

PRESCRIBING PRINCIPLES FOR ANTIMICROBIALS

Antimicrobial resistance (AMR) is a global concern in both human and veterinary medicine. Whilst the Pig Veterinary Society believes that the primary responsibility of the prescribing veterinary surgeon is to the animals under their care, it is also vital that measures are implemented to minimise the use of antimicrobials, and where antimicrobial therapy is necessary, that prescribing is done responsibly.

PVS ANTIMICROBIAL PRESCRIBING GUIDANCE

- The basis for responsible prescribing is a diagnosis, and the prescribing vet should confirm a clinical diagnosis with diagnostic investigations, including antimicrobial sensitivity testing, as appropriate.
- 2. The use of antimicrobials should be minimised by:
 - a. Encouraging good farm management
 - b. Optimising the animal's environment
 - c. Implementing appropriate vaccination regimes
 - d. Considering routes of administration in order that any course of treatment can be administered most effectively, and to the minimum number of pigs requiring treatment, and with due regard to possible environmental contamination

The above should be addressed by on-going veterinary advice, including the Veterinary Health Plan, which should be regularly reviewed and updated.

- 3. Whenever possible, the prescribing vet should adhere to the product's SPC and to the prescribing cascade: https://www.gov.uk/guidance/the-cascade-prescribing-unauthorised-medicines
- 4. Antimicrobial use should be reviewed on every farm at least annually and this review should be recorded in the farm's Veterinary Health Plan.
- 5. Antimicrobial selection should be considered in accordance with the European Medicine Agency's (EMA) antimicrobial classification guidelines; a summary of this and PVS recommendations can be found in the table overleaf.

Further Resources

EMA's categorisation of antibiotics in the European Union:

https://www.ema.europa.eu/en/documents/report/categorisation-antibiotics-european-union-answer-request-european-commission-updating-scientific_en.pdf

RUMA advice on antimicrobials in pig production:

https://www.ruma.org.uk/pigs/responsible-use-antimicrobials-pig-production/

BVA responsible use of antimicrobials policy:

https://www.bva.co.uk/take-action/our-policies/responsible-use-of-antimicrobials/



PVS Category	Group	Examples
Class One (EMA Category D) Use with Prudence	TETRACYCLINES	TETRACYCLINE OXYTETRACYCLINE CHLORTETRACYCLINE DOXYCYCLINE
First line treatment choice whenever possible	DIAMINOPYRIMIDINES & SULPHONAMIDES	TRIMETHOPRIM/SULPHA
Use prudently and only when clinically required	PENICILLINS	PHENOXYMETHYL PENICILLIN PROCAINE PENICILLIN AMOXYCILLIN AMPICILLIN
	AMINOGLYCOSIDE	SPECTINOMYCIN
Class Two (EMA Category C) Use with Caution • Choose only when there are no clinically effective alternatives in Category D	BETALACTAMS PLUS BETA LACTAMASE INHIBITORS AMINOGLYCOSIDES	AMOXYCILLIN + CLAVULANIC ACID APRAMYCIN NEOMYCIN PAROMOMYCIN STREPTOMYCIN
	PLEUROMUTILINS	TIAMULIN VALNEMULIN
 Prescribing should ideally be supported by sensitivity testing 	PHENICOLS	FLORFENICOL
It should be considered that equivalents used within human medicine exist within this category	LINCOSAMIDES	LINCOMYCIN
	MACROLIDES	TYLOSIN TYLVALOSIN TILMICOSIN TULATHROMYCIN TILDIPIROSIN
Class Three (EMA Category B)- HP-CIA's	FLUOROQUINOLONES	ENROFLOXACIN MARBOFLOXACIN
Restrict Use	3 RD /4 TH GEN CEPHALOSPORINS	CEFTIOFUR CEFQUINOME
Last resort products	POLYMYXINS	COLISTIN
Only consider use when no antibiotics in Category C or D could be clinically effective		
Sensitivity testing to support use		
Antibiotics in this category are <u>critically</u> <u>important</u> in human medicine and use in animals should be restricted to mitigate the risk to public health		
EMA Category A	These medicines may not be used in food-producing animals	

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GUIDANCE NOTE FOR THE USE OF ANTIMICROBIALS UNDER THE CASCADE

This document outlines examples of the use of products under the Cascade and offers guidance on the implementation thereof.

The Cascade permits veterinary surgeons the clinical freedom, on a case-by-case basis, to prescribe the most appropriate product for the animals under their care.

In departing from the clinical particulars (section 4) of the SPC, the veterinary surgeon must balance the benefits against the risks of doing so, and thus take responsibility for their clinical decision.

(*)Risk could relate to the animal, the owner or person administering the product, consumers (where veterinary medicine residues in food might be affected), AMR, the environment and even wider public health.

Considerations for the proposed use of a product must include full understanding of the SPC, (*)risk factors, advice from the MAH and pharmacological properties (section 5 of the SPC). Additional considerations and guidance are shown in the table below.

Contraindications, special warnings and precautions (SPC sections 4.4 - 4.8 and 4.10) should be adhered to.

This whole document is intended as a practical guide to support members in their prescribing decisions and is not intended as a complete summary of the legislative or other framework surrounding these issues, nor as a legal guide.

Examples of use of products under the Cascade are given overleaf together with guidance to consider.

Further Resources:

The Veterinary Medicines Regulations (2013) and Amendment Regulations (2014): http://www.legislation.gov.uk/uksi/2013/2033/contents/made https://www.legislation.gov.uk/uksi/2013/2033/contents/made

Pig Veterinary Society: Guidance for members in relation to medicated feed and MFSp completion: https://www.pigvetsoc.org.uk/resources/pvs-documents

Responsible use of antibiotics under the 'Cascade':

https://www.gov.uk/guidance/responsible-antibiotic-use-under-the-prescribing-cascade

Glossary:

MAH Marketing Authorisation Holder

WDP Withdrawal Period

SPC Summary of Product Characteristics

SIC Special Import Certificate
STC Special Treatment Certificate

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Clinical	Details	Examples	WDP and other
Particulars Indication	The indication is outwith	Use of Amoxicillin to treat	Advice The specified
mulcation	that specified in section 4.2 of the SPC, but the species, dose rate and length of treatment are consistent with the SPC	Streptococcal meningitis	withdrawal period on the SPC may be applied
Administration route	The route of administration of the proposed product falls outwith section 4.9 of the SPC	Administration of a product via liquid feed	Consider increasing WDP above that on SPC and consult MAH for further advice
		Administration via top dressing, when not specified on the SPC	Not supported for use in the herd. Consult the MAH
		A molecule or brand is licensed in feed but not via water medication (or <i>vice versa</i>)	Consider increasing WDP above that on SPC and consult MAH for further advice
Species	The product is licensed for use in another livestock species but not pigs	Use of Neomycin/ Streptomycin oral solution	Apply the statutory WDP of the Cascade (28 days), or longer if appropriate.
Dose Rate	The proposed dose rate is below that of the SPC	PVS would only consider acceptable to use any POM-V at dose rates below those indicated on the SPC where there is sound clinical and/or research evidence	Based on sound clinical evidence, the specified WDP on the SPC may be applied
	The proposed dose rate is above that of the SPC	Elimination programmes	Consider increasing WDP above that on SPC and consult MAH for further advice
Duration of Treatment	The proposed duration of treatment is shorter than the SPC	PVS would only consider acceptable to use any POM-V for shorter periods than those indicated on the SPC where there is sound clinical and/or research evidence	Based on sound clinical evidence, the specified WDP on the SPC may be applied
	The proposed duration of treatment is longer than the SPC	Elimination programmes	Apply the statutory WDP of the Cascade (28 days) or longer, if appropriate
Concurrent Use of Products	Use of two licensed products	Tiamulin + Chlortetracycline to treat enzootic pneumonia	Apply the longer of two WDPs, if the two products are used within the terms of the SPC
No VM Number Exists	The product is licensed in another EU country and imported under SIC	No national (UK) licence exists e.g. there is supply chain disruption	Apply WDP as per SPC in other EU country

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