

TOPIC 1 – Antimicrobial Stewardship

KEY STEPS



- Antimicrobial resistance (AMR) is a growing threat to modern human and veterinary healthcare.
- This one-health problem requires concerted effort by us all.
- We must **replace**, **reduce** and **refine** our antibiotic use.

Dr. Tim Nuttall

HEAD OF DERMATOLOGY, ROYAL (DICK)
SCHOOL OF VETERINARY STUDIES,
UNIVERSITY OF EDINBURGH



Dr Tim Nuttall is the Head of Dermatology at the Dick Vet School and is a globally recognized veterinary specialist in dermatology, canine atopic dermatitis and antimicrobial resistance.

Tim has written over 100 clinical / scientific publications and has served on RCVS, BSAVA, ESVD and DEFRA scientific committees as well as the Controlling Antimicrobial Resistance in Scotland (CARS) Steering Group. His current research programme is studying antimicrobial resistance and skin infections. In 2014 he received the BSAVA Woodrow Award for outstanding contributions to veterinary medicine.

Improving antimicrobial stewardship – The 3 R's

| REPLACE | REDUCE | REFINE |
|--|---|---|
| Use alternatives to antibiotics | Reduce antibiotic prescribing – audit & benchmark use Reduce owner expectations for antibiotic treatment | Diagnose bacterial infections |
| Practice preventative care | | Use the right drug for the right bug |
| Treat the underlying disease | | Use 1st tier & narrow-spectrum antibiotics |
| Use symptomatic treatment - let the immune system tackle the infection | | Justify use of 2nd /3rd tier & broad-spectrum antibiotics |
| Use antiseptics | | Treat to clinical cure – avoid over-treating |

Key steps in antimicrobial stewardship

| | |
|---|---|
| 1 | <ul style="list-style-type: none"> Confirm a bacterial infection Avoid 'just in case' treatment Avoid use of antibiotics when unlikely to impact outcomes e.g. vomiting, diarrhea, etc |
| 2 | <ul style="list-style-type: none"> Does the infection need systemic antibiotics? Think about alternatives |
| 3 | <ul style="list-style-type: none"> For systemic treatment always use the lowest tier & most narrow spectrum drug Don't use broad-spectrum drugs by default |
| 4 | <ul style="list-style-type: none"> Treat to clinical cure Avoid overly long courses of antibiotics |
| 5 | <ul style="list-style-type: none"> Animals with recurrent infections do not have antibiotic deficiencies Diagnose and manage the underlying condition |

Diagnosis of bacterial infections

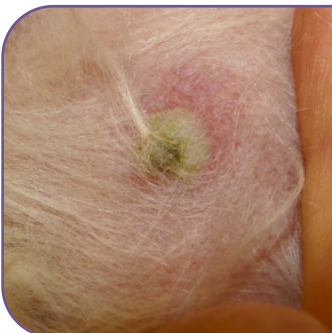
The approach to diagnosis should follow:



SITE

Site of infection – think about the most likely bacteria found there.

| INFECTION | COMMON PATHOGEN(S) |
|---|---|
| Pyoderma | Staphylococci |
| Otitis externa | Staphylococci; <i>Pseudomonas</i> ; <i>Malassezia</i> ; occ. other Gram+ or Gram- bacteria |
| Otitis media | Staphylococci; <i>Pseudomonas</i> ; occ. other Gram+ or Gram- bacteria |
| Bites | Staphylococci; streptococci, <i>Pasteurella</i> ; <i>Prevotella</i> ; fusobacteria; other oral flora; mixed infections common |
| Cystitis | <i>E. coli</i> ; other coliforms; staphylococci; enterococci; streptococci |
| Prostatitis | As above |
| Gingivitis & periodontitis | Spirochetes; <i>Porphyromonas</i> ; <i>Prevotella</i> ; and other oral flora inc. anaerobes; mixed infections common |
| Cholangio-hepatitis | <i>E. coli</i> ; less commonly other coliforms; enterococci; staphylococci; streptococci; clostridia; <i>Actinomyces</i> ; corynebacteria; and many others |
| Anal sacs | <i>E. coli</i> ; <i>Enterococcus</i> ; <i>Proteus</i> ; clostridia; occ. other Gram+ or Gram- bacteria; mixed infections common |
| Conjunctivitis | Cats: <i>Chlamydophila felis</i> ; viruses; <i>Mycoplasma</i> ; occ. Gram+ or Gram- bacteria. Dogs: staphylococci; streptococci; <i>E. coli</i> ; <i>Proteus</i> |
| Upper respiratory tract | <i>Bordetella</i> ; <i>Mycoplasma</i> ; viruses; occ. Gram+ or Gram- bacteria |
| Pneumonia Pyothorax | Dogs: usually Gram- aerobes; occ. anaerobes; mixed infections common Cats: <i>Pasteurella</i> ; <i>Actinomyces</i> ; other Gram- aerobes/anaerobes; occ. <i>Nocardia</i> . |
| Mastitis | <i>E. coli</i> ; staphylococci; streptococci |
| Osteomyelitis Discospondylitis Septic arthritis | Gram+ cocci most likely but a variety of organisms can be seen (inc. anaerobes) |
| Septicaemia | Staphylococci and <i>E. coli</i> most common but a wide variety of aerobes and anaerobes can be seen |



This dog has a bullous pustule

– These are almost always associated with *Staphylococcus pseudintermedius*.

SIGNS

Signs - bacterial infections have a consistent clinical presentation – this is especially true for the skin (see table).

| CLINICAL SIGNS | LIKELY DIAGNOSIS | DEPTH |
|--|--|---|
| Seborrheic | Bacterial overgrowth syndrome | Surface |
| Papules, pustules, epidermal collarettes and scaling | Superficial folliculitis Impetigo Superficial spreading pyoderma | Superficial |
| Erosions and ulcers | Pyotraumatic dermatitis Intertrigo | Surface to superficial Deep if ulcerated |
| Ulcers and draining sinus tracts | Furunculosis Deep pyoderma | Deep |
| Nodules and swelling | Cellulitis Abscess | Deep |



Epidermal collarettes are distinctive lesions that are highly specific for staphylococcal skin infections.

CYTOLOGY & CULTURE

- **Cytology** will reliably confirm the presence of infection and indicate the most likely type of bacteria (e.g. rods or cocci); most techniques are easy, quick and cheap.
- Where necessary, **culture and antimicrobial susceptibility tests (ASTs)** will identify the bacteria and their antimicrobial susceptibility/resistance.
- **Do not** use ASTs to select topical treatment – AST results and clinical outcomes are poorly correlated.

| EMPIRICAL THERAPY IS APPROPRIATE | WHEN IS AMR MORE LIKELY? |
|--|---|
| Topical therapy | After multiple antibiotic courses |
| First episode of infection | Non-healing wounds |
| No systemic antimicrobial therapy within 3 months | Post-operative infections |
| Simple infections (e.g. surface/superficial pyodermas) | Nosocomial (i.e. healthcare associated) infections |
| Cytology shows bacteria with predictable antibiotic susceptibility (e.g. staphylococci in skin infections) | Prolonged hospitalisation and/or ICU care |
| AMR is unlikely | Implants (e.g. catheters, urinary devices, feeding tubes, surgical implants etc.) |
| LIFE-THREATENING INFECTIONS THAT NEED IMMEDIATE TREATMENT | CULTURE AND ASTS ARE NECESSARY |
| Start empirical treatment based on history, clinical signs and cytology | Deep and/or complex infections |
| Collect samples for culture & AST | Unusual clinical signs/cytology |
| Review treatment when results available | Rod bacteria |
| | If empirical treatment fails |
| | If AMR is more likely |

Does the infection need antibiotic therapy?

- Systemic antibiotics affect the whole microbiome not just the site of infection.
- For mild, non-life threatening, focal, surface or superficial infections:
 - Manage the primary disease
 - Use supportive care
 - Consider topical antiseptics
 - Consider topical antibiotics



This dog has a pyotraumatic skin lesion with a mixed bacterial infection; however, it doesn't need systemic antibiotics

- It will respond to topical cleansing and management of its atopic dermatitis.

Antibiotic selection

If systemic treatment is justified, select an appropriate antibiotic using the site, signs, cytology and culture approach.



Follow treatment guidelines: always aim for the lowest tier & most narrow spectrum antibiotic appropriate for the infection.

Also consider:

Can the antibiotic penetrate to the site of infections? Think about the impact of biofilms, pus, necrosis, scarring & blood-tissue barriers.

Can the owner safely and reliably administer the antibiotic? Use precise instructions (e.g. 'every 12 hours' not 'twice daily') and help make administration easier (flavoured medication, pill pockets or treats, demonstration & training etc.).

Only use long acting injectable antibiotics as a last resort.

TREAT TO CLINICAL CURE

Always treat to clinical cure – i.e. resolution of clinical signs and normal cytology.

There may be residual inflammation or clinical signs associated with underlying condition (e.g. atopic dermatitis) – this does not justify prolonged courses of treatment.

Culture can identify the end-point in some cases (e.g. in normally sterile sites and/or with non-commensal bacteria such as *Pseudomonas*).

Avoid overly long courses of treatment – these do not improve the clinical outcome and increase the selection pressure for AMR.

RECURRENT BACTERIAL INFECTIONS

1

Recurrent bacterial infections are always secondary.

2

Animals with recurrent infections do not have antibiotic deficiencies; they have underlying conditions.

3

The underlying condition must be diagnosed and managed.

4

Repeated antibiotic courses are the biggest single driver of AMR – these must be avoided.

REFERENCES AND RESOURCES

FECAVA posters and guidelines - <https://www.fecava.org/policies-actions/guidelines/>

BSAVA PROTECT ME antibiotic use guidelines - <https://www.bsava.com/Resources/Veterinary-resources/PROTECT-ME>

International Society for Companion Animal Infectious Diseases - www.iscaid.org

RCVS Knowledge AMR Hub - <https://knowledge.rcvs.org.uk/amr/>

USE MEDICINES RESPONSIBLY | www.noah.co.uk/responsible

For further information contact your local sales representative or customer services. Krka UK Ltd, Thames House, Waterside Drive, Langley, SL3 6EZ, United Kingdom
Tel: 020 7164 6156 | Email: info.uk@krka.biz | www.krka.co.uk | Copyright © KRKA 2023, All rights reserved. ® Registered trademarks of Krka d.d. Novo Mesto.

