST (Disc Diffusion or MIC, as appropriate)	<ul><li>☐ Skin/ear BACT. and fungal examination (culture, ID, SENS)</li><li>☐ Dermatophyte (direct microscopy only)</li></ul>			
☐ General FUNGAL culture & ID	☐ Dermatophyte (direct microscopy, culture and ID)			
☐ MRSA/MRSP screen (culture only, 1 swab/pooled)	☐ Skin/ear fungal culture only			
☐ Strangles culture screen (culture & ID only)				
☐ Strangles culture screen and <i>S.equi</i> MIC	<u>Faeces</u>			
☐ Bordetella bronchiseptica culture	□ Faecal bacteriology (general screen for Salmonella spp, Campylobacter spp, C. perfringens and C. difficile)			
☐ Cryptococcus neoformans (India ink stain, culture, ID)	☐ Yersinia spp.			
☐ Direct smear examination (special stains):	☐ Salmonella spp only screen			
☐ Gram ☐ Ziehl Neelsen ☐ Helicobacter spp.	AMR phenotypic screening:			
☐ MALDI-TOF ID only	_	☐ Isolate ID ESBL phenotypic testing		
☐ MIC (additional to tests where not included)	☐ Isolate ID <b>CRE</b> phenotypic testing			
PCR-based diagnostics:	Molecular detection of antimicrobial resistance:			
□ MRSA/MRSP PCR (mecA and/or mecC)	□ ESBL/pAmpC resistance genes			
□ Strangles PCR	□ CRE screening of most common CRE resistance genes			
□ Clostridium perfringens PCR (toxinotyping, enterotoxin and β2toxin detection)	□ Combined ESBL/pAmpC/CPE resistance genes screening			
Note:		For laboratory us	se only:	
Bacterial/fungal isolates are identified (ID) by MALDI-TO		Lab. no:		
itive cultures are reported less than 24h from reception of	samples	Received:		