Sample Guide - Microbiology

A guide to help you provide the correct samples. Photographs of swabs or receptacles are accompanied by their description and major indications for use

Routine Culture

Culture of fluid, washes or tissue Sterile (wrap tissue in a sterile, saline soaked plain tube swab to avoid dessication) Use for all submissions requiring Charcoal Charcoal cultures. Essential for organisms transport Swab destroyed by light (CEMO); preferred media for Bordetella bronchiseptica No Use for PCR. Dry swabs can be Plain Swab moistened in sterile saline prior to transport media sampling Viral Virology transport **Use for Virus isolation** Swab media

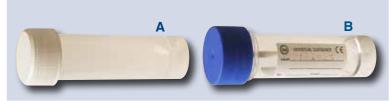
Urinalysis and culture

Fill the tube to the line to avoid Boric Acid Bacteriostatic Urine culture, when transit time (Red top) to the laboratory >2 hrs Gram-ve organisms Plain Tube Urine sediment For samples obtained by examination and cystocentesis preservative chemical analysis may not be required

too high a concentration of boric acid which may have a bactericidal effect on some

Faecal analysis and culture

Universal Containers (A) without spatula or (B) with spatula



A minimum of 10g of faeces is required

Faecal pathogens may be shed intermittently, where possible submit 3 samples collected at 48 hour intervals as a single pooled sample

If submitting feline faeces for Tritrichomonas PCR, multiple samples should be collected over 24 hours and submitted without cat litter

Blood and Joint fluid culture

Samples will be incubated for up to 10 days prior to culture

Joint fluid culture

Inoculate 5ml of Fast Anaerobe broth with 0.5ml of synovial fluid



Blood culture

Inoculate 50ml of Fast Anaerobe broth with 5ml of blood

