European Union comments on
Codex Circular Letter CL 2017/83-AMR:

PROPOSED DRAFT REVISION OF THE CODE OF PRACTICE TO MINIMIZE AND CONTAIN ANTIMICROBIAL RESISTANCE (CAC/RCP 61-2005)

(for comments at Step 3 through CL 2017/83-AMR)

Mixed Competence
European Union Vote

This document contains contributions from Codex Members and Observers, including potential areas of consensus, identified in the Electronic Working Group (EWG) held 26 July to 31 August 2017. New and revised text in [square brackets] is proposed for further comment and potential adoption at TFAMR5. Similarly, the working group identified potential consensus around existing text that could be deleted. All text proposed for replacement or deletion is accordingly noted by strikethrough for further comment and consideration at TFAMR5.

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[List of Acronyms Used in the Document]
ADI Acceptable Daily Intake
AMR Antimicrobial Resistance
AMU Antimicrobial Use
CAC Codex Alimentarius Commission
CAC/RCP Codex Alimentarius Commission/Recommended Code of Practice
CCRVDF Codex Committee on Residues of Veterinary Drugs in Foods
FAO Food and Agriculture Organization of the United Nations
MRL Maximum Residue Limit
OIE Office International des epizooties/International Office of Epizooties, World Organisation for Animal Health
VICH International Cooperation on Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products
VMP Veterinary Medicinal Product
WHO World Health Organization
Introduction

1. Antimicrobial resistance poses a complex, global public health challenge. Within the food production to consumption continuum, there is a need to address the emergence, selection and dissemination of resistant microorganisms and resistance determinants. The development of strategies for good practices in agriculture (crops), aquaculture and animal husbandry including the responsible and prudent use of antimicrobials agents in all sectors following a One Health approach will form a key part of multi-sectoral national action plans to address risks of foodborne antimicrobial resistance.

2. This Code of Practice is an integral part of risk analysis focusing on risk management options and should be read in conjunction with other Codex texts including the Proposed Draft Guidelines on Integrated Surveillance (CAC/GL xx-xxxx) and the Guidelines for Risk Analysis of Foodborne Antimicrobial Resistance CAC/GL 77-2011. In addition, the Codex Code of Hygienic Practice for Fresh Fruits and Vegetables (CAC/RCP 53-2003) and the Code of Practice on Good Animal Feeding (CAC/RCP 54-2004), and Principles and Guidelines for the Conduct of Microbiological Risk Management (CAC/GL 63-2007) are particularly relevant for use of agricultural chemicals on crops and animal feed, respectively. WHO guidelines on Integrated surveillance of antimicrobial resistance in foodborne bacteria, Application of a One Health Approach (2017) and Critically Important Antimicrobials for Human Medicine (2016) and relevant chapters of the OIE Terrestrial and Aquatic Animal Health Codes and the List of Antimicrobials of Veterinary Importance should also be considered.

3. Where available, national and local guidelines to minimize and contain antimicrobial resistance should be taken into consideration. Best practices and guidelines on the responsible and prudent use of antimicrobials developed by governmental and professional organizations should also be considered.

4. This document provides additional guidance for on relevant measures along the food chain to minimize the development and spread of foodborne antimicrobial resistance, including guidance on] the responsible and prudent use of antimicrobials in [the] food [chain]-producing animals, and should be read in conjunction with the Recommended International Code of Practice for Control of the Use of Veterinary Drugs CAC/RCP 38-1993. Its objectives are [part of a One Health approach] to minimize the potential adverse impact on human health resulting from the use of antimicrobial agents in [the] food [chain]-producing animals, in particular the development of antimicrobial resistance. It is also important to provide for the safe and effective use of veterinary antimicrobial [agents] drugs in veterinary medicine by maintaining their efficacy.

This document defines the respective responsibilities of authorities and relevant stakeholders involved in the authorization, production, control, distribution and use of veterinary antimicrobials such as the national regulatory authorities, the veterinary pharmaceutical industry[manufacturers], veterinarians and plant health professionals, [wholesale and retail] distributors[,] and [food] producers[,] and consumers of food-producing animals.

2. The marketing authorization procedure has a significant role in establishing the basis for [the responsible and] prudent use of veterinary antimicrobial drugs [agents] in food-producing animals through clear label indications, directions and warning statements.

3. A number of codes of practice relating to the use of veterinary antimicrobial drugs and the conditions thereof have been developed by different organisations. These codes were taken into consideration and some elements were included in the elaboration of this Code of Practice to Minimize and Contain Antimicrobial Resistance.

4. In keeping with the Codex mission, this Code [of Practice] focuses on antimicrobial use in [the] food-producing animals[chain]. It is recognized that [the use of antimicrobial agents in the food chain may result in exposure in the environment]. As part of a One Health strategy to minimize and contain antimicrobial resistance, only authorized products should be used and best practices in the food production sector should be followed to minimize the occurrence/persistence in the environment of antimicrobials and their metabolites from anthropogenic sources, and to minimize the risks associated with the selection (enhancement) and dissemination of resistant microorganisms and resistance determinants in the environment. Antimicrobial resistance is also an ecological problem and that management of antimicrobial resistance may require addressing the persistence of resistant microorganisms in the environment. Although this issue is most relevant for CCRVDF with respect to food-producing animals, the same principles apply to companion animals, which also harbor resistant microorganisms.
[Scope]

[7. This Code of Practice addresses the risk to human and animal health associated with the presence in food-producing animals, food and animal feed, and the transmission by direct contact with food-producing animals and through food and animal feed, of antimicrobial resistant microorganisms or resistance determinants. It provides [risk-based] guidance on relevant measures along the food chain to minimize and contain the development and spread of foodborne antimicrobial resistance, including guidance on good practices in agriculture (crops), aquaculture and animal husbandry and guidance on the responsible and prudent use of antimicrobial agents in agriculture (crops), animal husbandry, and aquaculture. Its objectives are to minimise the potential adverse impact on human and animal health resulting from the use of antimicrobial agents. Environmental aspects of AMR should be considered when implementing this Code. All actors involved in the authorisation, manufacture, sale and supply, prescription and use of antimicrobials in the food chain together with those involved in the handling, preparation, distribution and consumption of food have a role to play in optimizing the use of antimicrobial agents and/or limiting the spread of resistant microorganisms and resistance determinants.]

[8. As there are existing Codex or internationally recognized guidelines, the following areas related to antimicrobial agents or AMR are outside the scope of this document: residues of antimicrobial agents in food; AMR marker genes in recombinant-DNA plants and recombinant DNA microorganisms¹; nongenetically modified microorganisms (for example, starter cultures) intentionally added to food with a technological purpose ²; and certain food ingredients, which could potentially carry antimicrobial resistance determinants, such as probiotics³.]

1. **Alims and Objectives**

5. It is imperative that all who are involved in the authorisation, manufacture, sale and supply, prescription and use of antimicrobials in food-producing animals act legally, responsibly and with the utmost care in order to limit the spread of resistant microorganisms among animals so as to protect the health of consumers.

6. Antimicrobial drugs are powerful tools for the management of infectious diseases in animals and humans. This Code and existing guidelines for the responsible use of antimicrobial drugs in food-producing animals include recommendations intended to prevent or reduce the selection of antimicrobial-resistant microorganisms in animals and humans in order to:

   - Protect consumer health by ensuring the safety of food of animal origin intended for human consumption.
   - Prevent or reduce as far as possible the direct and indirect transfer of resistant microorganisms or resistance determinants within animal populations and from food-producing animals to humans.
   - Prevent the contamination of animal-derived food with antimicrobial residues which exceed the established MRL.
   - Comply with the ethical obligation and economic need to maintain animal health.

7. This Code does not address environmental issues related to antimicrobial resistance from the use of veterinary antimicrobial drugs but it encourages all those involved to consider the ecological aspects when implementing the Code. Efforts should be made to ensure that environmental reservoirs of veterinary antimicrobial drugs, antimicrobial-resistant organisms and resistance determinants are kept to a minimum. In particular:

   - Regulatory authorities should assess the impact of proposed veterinary antimicrobial drug use on the environment in accordance with national guidelines or recognized international guidelines.²
   - Research should be conducted on resistant microorganisms in the environment and the magnitude of resistance determinant transfer among microorganisms in the environment.

8. The responsible use of veterinary antimicrobial drugs in food-producing animals:

   1. The food safety assessment on the use of antimicrobial resistance marker genes in recombinant-DNA plants is addressed in the Guideline for the Conduct of Food Safety Assessment of Foods Derived from Recombinant-DNA Plants (CAC/GL 45-2003).]
   2. The food safety assessment on the use of antimicrobial resistance marker genes in recombinant-DNA microorganisms is addressed in the Guideline for the Conduct of Food Safety Assessment of Foods Produced Using Recombinant-DNA Microorganisms (CAC/GL 46-2003).]
is controlled by the veterinary profession or other parties with the required expertise, and by the relevant authorities.

- is part of good veterinary and good animal husbandry practice and takes into consideration disease prevention practices such as the use of vaccination and improvements in husbandry conditions.

- aims to limit the use of veterinary antimicrobial drugs according to their approved and intended uses, and takes into consideration on-farm sampling and testing of isolates from food-producing animals during their production, where appropriate, and makes adjustments to treatment when problems become evident.

- should be based on the results of resistance surveillance and monitoring (microbial cultures and antimicrobial sensitivity testing), as well as clinical experience.

- does not include the use for growth promotion of veterinary antimicrobial drugs that belong to or are able to cause cross resistance to classes of antimicrobial agents used (or submitted for approval) in humans in the absence of a risk analysis. This risk analysis should:
  - be undertaken by the appropriate national regulatory authority;
  - be based on adequate scientific evidence; and
  - focus on the potential to impact resistance to antimicrobials used in human medicine.

- is aimed at all the relevant parties, such as:
  - regulatory and scientific authorities;
  - the veterinary pharmaceutical industry;
  - distributors and others handling veterinary antimicrobial drugs;
  - veterinarians, pharmacists and producers of food-producing animals.

[Definitions]

[Antimicrobial agent: Any substance of natural, semi-synthetic, or synthetic origin that at in vivo concentrations kills or inhibits the growth of microorganisms by interacting with a specific target. The term antimicrobial is a collective for antiviral, antibacterial (i.e. antibiotics), antifungal, and antiprotozoal agents.]

[Antimicrobial Resistance (AMR): The ability of a microorganism to multiply or persist in the presence of an increased level of an antimicrobial agent relative to the susceptible counterpart of the same species.]

[Antimicrobial Resistance Determinant: The genetic element(s) encoding for the ability of microorganisms to withstand the effects of an antimicrobial agent. They are located either chromosomally or extra-chromosomally and may be associated with mobile genetic elements such as plasmids, integrons or transposons, thereby enabling horizontal transmission from resistant to susceptible strains.]

[Antibiotic: A naturally derived substance that acts against microorganisms, specifically bacteria.]

[Antibiotic resistance: The ability of a microorganism, specifically bacteria, to multiply or persist in the presence of an increased level of an antibiotic relative to the susceptible counterpart of the same species.]

[Antibacterial: A substance that acts against bacteria. The term does not include biocides (antiseptics and disinfectants).]

[Medically important antimicrobials: Antimicrobial agents important for therapeutic use in humans.]

[Infectious diseases-related use: Administration of antimicrobial agents for the treatment, control/metaphylaxis and prevention/prophylaxis of disease.]

[Treatment of disease: Administration of antimicrobial agents to infected individuals or populations to resolve clinical signs, infection or illness.]

[Control of disease/metaphylaxis: Administration of antimicrobial agents to populations which contain healthy and infected individuals to minimize or resolve clinical signs, infection or illness.]

It could be useful to include an explanatory diagram that could further clarify the proposed definitions and distinguish between infectious diseases-related uses and uses not related to infectious diseases.
[**Prevention of disease/prophylaxis:** Administration of antimicrobial agents to healthy individuals in a population at risk of a specific disease, prior to the onset of clinical signs, with appropriate professional oversight, dose, and duration.]

**Growth promotion:** Administration of antimicrobial agents to increase the rate of weight gain and/or the efficiency of feed utilization in animals by other than purely nutritional means. The term does NOT apply to the use of antimicrobials for the specific purpose of treating, controlling, or preventing infectious diseases, even when an incidental growth response may be obtained.

**Cross-Resistance:** The ability of a microorganism to multiply or persist in the presence of other members of a particular class of antimicrobial agents or across different classes due to a shared mechanism of resistance.

**Marketing Authorization:** Process of reviewing and assessing a dossier to support a medicinal product to determine whether to permit its marketing (also called licensing, registration, approval, etc.), finalized by granting of a document also called marketing authorization (MA) (equivalent: product license).]

**One Health approach:** A collaborative, multisectoral, and trans-disciplinary approach - working at the local, regional, national, and global levels - with the goal of achieving optimal health outcomes recognizing the interconnection between humans, animals, plants, and their shared environment.

**General Principles to Minimize and Contain Antimicrobial Resistance**

**Principle 1:** A One Health approach should be considered, wherever possible and applicable, when identifying, selecting, implementing and evaluating AMR risk management options.

**Principle 2:** Good production practices and disease prevention practices such as biosecurity, adequate nutrition, vaccination, improved production practices, and alternatives to antimicrobial agents should first be considered to reduce the need for use of antimicrobial agents.

**Principle 3:** Species or sector-specific guidelines on responsible and prudent use of antimicrobial agents should be developed, implemented, and reviewed on a regular basis to maintain their effectiveness in reducing the risk of foodborne antimicrobial resistance. Such guidelines should be developed on a national or regional level according to the local circumstances of AMR and availability of treatments. These guidelines could be included as a part of national action plans on antimicrobial resistance with development and dissemination shared among countries and organisations.

**Principle 4:** The WHO list of Critically Important Antimicrobials, the OIE List of Antimicrobials of Veterinary Importance, and national lists, where available, should be used to set priorities for risk assessment and risk management. The lists should be regularly updated and consolidated.

**Principle 5:** Responsible and prudent administration in food-producing animals does not include the use for growth promotion of antimicrobial drugs.

**Principle 6:** Medically important antimicrobial agents should only be used for infectious diseases-related purposes (treatment, control/metaphylaxis or prevention/prophylaxis of disease); or in certain circumstances for research and conservation (e.g. skeletal marking in fish).

**Principle 7:** Antimicrobial agents should only be used in well-defined circumstances for the prevention/prophylaxis and control/metaphylaxis of a specific disease and follow appropriate professional oversight, dose, and duration.

**Principle 8:** Only legally authorized antimicrobial agents should be used and all applicable label directions should be followed; except where specific legal exemptions apply.

**Principle 9:** Foodborne AMR risk management measures should be implemented in a way that is proportionate to the risk and reviewed on a regular basis as described in CAC/GL77. Risk managers should consider potential unintended consequences to human and animal health of recommended risk management measures.

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6 [Could include ethnoveterinary and other alternative approaches, e.g. herbal medicine, probiotics, prebiotics, competitive exclusion, bacteriophages, immunomodulators, organic acids and teat sealants.]
Principle 10: Monitoring and surveillance of the use of antimicrobial agents and the incidence or prevalence, and in particular trends, of foodborne antimicrobial resistant microorganisms and resistance determinants, with specific consideration being given to temporal trends, are among the critical factors to consider when developing risk management measures and evaluating the effectiveness of implemented risk management measures. Use of medically important antimicrobial drugs in humans, animals and crops, and transmission of pathogens and resistance genes between humans, animals, and the environment are additional factors to consider.

Principle 11: This document is designed to provide a framework, for the development of measures to mitigate the risk of foodborne AMR, that countries may implement, as part of their national strategy on AMR, in accordance with their capabilities, based on their national situation/capacities, and within a reasonable period of time. A stepwise approach may be utilized by some countries to properly implement all of the elements in this document.

Principle 12: Medically important antimicrobials should be administered or applied only by veterinarians, plant health professionals or other suitably trained person authorized in accordance with national legislation.

Principle 13: Administration of antimicrobial agents should take into consideration sampling and susceptibility testing of isolates from the production setting, where appropriate, and make adjustments to the administration when problems become evident.

Principle 14: Administration of antimicrobial agents should be based on sound clinical judgement and where feasible on the results of integrated resistance surveillance and monitoring (bacterial cultures and antimicrobial susceptibility testing), as well as relevant experience.

Principle 15: The reduce, replace and rethink (RRR) strategy should be actively promoted within all sectors.

Principle 16: On a continuous and stepwise implementation of risk management measures along the food chain to minimize the possible risks associated with foodborne AMR, priority should be given to the most relevant elements as from a public health perspective.

Principle 17 (new): Animal health and welfare, and the environment should also be taken into account when considering risk management measures. Efforts should be made to ensure that environmental reservoirs of antimicrobial agents, antimicrobial resistant organisms and resistance determinants are kept to a minimum.

Principle 18 (new): Responsible and prudent preventive/prophylactic use of antimicrobials should be limited to exceptional cases, only when the risk of bacterial disease is high and consequences are severe and should be based on veterinarian oversight (or by other suitably trained and authorised person in accordance with national legislation). This use should not be systematic, nor routine, nor applied to compensate for poor hygiene or inadequate animal husbandry/plant production practices, and it should be prescribed only for a limited duration to cover the period of risk. It should always be based on epidemiological and clinical knowledge, with documented justification. When considering preventive use in populations; it should be focused on subsets at highest risk. Preventive use of antibiotics should be limited to individual animals only. Preventive use should always represent a very small proportion of total infectious disease-related use.

Principle 19 (new): Responsible and prudent control/metaphylactic use should not be systematic, nor routine, nor applied to compensate for poor hygiene or inadequate animal husbandry/plant production practices. The decision to administer antimicrobials metaphylactically should be based on a diagnosis and prescribed by or on the order of a veterinarian or other suitably trained person authorised in accordance with national legislation, with documented justification. It should be based on epidemiological and clinical knowledge, an understanding of risk factors associated with the group, and in accordance with pre-established criteria (where available) for initiation of administration of antimicrobials.
Responsibilities of the Regulatory Authorities

9. The national regulatory authorities, which are responsible for granting the marketing authorisation for antimicrobials for use in food-producing animals [the food chain], have a significant role in specifying the terms of this authorisation and in providing the appropriate information to the veterinarian [and plant health professionals] through product labelling and/or by other means, in support of [the responsible and] prudent use of veterinary antimicrobial drugs[agents] in food-producing animals [the food chain]. It is the responsibility of regulatory authorities to develop up-to-date guidelines on data requirements for evaluation of veterinary antimicrobial drug[agent] applications. National governments in cooperation with animal [plant,] and public health professionals should adopt a proactive [One Health] approach to promote [the responsible and] prudent use of antimicrobials in food-producing animals [the food chain] as an element of a national strategy for the containment of antimicrobial resistance. Other elements of the national strategy should include good animal husbandry [and plant production] practices, vaccination [and biosecurity] policies and development of animal [and plant] health care at the farm level, all of which should contribute to reduce the prevalence of animal [and plant] disease requiring antimicrobial treatment. Use of veterinary antimicrobial drugs for growth promotion that belong to classes of antimicrobial agents used (or submitted for approval) in humans and animals should be terminated or phased out in the absence of risk-analysis, as described in Paragraph 8.

10. It is the responsibility of the pharmaceutical company or sponsor\(^2\) to submit the data requested by the regulatory authorities for granting marketing authorisation.

11. The use of antimicrobial agents in food-producing animals requires a marketing authorisation, granted by the competent authorities when the criteria of safety, quality and efficacy are met.

- The examination of dossiers/drug applications should include an assessment of the risks to both animals and humans resulting from the use of antimicrobial agents in food-producing animals. The evaluation should focus on each individual veterinary antimicrobial drug but take into consideration the class of antimicrobials to which the particular active principle belongs.

- The safety evaluation should include consideration of the potential impact of the proposed use in food-producing animals on human health, including the human health impact of antimicrobial resistance developing in microorganisms found in food-producing animals and their environment associated with the use of veterinary antimicrobial drugs.

12. [\(10.\).] If dose ranges or different durations of treatment are indicated, the national regulatory authorities should give guidance on the approved product labelling regarding the conditions that will minimize the development of resistance in target pathogens, and minimize the exposure to the antimicrobial of other microorganisms that might be of public health relevance, when this information is available.

11. For more information on antimicrobial drugs for food-producing animals see the OIE Aquatic Animal Health Code Chapter 6.2.4 - Responsibilities of competent authorities and the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 1. Marketing authorization, and other Sections.]

13. The relevant authorities should make sure that all the antimicrobial agents used in food-producing animals are prescribed by a veterinarian or other suitably trained person authorized in accordance with national legislation or used under conditions stipulated in the national legislation. (See OIE Guidelines for Antimicrobial Resistance: Responsible and Prudent Use of Antimicrobial Agents in Veterinary Medicine (Terrestrial Animal Health Code, Appendix 3.9.3)

14. No veterinary antimicrobial drug should be administered to animals unless it has been evaluated and authorized for such use by the relevant authorities or the use is allowed through off-label guidance or legislation. Regulatory authorities should, where possible, expedite the market approval process of new veterinary antimicrobial drug formulations considered to have the potential to make an important contribution in the control of antimicrobial resistance.

15. Countries without the necessary resources to implement an efficient authorisation procedure for veterinary antimicrobial drugs and whose supply of veterinary antimicrobial drugs mostly depends on imports from foreign countries should:

- ensure the efficacy of their administrative controls on the import of these veterinary antimicrobial drugs,
- seek information on authorizations valid in other countries, and

develop the necessary technical cooperation with experienced authorities to check the quality of imported veterinary antimicrobial drugs as well as the validity of the recommended conditions of use. Alternatively, a national authority could delegate a competent institution to provide quality certification of veterinary antimicrobial drugs.

16. All countries should make every effort to actively combat the manufacture, advertisement, trade, distribution and use of illegal and/or counterfeit bulk active pharmaceutical ingredients and products. Regulatory authorities of importing countries could request the pharmaceutical industry to provide quality certificates or, where feasible, certificates of Good Manufacturing Practices prepared by the exporting country's national regulatory authority.

QUALITY CONTROL OF ANTIMICROBIAL AGENTS

17. Regulatory authorities should ensure that quality controls are carried out in accordance with international guidance and in compliance with the provisions of good manufacturing practices, in particular:

- to ensure that the quality and concentration (stability) of veterinary antimicrobial drugs in the marketed dosage form(s) is maintained and properly stored up to the expiry date, established under the recommended storage conditions.
- to ensure the stability of veterinary antimicrobial drugs when they are mixed with feed or drinking water.
- to ensure that all veterinary antimicrobial drugs are manufactured to the appropriate quality and purity.

[13. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 2. Quality control of antimicrobial agents and VMP containing medically important antimicrobial agents. Relevant Chapters of the OIE Aquatic Animal Health should be quoted.

ASSESSMENT OF EFFICACY

[14. Assessment of efficacy is important to assure adequate response to the administration of antimicrobials.]

[15. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 3. Assessment of therapeutic efficacy. Relevant Chapters of the OIE Aquatic Animal Health should be quoted.

18. Preclinical data should be generated to establish an appropriate dosage regimen necessary to ensure the efficacy of the veterinary antimicrobial drug and limit the selection of microbial-resistant microorganisms. Such preclinical trials should, where applicable, include pharmacokinetic and pharmacodynamic studies to guide the development of the most appropriate dosage regimen.

19. Important pharmacodynamic information may include:

- mode of action;
- the spectrum of antimicrobial activity of the substance;
- identification of bacterial species that are naturally resistant relevant to the use of the veterinary antimicrobial drugs;
- antimicrobial minimum inhibitory and/or bactericidal concentrations;
- determination of whether the antimicrobial exhibits time or concentration-dependent activity or co-dependency;
- evaluation of activity at the site of infection.

20. Important pharmacokinetic information may include:

- bioavailability according to the route of administration;
- concentration of the veterinary antimicrobial drug at the site of infection and its distribution in the treated animal;
- metabolism which may lead to the inactivation of veterinary antimicrobial drugs;
- excretion routes.

21. The use of fixed combinations of veterinary antimicrobial drugs should be justified taking into account:
pharmacodynamic (additive or synergistic effects towards the target microorganism);
pharmacokinetics (maintenance of the concentrations of associated antimicrobials responsible for additive or synergistic effects at the site of infection throughout the treatment period).

22. Clinical data should be generated to confirm the validity of the claimed indications and dosage regimens established during the preclinical phase.

23. Criteria to be considered include:
- parameters for qualitatively and quantitatively assessing efficacy;
- diversity of the clinical cases met when carrying out clinical trials;
- compliance of the protocols of clinical trials with good clinical practice, such as VICH guidelines;
- eligibility of the studied clinical cases based on appropriate clinical and microbiological criteria.

ASSESSMENT OF THE POTENTIAL OF VETERINARY ANTIMICROBIAL DRUGS[AGENTS] TO SELECT FOR RESISTANT MICROORGANISMS

[16. The competent authority should assess the potential of medically important antimicrobial drugs to select for resistant microorganisms taking into account CAC/GL77.]

[17. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 4. Assessment of the potential of antimicrobial agents to select for resistance and Chapter 6.10 Risk Analysis for Antimicrobial Resistance Arising from the Use of Antimicrobial Agents in Animals.] Relevant Chapters of the OIE Aquatic Animal Health should be quoted.

24. Where applicable, data from preclinical or clinical trials should be used to evaluate the potential for target microorganisms, foodborne and/or commensal microorganisms to develop or acquire resistance.

25. Appropriate information should be provided to support an adequate assessment of the safety of veterinary antimicrobial drugs being considered for authorisation in food-producing animals. The regulatory authorities should develop criteria for conducting such assessments and interpreting their results. Existing guidelines for antimicrobial resistance risk assessment, such as the OIE Guideline may be used for more comprehensive information. The type of information to be evaluated in these assessments may include, but is not limited to, the following:
- the route and level of human exposure to food-borne or other resistant microorganisms;
- the degree of cross resistance within the class of antimicrobials and between classes of antimicrobials;
- the pre-existing level of resistance, if available, in pathogens causing gastrointestinal infections in humans (baseline determination);
- the concentration of active compound in the gut of the animal at the defined dosage level.

ESTABLISHMENT OF ADIS (ACCEPTABLE DAILY INTAKE), MRLS (MAXIMUM RESIDUE LIMIT), AND WITHDRAWAL PERIODS FOR VETERINARY ANTIMICROBIAL DRUGS

26. When setting ADIs and MRLs for veterinary antimicrobial drugs, the safety evaluation is carried out in accordance with international guidelines and should include the determination of microbiological effects (e.g., the potential biological effects on the human intestinal flora) as well as toxicological and pharmacological effects.

27. An acceptable daily intake (ADI) and a maximum residue limit (MRL) for appropriate food stuffs (i.e., meat, milk, eggs, fish and honey) should be established for each antimicrobial agent. MRLs are necessary in order that officially recognised control laboratories can monitor that the veterinary antimicrobial drugs are being used as approved. Withdrawal periods should be established for each veterinary antimicrobial drug, which make it possible to produce food in compliance with the MRLs.

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28. Withdrawal periods have to be established for each veterinary antimicrobial drug by taking into account:

- the MRLs established for the considered veterinary antimicrobial drug;
- the pharmaceutical form;
- the target animal species;
- the dosage regimen and the duration of treatment;
- the route of administration.

[ASSESSMENT OF ENVIRONMENTAL IMPACT]

[18. Regulatory authorities should assess the impact of proposed veterinary antimicrobial drug use on the environment in accordance with national guidelines or recognized international guidelines.]

[19. For more information on antimicrobial drugs for food-producing animals see the International Cooperation on Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products guidelines.]

[20. Regulatory authorities should consider addressing the environmental aspects of AMR (e.g. pollution from pharmaceutical manufacture, impacts of reusing waste water for irrigation and using manure for soil fertilization, harmonized monitoring and establishment of maximum admissible levels, etc.)]

ESTABLISHMENT OF A SUMMARY OF PRODUCT CHARACTERISTICS FOR EACH VETERINARY ANTIMICROBIAL DRUG FOR FOOD-PRODUCING ANIMALS

[21. Regulatory authorities should establish a Summary of Product Characteristics that can be utilized in labelling and as a package insert.]

[22. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 7. Establishment of a summary of product characteristics for each VMP containing antimicrobial agents.] Relevant Chapters of the OIE Aquatic Animal Health should be quoted.

29. The summary of product characteristics contains the information necessary for the appropriate use of veterinary antimicrobial drugs. It constitutes, for each veterinary antimicrobial drug, the official reference of the content of its labelling and package insert. This summary contains the following items:

- pharmacological properties;
- target animal species;
- indications;
- target microorganisms;
- dosage and administration route;
- withdrawal periods;
- incompatibilities;
- shelf-life;
- operator safety;
- particular precautions before use;
- instructions for the return or proper disposal of unused or out-of-date products;
- any information on conditions of use relevant to the potential for selection of resistance should be included, for the purpose of guidance on prudent use;
- class and active ingredient of the veterinary antimicrobial drug.

SURVEILLANCE AND MONITORING PROGRAMMES

[23. Regulatory authorities should establish systems for the surveillance and monitoring of antimicrobial resistance and antimicrobial use following the Codex Proposed Draft Guidelines on Integrated Surveillance (CAC/GL xx-xxxx), taking into consideration relevant sections of Guidelines for Foodborne Antimicrobial Resistance CAC/GL 77-2011; WHO guidelines on Integrated surveillance of antimicrobial resistance in foodborne bacteria, Application of a One Health Approach (2017); and OIE Terrestrial Animal Health Code Chapter 6.7 Harmonisation of national antimicrobial resistance surveillance and monitoring programmes and Chapter 6.8 Monitoring of the quantities and usage patterns of antimicrobial agents used in food-producing animals.] Relevant Chapters of the OIE Aquatic Animal Health should be quoted.

[24. The surveillance and monitoring of antimicrobial resistant microorganisms in different production sectors and in different products is necessary for understanding the development and dissemination of antimicrobial resistance, providing relevant risk assessment data, and assessing the effectiveness of interventions. Surveillance and monitoring programmes involve specific and continuous data collection, analysis and reporting that quantitatively monitor temporal trends in the occurrence and/or prevalence and distribution of resistance to antimicrobial agents; it also allows the identification of emerging or specific patterns of multiple resistance.]

30. The relevant authorities should develop a structured approach to the investigation and reporting of the incidence and prevalence of antimicrobial resistance. For the purposes of this Code, priority should be given to the evaluation of antimicrobial resistance in foodborne microorganisms.

For reasons of efficiency, the methods used to establish such programmes (laboratory techniques, sampling, choice of veterinary antimicrobial drug(s) and microorganism(s)) should be harmonized as much as possible at the international level (e.g. OIE documents on “Harmonisation of National Antimicrobial Resistance Monitoring and Surveillance Programmes in Animals and Animal Derived Food” http://www.oie.int/eng/publicat/r/2003/a_r20318.htm and “Standardisation and Harmonisation of Laboratory Methodologies Used for the Detection and Quantification of Antimicrobial Resistance” http://www.oie.int/eng/publicat/r/2003/a_r20317.htm).

31. Preferably, epidemiological surveillance of antimicrobial resistance should be accompanied by data on the amounts of veterinary antimicrobial drugs used by veterinarians and other authorized users in food-producing animals. These data could be collected using one or more of the following sources:

- production data from manufacturers;
- importers and exporters;
- if possible, data on intended and actual usage from manufacturers, wholesale and retail distributors including feed mills, and veterinary prescription records;
- surveys of veterinarians, farmers and producers of food-producing animals.

32. Regulatory authorities should have in place a pharmacovigilance programme for the monitoring and reporting of adverse reactions to veterinary antimicrobial drugs, including lack of the expected efficacy related to microbial resistance. The information collected through the pharmacovigilance programme should form part of the comprehensive strategy to minimize microbial resistance.

33. In cases, where the assessment of data collected from pharmacovigilance and from other post-authorization surveillance including, if available, targeted surveillance of antimicrobial resistance, suggests that the conditions of use of the given veterinary antimicrobial drug should be reviewed, regulatory authorities shall endeavour to achieve this re-evaluation.

DISTRIBUTION OF VETERINARY ANTIMICROBIAL DRUGS/AGENTS IN VETERINARY MEDICINE

[25. Regulatory authorities should make sure antimicrobial agents are distributed through appropriate distribution systems in accordance with national legislation and medically important antimicrobials are distributed to appropriately credentialed veterinarians, plant health professionals, or other suitably trained person authorized in accordance with national legislation.]

[26. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 9. Supply and administration of the VMP containing antimicrobial agents.] Relevant Chapters of the OIE Aquatic Animal Health should be mentioned.

34. The relevant authorities should make sure that all veterinary antimicrobial drugs used in food-producing animals are, to the extent possible:
prescribed by a veterinarian or other suitably trained person authorized in accordance with national legislation or used under conditions stipulated in the national legislation;

 supplied only through licensed/authorized distribution systems;

 administered to animals by a veterinarian or, under the supervision of a veterinarian or other suitably trained person authorized in accordance with national legislation; and that

 proper records are kept of their administration (see Paragraph 58, Responsibilities of Veterinarians: Recording section).

 [27. Distribution and use should be regularly controlled by the competent authorities, and targeted checks should be carried out, where appropriate, on prescribers with high levels or concerning patterns of prescriptions.]

 CONTROL OF ADVERTISING

 [28. Regulatory authorities should assure that advertising of antimicrobial agents is done in accordance to national legislation.]

 [29. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 10. Control of advertising.] Relevant Chapters of the OIE Aquatic Animal Health should be mentioned.

 35. Advertising of veterinary antimicrobial drugs[agents] should be done in a manner consistent with prudent use guidelines and any other specific regulatory recommendation for the product.

 All advertising of veterinary antimicrobial drugs[agents] should be controlled by the relevant authorities.

 - The authorities should ensure that advertising of veterinary antimicrobial drugs[agents]:
   - complies with the marketing authorisation granted, in particular with the content of the summary of product characteristics; and
   - complies with each country’s national legislation.

 TRAINING OF USERS OF VETERINARY ANTIMICROBIAL DRUGS[AGENTS]

 36. Training should be undertaken to assure the safety to the consumer of animal derived food and therefore the protection of public health. Training should involve all the relevant professional organisations, regulatory authorities, the pharmaceutical industry[marketing authorization holders], veterinary schools, research institutes, professional associations[ trade associations] and other approved users such as farmers and producers of food animals and should focus on:

 - information on disease prevention and management strategies to reduce the need to use veterinary antimicrobial drugs[agents];
 - relevant pharmacokinetic and pharmacodynamic information to enable the veterinarian [and plant health professionals] to use veterinary antimicrobial drugs[agents] responsibly and prudently;
 - the ability of veterinary antimicrobial drugs[agents] to select for resistant microorganisms in food-producing animals that may contribute to animal[, plant], human health and environmental problems; and
 - the need to observe responsible and prudent use recommendations and using veterinary antimicrobial drugs[agents] in animal husbandry[production settings] in agreement with the provisions of the marketing authorisations and veterinary[professional] advice.

 [32. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.3 - Responsible and prudent use of antimicrobial agents in veterinary medicine section 11. Training on the usage of antimicrobial agents.] Relevant Chapters of the OIE Aquatic Animal Health should be mentioned.

 DEVELOPMENT OF RESEARCH

 37. The relevant authorities should encourage public and private research to:

 - improve the knowledge about the mechanisms of action of antimicrobials in order to optimise the dosage regimens and their efficacy;
• improve the knowledge about the mechanisms of selection, emergence and dissemination of resistance determinants;
• develop practical models for applying the concept of risk analysis to assess the public health concern precipitated by the development of resistance;
• further develop protocols to predict, during the authorisation process, the impact of the proposed use of the veterinary antimicrobial drugs[agents] on the rate and extent of resistance development; and
• develop and encourage [good animal husbandry and plant production practices and] alternative methods to prevent [and treat] infectious diseases [that would reduce the need to use antimicrobials]
• [develop alternatives to antimicrobials, new antimicrobials, rapid diagnostics, and vaccines, including autogenous vaccines.]
• [determine the potential transfer to fresh produce and other crops of resistant microorganisms and determinants from animal manures used as fertilizer.]

[34. Research should be conducted, as resources permit, on antimicrobials, their metabolites, resistant microorganisms and resistance determinants in the environment, and if feasible, factors affecting and the magnitude of resistance determinant transfer among microorganisms in the environment.]

COLLECTION AND DESTRUCTION OF UNUSED OR [OUT-OF-DATE] VETERINARY ANTIMICROBIAL DRUGS[AGENTS]

38-35. The relevant authorities should develop effective [and compulsory] procedures for the safe collection and destruction of unused or out-of-date veterinary antimicrobial drugs[agents].

Responsibilities of the Veterinary Pharmaceutical Industry [Manufacturers]

MARKETING AUTHORISATION OF VETERINARY ANTIMICROBIAL DRUGS [AGENTS] FOR FOOD-PRODUCING ANIMALS

39-36. It is the responsibility of the veterinary pharmaceutical industry—antimicrobial agent marketing authorization holders:
• to supply all of the information requested by the national regulatory authority in order to establish objectively the quality, safety and efficacy of veterinary antimicrobial drugs[agents]; and
• to ensure the quality of this information on the basis of the implementation of procedures, tests and trials in compliance with the provisions of good manufacturing, good laboratory and good clinical practices.

MARKETING AND EXPORT OF VETERINARY ANTIMICROBIAL DRUGS [AGENTS]

40-37. Only officially licensed/authorized veterinary antimicrobial drugs[agents] should be marketed, and then only through approved distribution systems.
• Only veterinary antimicrobial drugs[agents] meeting the quality standards of the importing country should be exported from a country in which the products were produced;
• The information necessary to evaluate the amount of veterinary antimicrobial drugs[agents] marketed should be provided to the national regulatory authority.

[38. Package size and the strength of antimicrobial formulations should be adapted as far as possible to the approved indications of use (in order to avoid improper dosing, overuse and leftovers).]

ADVERTISING

41-39. It is the responsibility of the veterinary pharmaceutical industry [marketing authorization holders] to advertise veterinary [medically important] antimicrobial[s] drugs in accordance with the provisions of Paragraphs 35-28-30 on the Responsibilities of the Regulatory Authorities, Control of Advertising and to not inappropriately advertise [medically important] antimicrobials directly to the food animal producer. [Advertising should only be allowed to persons permitted to prescribe or supply antimicrobial drugs. Promotional campaigns involving economic or material benefits for prescribers or suppliers of antimicrobials should be prohibited.]

TRAINING

42-40. It is the responsibility of the veterinary pharmaceutical industry [marketing authorization holders] to participate in the training of users of veterinary antimicrobial drugs[agents] as defined in Paragraph 36.31.
RESEARCH

43. [41.] It is the responsibility of the veterinary pharmaceutical industry [marketing authorization holders] to contribute to the development of research as defined in Paragraph 37 [including research on the development of alternatives to the use of antimicrobials, new antimicrobials, rapid diagnostics and vaccines].

Responsibilities of Wholesale and Retail Distributors

44.[42.] Retailers distributing veterinary [medically important] antimicrobial[s] drugs should only do so on the prescription of a veterinarian, [plant health professional] or other suitably trained person authorized in accordance with national legislation and all products should be appropriately labelled.

45.[43.] Distributors should encourage compliance with the national guidelines on the responsible use of veterinary [medically important] antimicrobial[s] drugs and should keep detailed records of all [medically important] antimicrobials supplied according to the national regulations including:

- date of supply
- name of prescribing veterinarian [plant health professional, or other suitably trained and authorized person in accordance with national legislation]
- name of user
- name of medicinal product
- batch number
- quantity supplied

46.[44.] Distributors should participate in the training of users of veterinary antimicrobial drugs[agents] as defined in Paragraph 36 [30].

Responsibilities of Veterinarians [11] [and Plant Health Professionals]

47.[45.] The veterinarian[ and plant health professionals are] responsible for identifying recurrent disease problems and developing alternative strategies to prevent or treat infectious disease. These may include changes in husbandry conditions and vaccination programs where vaccines are available/biosecurity, improved production practices, and alternatives to antimicrobials including vaccinations where applicable/available.

[46. For more information on antimicrobial drugs for food-producing animals see the OIE Terrestrial Animal Health Code Chapter 6.9.6 – Responsibilities of veterinarians.] Relevant Chapters of the OIE Aquatic Animal Health should be mentioned.

48. Veterinary antimicrobial drugs should only be prescribed for animals under his/her care, which means that:

- the veterinarian has been given responsibility for the health of the animal or herd/flock by the producer or the producer’s agent;
- that responsibility is real and not merely nominal;
- that the animal(s) or herd/flock have been seen immediately before the prescription and supply, or recently enough for the veterinarian to have personal knowledge of the condition of the animal(s) or current health status of the herd or flock to make a diagnosis and prescribe; and
- the veterinarian should maintain clinical records of the animal(s) or the herd/flock.

49.[47.] It is recommended that veterinary professional organizations develop for their members species-specific clinical practice guidelines on the responsible use of veterinary antimicrobial drugs[species or sector-specific guidelines for responsible and prudent use of antimicrobial agents].

[Within the national action plans, which countries are developing under the Global Action Plan, there should be the recommendation to develop sector and species-specific clinical practice guidelines on the responsible use of veterinary antimicrobials agents. These guidelines would be created by the sector specific veterinary professional organizations depending on local AMR circumstances and availability of treatment]

[11] Under some circumstances, this may refer to a suitably trained person authorized in accordance with national legislation.
Veterinary antimicrobial drugs should only be used when necessary and in an appropriate manner:

- A prescription [or order for application] for veterinary [medically important] antimicrobial[s] drugs must precisely indicate the treatment regimen, the dose, the dosage intervals, the duration of the treatment, the withdrawal period[ when appropriate,] and the amount of antimicrobial to be delivered depending on the dosage, the number, and the weight of the animals[ and the characteristics of the individual or population] to be treated;

- [The delivered amount should be limited only for the treatment concerned. Prescriptions should also indicate the animal keeper/owner and the identification of the animal(s) or plants to be treated;]

- All veterinary [medically important] antimicrobial[s] drugs should be prescribed [or applied] and used according to [label directions and] the conditions stipulated in the national legislation.

For food-producing animals, the appropriate use of veterinary [medically important] antimicrobial[s] drugs in practice is a clinical decision which should be based on the experience and local expertise of the prescribing veterinarian, and the accurate diagnosis, based on adequate diagnostic procedures. There will be occasions when a group of animals, which may have been exposed to pathogens, may need to be treated without recourse to an accurate diagnosis and antimicrobial susceptibility testing in order to prevent the development of clinical disease and for reasons of animal welfare.

Determination of the choice of a veterinary antimicrobial [agents] drug by:

- The expected efficacy of the treatment based on:
  - the clinical experience of the veterinarian [, plant health professional or suitably trained and authorized person in accordance with national legislation];
  - the spectrum of the antimicrobial activity towards the pathogens involved;
  - the epidemiological history of the rearing[production] unit particularly in regards to the antimicrobial resistance profiles of the pathogens involved. Ideally[Whenever possible], the antimicrobial profiles should be established before the commencement of treatment. [If this is not possible, samples should nevertheless be taken before start of the treatment to allow, if necessary, for adjustment of therapy based on sensitivity testing.] Should a first antimicrobial treatment fail or should the disease recur, the use of a second veterinary antimicrobial drug[agent] should be based on the results of microbiological/susceptibility tests;
  - the appropriate route of administration;
  - results of initial treatment;
  - [ previous published scientific information on the treatment of the specific disease:]
    - known pharmacokinetics/tissue distribution to ensure that the selected veterinary antimicrobial drug is active at the site of infection;
    - prognosis[the likely course of the disease].

- The need to minimize the adverse health impact from the development of antimicrobial resistance based on:
  - the choice of the activity spectrum of the veterinary antimicrobial drug[agent] [(narrow-spectrum antimicrobials should be preferred whenever possible/appropriate)];
  - the targeting of specific microorganism;
  - known or predictable susceptibilities using antimicrobial susceptibility testing;
  - optimized dosing regimens;
  - the use of effective combinations of veterinary antimicrobial drugs[agents];
  - the importance of the antimicrobial drugs to veterinary and human medicine; and,
  - the route of administration.

If the label conditions allow for some flexibility, the veterinarian should consider a dosage regimen that is long enough to allow an effective recovery of the animal but is short enough to limit the selection of resistance in foodborne and/or commensal microorganisms.
OFF-LABEL USE

54.[51.] [For food-producing animals, the] The off-label use of a veterinary antimicrobial drug may be permitted in appropriate [(exceptional)] circumstances and should be in agreement with the national legislation in force including the administrative withdrawal periods to be used. It is the veterinarian’s responsibility to define the conditions of responsible use in such a case including the therapeutic regimen, the route of administration, the duration of the treatment and the withdrawal period. Off-label use of [medically important] antimicrobial for growth promotion purposes should not be permitted.

[Human health risk related to foodborne antimicrobial resistance should be an important factor when considering the off-label use of veterinary antimicrobial agents.]

RECORDING

55.[52.] [For food-producing animals, records] Records on veterinary antimicrobial drugs should be kept in conformity with national legislation. Veterinarians may refer to recording information as covered in the relevant national legislation.

In particular, for investigation of antimicrobial resistance, veterinarians should:

- record the antimicrobial susceptibility testing results;
- record the antimicrobial used, the dosage regimen and the duration;
- investigate adverse reactions to veterinary antimicrobial drugs, including lack of expected efficacy due to antimicrobial resistance, and report it, as appropriate, to the regulatory authorities [(through a pharmacovigilance system)].

56.[53.] [For food-producing animals, veterinarians] Veterinarians should also periodically review farm records on the use of veterinary antimicrobial drugs to ensure compliance with their directions.

TRAINING

57.[54.] Veterinary professional organizations should participate in the training of users of veterinary antimicrobial drugs as defined in Paragraph 36.[31].

Responsibilities of [Food] Producers

58.[55.] Producers are responsible for preventing disease outbreaks and implementing health and welfare programmes on their farms. They may, as appropriate, call on the assistance of their veterinarian[, plant health professional, or other suitably trained person authorized in accordance with national legislation. All people involved with food-producing animals[the food chain] have an important part to play in [preventing disease and] ensuring the responsible [and prudent] use of veterinary antimicrobial drugs[agents].

59.[56.] Producers of food-producing animals have the following responsibilities:

- to use veterinary antimicrobial drugs[agents] only when necessary and not as a replacement for good management and farm hygiene, or other disease prevention methods such as vaccination;
- to implement a health plan in cooperation with the veterinarian [, plant health professional, or other suitably trained person authorized in accordance with national legislation] in charge of the animals that outlines preventative measures (e.g. mastitis plan, worming and vaccination programmes, etc.);
- to use veterinary antimicrobial drugs[agents] in the species, for the uses and at the doses on the approved labels and in accordance with the prescription, product label instructions or the advice of a veterinarian [, plant health professional or other suitably trained person authorized in accordance with national legislation] familiar with the animals and[or] the production site;
- to isolate sick animals and dispose of dead or dying animals promptly under conditions approved by relevant authorities;
- to comply with the storage conditions of veterinary antimicrobial drugs[agents] according to the approved product labelling;
- to address hygienic conditions regarding contacts between people (veterinarians, [plant health professionals,] breeders, owners, children) and the animals[populations] treated;

Veterinarians can also refer to the “Recommended International Code of Practice for Control of the Use of Veterinary Drugs CAC/RCP 38-1993.”
• to comply with the recommended withdrawal periods to ensure that residue levels in animal derived food do not present a risk for the consumer;

• to not use out-of-date veterinary antimicrobial drugs[agents] and to dispose of all unused veterinary antimicrobial drugs[agents] in accordance with the provisions on the product labels [and national legislation];

• to inform the veterinarian, plant health professional, or other suitably trained person authorized in accordance with national legislation in charge of the [production] unit of recurrent disease problems;

• to maintain all clinical and laboratory records of microbiological and susceptibility tests if required by the national regulatory authority. These data should be made available to the veterinarian[professional] in charge of treating the animals[treatment] in order to optimize the use of veterinary antimicrobial drugs[agents].

• to keep adequate records of all veterinary antimicrobial drugs[agents] used, including the following:
  o name of the veterinary antimicrobial drug[agent]/active substance and batch number;
  o name of supplier;
  o date of administration;
  o identification of the animal or group of animals[production unit] to which the veterinary antimicrobial drug[agent] was administered;
  o clinical conditions[disease] treated;
  o quantity and duration of the antimicrobial agent administered;
  o withdrawal periods;
  o result of laboratory tests;
  o result of treatment;
  o name of the prescribing veterinarian [, plant health professional] or other suitably trained person authorized in accordance with national legislation.

• To ensure sound management of animal wastes and other materials to avoid [minimize] dissemination of antimicrobial agents and resistance determinants into the environment;

• To prevent the unnecessary contact with and transmission of resistant bacteria to all personnel, including farm workers;

• To assist the relevant authorities in surveillance programs related to antimicrobial resistance.

[57. The responsible and prudent use of antimicrobials must be supported by continuous efforts in disease prevention to minimise infection during production and decrease the volume of antibiotics used. Efforts should aim to improve health, thereby reducing the need for antibiotics. This can be achieved by improving hygiene, biosecurity and health management on farms, and implementing national or international good animal husbandry, aquaculture, or agricultural practices. Disease prevention through the use of vaccines and other measures such as probiotics (beneficial bacteria found in various foods), prebiotics (non-digestible foods that help probiotic bacteria grow and flourish) or competitive exclusion products (intestinal bacterial flora that limit the colonisation of some bacterial pathogens) should be considered and applied wherever appropriate and available.]

[58. Concerted efforts of all stakeholders within the entire food chain is required to minimize and contain foodborne antimicrobial resistance. While such efforts mainly focus on responsible and prudent use of antimicrobial agents in primary production at the farm level, the later phase of the food chain also plays a significant role in preventing transmission and spread of resistant bacteria and resistance determinants.

Food processing industry, food retailers and consumers should take necessary action in accordance with the Principles and Guidelines for the Conduct of Microbiological Risk Management (CAC/GL 63-2007).

Food retailers should favor food produced in accordance with quality schemes and systems of production and supply that apply principles of prudent use i.e. that minimize the use of antimicrobials (and, in case of food producing animals, promote high standards of animal health and welfare).]
**Responsibilities of Consumers**

[59. Consumers have an important role to play to minimize and control antimicrobial resistance. By practicing safe food handling techniques, following health recommendations, and maintaining awareness of antimicrobial resistance information, consumers can minimize the risk of contracting and spreading infectious bacteria thereby further reducing the need for antibacterials. Consumer should:

- Take antibiotics only when needed in accordance with medical prescription;
- Use a food thermometer to ensure that foods are cooked to a safe internal temperature: 145°F (63°C) for whole beef, pork, lamb, and veal (allowing the meat to rest for 3 minutes before carving or consuming), 160°F (71°C) for ground meats, and 165°F (74°C) for all poultry, including ground chicken and ground turkey;
- Wash hands after touching raw meat, poultry, and seafood. Also wash work surfaces, cutting boards, utensils, and grill before and after cooking;
- Keep food below 40°F and refrigerating foods within 2 hours of cooking (1 hour during the summer heat);
- Separate raw meat, poultry, seafood and eggs from fresh produce and ready-to-eat foods to avoid cross contamination. Use different cutting boards to prepare raw meat or poultry and any food that will be eaten without cooking;
- Wash hands after contact with feces, animals or animal environments;
- Report suspected outbreaks of illness from food to local health department; and
- Review Competent Authority’s traveler’s health recommendations when preparing to travel to a foreign country.

**Responsibilities of feed business operators**

Reference should be made to chapter 6.9 Article 6.9.8 of the OIE Animal Health Code

**Advocacy and Communication**

[60. The successful control of antimicrobial resistance along the food chain requires the involvement and cooperation of all parties along the food chain. These include the relevant authorities and stakeholders such as the manufacturers, veterinarians and plant health professionals, wholesale and retail distributors, producers, and consumers who are involved in the authorisation, production, control, importation, exportation, distribution and use of antimicrobial agents.]

[61. Advocacy and communication strategies should identify relevant target audiences, such as policy makers, health, veterinary and agricultural professionals, farmers, players in the food industry, the media and the general public, who all have a responsibility in minimising antimicrobial resistance along the food chain.]

[62. Advocacy and communication efforts at the international and national levels should aim to raise awareness of the importance of antimicrobials in treating bacterial infections and the public health challenges of antimicrobial resistance, including within a food safety perspective.]

[63. Advocacy campaigns should be tailored to the specific stakeholder groups. Campaigns targeted at the agriculture and aquaculture sectors should include good animal husbandry, aquaculture or agricultural practices and the responsible and prudent use of antimicrobials. Those at the food industries should reinforce prevention of contamination and food hygiene practices. National guidelines and education programmes should promote best practices, including correct treatment, measures to prevent and reduce the transmission of pathogens, infection control and hygiene measures. Campaigns may also be targeted at consumers to encourage them to demand food that is produced in accordance with standards which require responsible use of antimicrobials i.e. where the amounts used are kept as low as possible (and, in case of food producing animals, where high standards of animal health and welfare are promoted).]

[64. The engagement and consultation of stakeholders prior to enforcement or introduction of prudent use policies or measures are critical for successful implementation. Regulatory authorities should engage all relevant stakeholder groups.]
[65. Establishment of an Adhoc scientific AMR Newsletter with the objective of collection of recent advances in AMR specially in the field of the tripartite organizations (FAO, WHO and OIE), Codex Alimentarius, universities and institutions; with relevance to the development and transmission of food-borne antimicrobial resistance in the food chain (with special emphasis on the genera of Enterobacteriaceae (STEC, Salmonella, Shigella, Campylobacter and Vibrio).

Conclusions
60. Veterinary antimicrobial drugs are very important tools for controlling a great number of infectious diseases in both animals and humans. It is vital that all countries put in place the appropriate systems to ensure that veterinary antimicrobial drugs are manufactured, marketed, distributed, prescribed and used responsibly, and that these systems are adequately audited.

61. This document is designed to provide the framework that countries may implement in accordance with their capabilities but within a reasonable period of time. A stepwise approach may be appropriate for a number of countries to properly implement all of the elements in this document.

62. The continued availability of veterinary antimicrobial drugs, which are essential for animal welfare and animal health and consequently human health, will ultimately depend on the responsible use of these products by all those involved in the authorisation, production, control, distribution and use of antimicrobials in food-producing animals.

End Notes

List of Abbreviations [Acronyms] Used in this Code
ADI—Acceptable Daily Intake
CAC—Codex Alimentarius Commission
CAC/RCP—Codex Alimentarius Commission/Recommended Code of Practice
CCRVDF—Codex Committee on Residues of Veterinary Drugs in Foods
FAO—Food and Agriculture Organization of the United Nations
MRL—Maximum Residue Limit
OIE—Office International des epizooties/International Office of Epizooties
VICH—International Cooperation on Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products
WHO—World Health Organization

Glossary of Definitions and Terms
Veterinary Antimicrobial Drug
Veterinary antimicrobial drug(s) refers to naturally occurring, semi-synthetic or synthetic substances that exhibit antimicrobial activity (kill or inhibit the growth of microorganisms). Where anticoccidial products have antibacterial activity, they should be considered as veterinary antimicrobial drugs, except where this is precluded by national legislation.

Disease Treatment/Therapeutic Use
Treatment/Therapeutic Use refers to use of an antimicrobial(s) for the specific purpose of treating an animal(s) with a clinically diagnosed infectious disease or illness.

Disease Prevention/Prophylactic Use
Prevention/Prophylactic Use refers to use of an antimicrobial(s) in healthy animals considered to be at risk of infection or prior to the onset of clinical infectious disease. This treatment includes:
• Control of the dissemination of a clinically diagnosed infectious disease identified within a group of animals, and
prevention of an infectious disease that has not yet been clinically diagnosed.

**Growth Promotion**

Growth Promotion refers to the use of antimicrobial substances to increase the rate of weight gain and/or the efficiency of feed utilization in animals by other than purely nutritional means. The term does NOT apply to the use of antimicrobials for the specific purpose of treating, controlling, or preventing infectious diseases, even when an incidental growth response may be obtained.