

Codex Alimentarius Efforts on Antimicrobial Resistance

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Codex Antimicrobial Resistance

- Process for Development
- Previous Codex Work
- Current Work

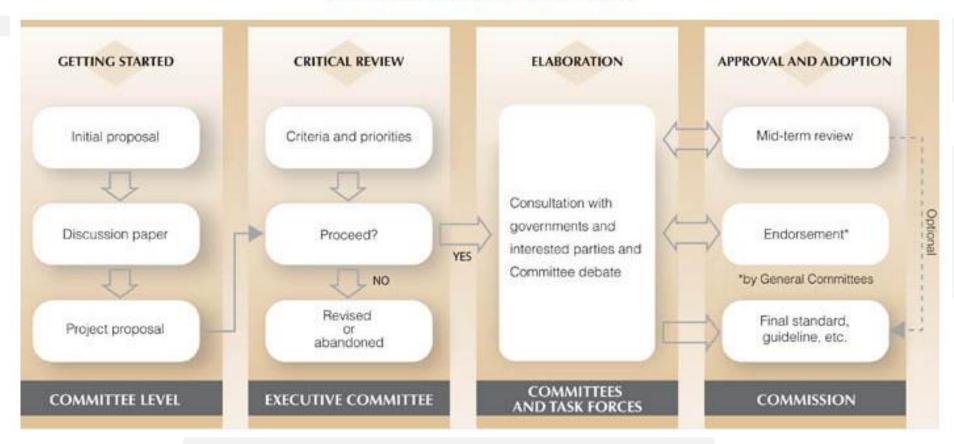


Codex Process

- Lengthy process to develop code
- Standing committees
- Working Groups
 - Physical
 - Electronic
- Task Forces
 - 3 to 4 year life



THE CODEX STANDARDS PROCESS





Codex Process – Steps Take Time

The Codex step procedure

Before a decision is made to undertake the development of a new standard or other text, a project proposal is prepared and discussed at Committee level.

STEP 1

The project proposal is reviewed by the Executive Committee and compared against the criteria and priorities established by the Commission.

STEPS 2, 3 AND 4

A draft text is prepared (Step 2) and circulated to member countries and all interested parties for comment (Step 3). The draft and the comments are reviewed at Committee level (Step 4) and, if necessary, a new draft is prepared.

STEP 5

The Commission reviews the progress made and agrees that the draft should go to finalization. After this stage, the draft is also endorsed by the relevant General Subject Committees so that it is consistent with Codex general standards.*

STEPS 6 AND 7

The approved draft is sent again to governments and interested parties for comment and finalized by the relevant Committee. The draft is submitted to the Commission for adoption.

STEP 8

Following a final round of comments, the Commission adopts the draft as a formal Codex text. The standard, guideline or other text is then published by the Codex Secretariat.

* Sometimes the text is considered to be ready for final adoption at this stage - often called Step 5/8.

Codex Prior AMR Work

- Committee on Residues of Veterinary Drugs in Food
- Code of Practice to Minimize and Contain Antimicrobial Resistance

CAC/RCP 61-2006	Page 1 of 18
CODE OF PRACTICE TO MINIMIZE AND CONTAIN ANTIMIC	PORIAL PROISTANCE
CACRCP 61-2005	ROBIAL RESISTANCE
INTRODUCTION	
AIMS AND OBJECTIVES	
RESPONSIBILITIES OF THE REGULATORY AUTHORITIES	
Quality Control of veterinary antimicrobial drugs	
Assessment of efficacy	
Assessment of the potential of veterinary antimicrobial drags to sel for resistant microorganisms	
Establishment of ADIs (acceptable daily intake). MRLs (maximum Withdrawal periods for veterinary antimicrobial drugs	
Establishment of a summary of product characteristics for each veterinary antimicrobial drug for food-producing animals	
Surveillance programmes	
Distribution of veterinary antimicrobial drugs in veterinary medicin	10
Control of advertising	
Training of veterinary antimicrobial drug users	
Development of research	
Collection and destruction of unused veterinary antimicrobial drug-	
RESPONSIBILITIES OF THE VETERINARY PHARMACEUTICAL	INDUSTRY
Marketing authorisation of veterinary antimicrobial drugs for food-	producing animals
Marketing and export of veterinary antimicrobial drugs	
Advertising	
Training	
Research	
RESPONSIBILITIES OF WHOLESALE AND RETAIL DISTRIBUT	ORS
RESPONSIBILITIES OF VETERINARIANS	
Off-label use	
Recording	
Training	
RESPONSIBILITIES OF PRODUCERS	
CONCLUSIONS	
Endnotes	
List of Abbreviations	
Glossary and Definitions of Terms	

Codex Prior AMR Work

- Code of Practice was completed in 2005
- Laid out responsibilities for regulation, distribution and use of antibiotics in food animals
- Much of it is duplicative of OIE Code of Practice



Codex Prior AMR Work

- First Task Force on AMR
- Guidelines for the Risk Analysis of Foodborne Antimicrobial Resistance
- Completed in 2011

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GUIDELINES FOR RISK ANALYSIS OF FOODBORNE ANTIMICROBIAL RESISTANCE

CAC/GL 77- 2011

Table of Contents

Introduction

scope

Definitions

General Principles for Foodborne AMR Risk Analysis

Framework for Foodborne AMR Risk Analysis

Preliminary Foodborne AMR Risk Management Activities

Identification of an AMR food safety issue

Development of a foodborne AMR risk profile

Ranking of the food safety issues and setting priorities for risk assessment and management

Establishment of preliminary risk management goals

Establishment of a risk assessment policy

Commission a foodborne AMR risk assessment

Foodborne AMR Risk Assessment

Sources of information

Process of foodborne AMR risk assessment

Hazard identification

Exposure assessment

Hazard characterization

Risk characterization

Foodborne AMR Risk Management

Consideration of the foodborne AMR risk assessment results

Identification of foodborne AMR RMOs

Evaluation of foodborne AMR RMOs

Selection of foodborne AMR RMOs

Implementation of foodborne AMR risk management decision(s)

Monitoring and review of foodborne AMR risk management measures

Surveillance of Use of Antimicrobial Agents and AMR Microorganisms and Determinants Foodborne AMR Risk Communication

Foodborne Risk Communication as a Risk Management Tool

Appendix 1. Elements for Consideration in a Foodborne AMR Risk Profile

Appendix 2, Suggested Elements for Consideration in a Foodborne AMR Risk Assessment

Appendix 3, Examples of Qualitative Foodborne AMR Risk Assessment

- Second Task Force on Antimicrobial Resistance (TFAMR) approved July, 2017
- Hosted by South Korea
- Physical Working Group to develop Scope hosted in London, December 2016
- E Working Groups on the documents
 - US*, China, Kenya and UK
 - Netherlands*, Chile, China and New Zealand

London Physical Working Group Developed Scope Documents

- Revise Code of Practice
- Develop a Surveillance and Monitoring Document
- Ask FAO for Additional Scientific Information



- Revision of the Code of Practice
 - Broaden scope beyond food animals
 - Risk-based guidance for entire food chain
 - Objective minimize risk to Public Health from development and spread of foodborne AMR
 - Scientifically supported, take into account new development
 - Consider lists of Critically Important Antimicrobials



- Surveillance and Monitoring
 - 1. Purpose provide guidance on design and implementation of integrated surveillance of AMR along the food chain
 - 2. Scope
 - Resistance
 - Use
 - Humans, Animals, Crops and Food



- E Working Groups have commented on a first draft of both documents
- Revisions made, out for comments (Due October 25)
- Will be considered at Step 3 at a TFAMR meeting in South Korea end of November
- FAO is also developing scientific information for consideration by the Task Force members

Challenges

- Scope Creep
 - Stray from Codex mandate of "through food"
 - Stray toward pre-harvest
- Cultural Differences on Regulation
 - Precautionary approach
- Different governments have different levels of interactions with their stakeholders
- Some country's Codex representatives may not have technical background



