

# codex alimentarius commission



FOOD AND AGRICULTURE  
ORGANIZATION  
OF THE UNITED NATIONS

WORLD  
HEALTH  
ORGANIZATION



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**Agenda Item 1**

**CX/RVDF 04/15/1**  
**June 2004**

## JOINT FAO/WHO FOOD STANDARDS PROGRAMME

### CODEX COMMITTEE ON RESIDUES OF VETERINARY DRUGS IN FOODS

#### Fifteenth Session

*Washington, DC (metro area), (United States of America), 26- 29 October 2004*

To be held at the Holiday Inn, Old Town Alexandria, 625 First Street, Alexandria, VA 22314  
(Washington, DC metro area) from Tuesday, 26 October at 10.00 hours to Friday, 29 October 2004

### PROVISIONAL AGENDA

Agenda Item	Subject Matter	Document Reference
1	Adoption of Agenda	CX/RVDF 04/15/1
2	Appointment of Rapporteur	
3	Matters Referred/of Interest to the Committee arising from the Codex Alimentarius Commission and Other Codex Committees and Task Forces	<a href="#">CX/RVDF 04/15/2</a>
4	Report of the 60 <sup>th</sup> and 62 <sup>nd</sup> Meetings of the Joint FAO/WHO Expert Committee on Food Additives	
5	Report of the OIE activities, including the Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products	<a href="#">CX/RVDF 04/15/3</a> <a href="#">CX/RVDF 04/15/3-Add.1</a>
6	Consideration of Maximum Residue Limits for Veterinary Drugs	
6 (a)	Draft Maximum Residue Limits for Veterinary Drugs <ul style="list-style-type: none"><li>Comments at Step 6 (<a href="#">CL 2003/24-RVDF</a>)</li></ul>	<a href="#">ALINORM 03/31A-App. IV and V</a> <a href="#">CX/RVDF 04/15/4</a> <a href="#">CX/RVDF 04/15/4-Add.1</a>
6 (b)	Proposed Draft Maximum Residue Limits for Veterinary Drugs <ul style="list-style-type: none"><li>Comments at Step 3 (<a href="#">CL 2004/17-RVDF</a>)</li></ul>	<a href="#">ALINORM 03/31A-App. VI</a> <a href="#">CX/RVDF 04/15/4A</a> <a href="#">CX/RVDF 04/15/4A-Add. 1</a>

- |        |                                                                                                                                                                                                                                                |                                                                             |
|--------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------|
| 7      | Proposed Draft Code of Practice to Minimize and Contain Antimicrobial Resistance                                                                                                                                                               | <a href="#">CL 2003/40-RVDF</a>                                             |
|        | <ul style="list-style-type: none"> <li>• Comments at Step 3 (CL 2003/40-RVDF)</li> </ul>                                                                                                                                                       | <a href="#">CX/RVDF 04/15/5</a><br><a href="#">CX/RVDF 04/15/5-Add.1</a>    |
| 8      | Proposed Draft Revised Guidelines for the Establishment of a Regulatory Program for the Control of Veterinary Drug Residues in Foods ( including Appendix on the Prevention and Control of Veterinary Drug Residues in Milk and Milk Products) | <a href="#">CX/RVDF 04/15/6</a>                                             |
|        | <ul style="list-style-type: none"> <li>• Comments at Step 3</li> </ul>                                                                                                                                                                         | <a href="#">CX/RVDF 04/15/6-Add. 1</a>                                      |
| 9      | Proposed Draft Revised Part II “General Consideration on Analytical Methods for Residue Control” of the Codex Guidelines for the Establishment of a Regulatory Program for the Control of Veterinary Drug Residues in Foods                    | <a href="#">CX/RVDF 04/15/7</a>                                             |
|        | <ul style="list-style-type: none"> <li>• Comments at Step 3</li> </ul>                                                                                                                                                                         | <a href="#">CX/RVDF 04/15/7-Add. 1</a>                                      |
| 10     | Discussion Paper on Risk Management Methodologies, Including Risk Assessment Policies, in the Codex Committee on Residues of Veterinary Drugs in Foods                                                                                         | <a href="#">CX/RVDF 04/15/8</a>                                             |
|        | <ul style="list-style-type: none"> <li>• Comments</li> </ul>                                                                                                                                                                                   | <a href="#">CX/RVDF 04/15/8-Add. 1</a>                                      |
| 11     | Methods of Analysis for Residues of Veterinary Drugs in Foods                                                                                                                                                                                  |                                                                             |
|        | <ul style="list-style-type: none"> <li>• Report of the <i>ad hoc</i> Working Group on Methods of Analysis and Sampling</li> </ul>                                                                                                              | CRD 1                                                                       |
| 11 (a) | Review of Performance-Based Criteria for Methods of Analysis for Residues of Veterinary Drugs in Foods                                                                                                                                         | <a href="#">CX/RVDF 03/10</a>                                               |
|        | <ul style="list-style-type: none"> <li>• Comments in response to <a href="#">CL 2003/11-RVDF</a>, Part C</li> </ul>                                                                                                                            | No comments received                                                        |
| 11 (b) | Identification of Routine Methods of Analysis for Veterinary Drug Residues in Foods                                                                                                                                                            | <a href="#">CX/RVDF 04/15/10</a>                                            |
|        | <ul style="list-style-type: none"> <li>• Comments in response to <a href="#">CL 2003/17-RVDF</a>, Part C</li> </ul>                                                                                                                            | <a href="#">CX/RVDF 04/15/10-Add.1</a>                                      |
| 12     | Consideration of the Priority List of Veterinary Drugs Requiring Evaluation or Re-evaluation                                                                                                                                                   |                                                                             |
|        | <ul style="list-style-type: none"> <li>• Comments in response to <a href="#">CL 2004/17-RVDF</a></li> </ul>                                                                                                                                    | <a href="#">CX/RVDF 04/15/11</a><br><a href="#">CX/RVDF 04/15/11-Add. 1</a> |
|        | <ul style="list-style-type: none"> <li>• Report of the <i>ad hoc</i> Working Group on Priority</li> </ul>                                                                                                                                      | CRD 2                                                                       |
| 13     | Other Business and Future Work                                                                                                                                                                                                                 |                                                                             |
| 13 (a) | Discussion Paper on Rounding of ADIs for Veterinary Drugs prior to Setting of MRLs                                                                                                                                                             | <a href="#">CX/RVDF 04/15/12</a>                                            |
| 14     | Date and Place of next Session                                                                                                                                                                                                                 |                                                                             |
| 15     | Adoption of the Report                                                                                                                                                                                                                         |                                                                             |

Working documents as prepared will be uploaded onto the Codex website. They can be downloaded and printed by accessing the following URL  
<http://www.codexalimentarius.net>  
Delegates are kindly requested to bring with them to the meeting all documents which have been distributed, as the number of additional copies which can be made available at the session is limited.

## NOTES ON THE PROVISIONAL AGENDA

### **Item 1 - Adoption of the Agenda (CX/RVDF 04/15/1)**

In accordance with Rule V.2 of the Rules of Procedure, the first item on the Provisional Agenda shall be the adoption of the Agenda.

### **Item 2 - Appointment of Rapporteur**

The Committee will be invited to appoint a Rapporteur to the Session.

### **Item 3 - Matters Referred/of Interest to the Committee arising from the Codex Alimentarius Commission and Other Codex Committees and Task Forces (CX/RVDF 04/15/2)**

Matters referred/of interest to the Committee arising from the 26<sup>th</sup> and 27<sup>th</sup> Sessions of the Codex Alimentarius Commission and other relevant Codex Committees and Task Forces are summarized in the working paper.

### **Item 4 - Reports of the 60<sup>th</sup> and 62<sup>nd</sup> Meetings of the Joint FAO/WHO Expert Committee on Food Additives (JECFA)**

The Secretariat of the Joint FAO/WHO Expert Committee on Food Additives will present a summary of the results of the 60<sup>th</sup> and 62<sup>nd</sup> JECFA Meeting convened in Geneva, Switzerland from 6 to 12 February 2003 and in Rome, Italy, from 4 to 12 February 2004 respectively. The reports are available online at: [http://www.fao.org/es/ESN/jecfa/whatisnew\\_en.stm](http://www.fao.org/es/ESN/jecfa/whatisnew_en.stm).

### **Item 5 - Report on OIE Activities, including the Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products (CX/RVDF 04/15/3)**

As at previous sessions of the Committee, the Committee will be presented with a report on the relevant activities of the OIE and VICH.

### **Item 6 - Consideration of Maximum Residue Limits for Veterinary Drugs**

#### **Item 6 (a) - Draft Maximum Residue Limits for Veterinary Drugs (ALINORM 03/31A-App. IV and App. V; CX/RVDF 04/15/4)**

The Committee will consider draft MRLs retained at Step 6 by the 14<sup>th</sup> Session of the CCRVDF for flumequine, neomycin, dicyclanil, melengestrol acetate and trichlorfon (metrifonate) (ALINORM 03/31A paras 57 and 63 and Appendix IV).

In addition, the Committee will consider draft MRLs returned to Step 6 (phoxim) and MRLs advanced to Step 6 (cyhalothrin and cefuroxime) by the 26<sup>th</sup> Session of the Codex Alimentarius Commission (ALINORM 03/41, paras. 112, 116 and 136 and Appendix VI).

Comments at Step 6 submitted in response to CL 2003/24-RVDF are summarized in CX/RVDF 04/15/4.

#### **Item 6 (b) - Proposed Draft Maximum Residue Limits for Veterinary Drugs (ALINORM 03/31A-App. VI; CX/RVDF 04/15/4A)**

The Committee will consider MRLs for cypermethrin and alpha-cypermethrin which were retained at Step 4 by its 14<sup>th</sup> Session (ALINORM 03/31A, para. 65 and Appendix VI).

In addition the Committee will consider proposed draft MRLs arising from the recommendations of 60<sup>th</sup> and 62<sup>nd</sup> Meeting of JECFA for: imidocarb, flumequine (Black tiger shrimp), pirlimycin, cypermethrin/alpha cypermethrin, doramectin and ractopamine.

Comments at Step 3 submitted in response to CL 2004/17-RVDF, Part A are summarised in CX/RVDF 04/15/4A.

**Item 7 - Proposed Draft Code of Practice to Minimize and Contain Antimicrobial Resistance (CL 2003/40-RVDF; CX/RVDF 04/15/5)**

The 14<sup>th</sup> Session of the CCRVDF agreed that a drafting group, under the direction of the United States, would prepare a revised version of the proposed draft Code of Practice for circulation, comments at Step 3 and further consideration at its 15<sup>th</sup> Session. The Committee agreed that the proposed draft Code of Practice would be revised by the drafting group on the basis of the discussion, written comments submitted at its 14<sup>th</sup> Session and comments submitted in response to CL 2003/11-RVDF, Part B (ii) (ALINORM 03/31A, paras 79-80).

The document prepared by the working group was circulated under CL 2003/40-RVDF. Comments submitted at Step 3 are summarized in CX/RVDF 04/15/5.

**Item 8 - Proposed Draft Revised Guidelines for the Establishment of a Regulatory Program for the Control of Veterinary Drug Residues in Foods ( including Appendix on the Prevention and Control of Veterinary Drug Residues in Milk and Milk Product) (CX/RVDF 04/15/6; CX/RVDF 04/15/6-Add. 1)**

The 14<sup>th</sup> Session of the CCRVDF agreed that a drafting group, led by New Zealand, would prepare a revised version of the guidelines for circulation, comments at Step 3 and further consideration at its 15<sup>th</sup> session. It was agreed that the Guidelines would be revised by the drafting group on the basis of the comments submitted and would include the proposed draft Appendix on the Control of Veterinary Drug Residues in Milk and Milk products (ALINORM 03/31A, para. 85).

Comments submitted at Step 3 are summarised in CX/RVDF 04/15/6-Add.1.

**Item 9 - Proposed Draft Revised Part II “General Consideration on Analytical Methods for Residue Control” of the Codex Guidelines for the Establishment of a Regulatory Program for the Control of Veterinary Drug Residues in Foods (CX/RVDF 04/15/7; CX/RVDF 04/15/7-Add. 1)**

The 14<sup>th</sup> Session of the Committee agreed that a drafting group would review Part II “General Considerations on Analytical Methods for Residue Control” of the Guidelines for the Establishment of a Regulatory Programme for Control of Veterinary Drug Residues in Foods (CAC/GL 16-1993) for circulation, comment at Step 3 and further consideration at its 15<sup>th</sup> meeting (ALINORM 03/31A, para. 105).

Comments submitted at Step 3 are summarised in CX/RVDF 04/15/7-Add.1.

**Item 10 - Discussion Paper on Risk Management Methodologies, Including Risk Assessment Policies, in the Codex Committee on Residues of Veterinary Drugs in Foods (CX/RVDF 04/15/8; CX/RVDF 04/15/8-Add. 1)**

The 14<sup>th</sup> Session of the Committee agreed that a working group, led by France, would prepare a revised version of the discussion paper on “Risk Management Methodologies, including Risk Assessment Policies in the Codex Committee on Residues of Veterinary Drugs in Foods” for circulation, additional comments and further consideration at its 15<sup>th</sup> Session. The Committee agreed that the revised document should specifically address the issue of substances with no ADI and/or MRL, should take account of the discussion and written comments received during the Session and the comments of the 60<sup>th</sup> meeting of JECFA on Annex I of CX/RVDF 01/9 (ALINORM 03/31A paras 95-96).

Comments submitted are summarised in CX/RVDF 04/15/8-Add.1.

**Item 11 - Methods of Analysis for Residues of Veterinary Drugs in Foods (CRD 1)**

The 14<sup>th</sup> CCRVDF agreed to convene the *ad hoc* Working Group prior to its 15<sup>th</sup> Session under the Chairmanship of Canada and the Netherlands. The report of the *ad hoc* Working Group will be circulated as Conference Room Document 1 (CRD 1).

**Item 11 (a) - Review of Performance-Based Criteria for Methods of Analysis for Residues of Veterinary Drugs in Foods (CX/RVDF 03/10; CX/RVDF 04/15/9)**

The 14<sup>th</sup> CCRVDF considered a document (CX/RVDF 03/10) on criteria relating to the selection of methods of analysis for veterinary drugs contained in the *Guidelines for the Establishment of a Regulatory Programme for Control of Veterinary Drugs in Foods* (CAC/GL 16-1993).

The 15<sup>th</sup> CCRVDF will continue its consideration of document CX/RVDF 03/10. Comments submitted in response to CL 2003/11, Part C requesting information on needs of developing countries related to validation of analytical methods, are summarised in CX/RVDF 04/15/9.

**Item 11 (b) - Consideration of the Identification of Routine Methods of Analysis for Veterinary Drug Residues in Foods (CX/RVDF 04/15/10)**

The 14<sup>th</sup> CCRVDF agreed that the *ad hoc* Working Group continue its work on the review and recommendation of methods of analysis and the updating of methods validation procedures (ALINORM 03/31A, para. 109) including additional information provided in response to CL 2004/17-RVDF, Part C.

**Item 12 - Consideration of the Priority List of Veterinary Drugs Requiring Evaluation or Reevaluation (CX/RVDF 04/15/11; CRD 2)**

The 14<sup>th</sup> CCRVDF agreed to convene the *ad hoc* Working Group on Priorities prior to its 15<sup>th</sup> Session under the Chairmanship of Australia to consider proposals for compounds to be evaluated or re-evaluated by JECFA (ALINORM 03/31A, para. 115). Recommendations of the *ad hoc* Working Group will be summarised in Conference Room Document 2 (CRD 2).

Comments submitted in response to CL 2004/17-RVDF, Part B, which will be considered by the *ad hoc* Working Group, are summarised in document CX/RVDF 04/15/11.

**Item 13 - Other Business and Future Work**

Other business and proposals for future work will be considered

**Item 13 (a) - Discussion Paper on Rounding of ADIs for Veterinary Drugs prior to Setting of MRLs (CX/RVDF 04/15/12)**

The paper has been added to the provisional agenda at the request of the US Government.

**Item 14 - Date and Place of Next Session**

The Chairperson will propose, on behalf of the host country, the tentative date and place of the next meeting.

**Item 15 - Adoption of the report**

The Committee shall adopt a report of its 15<sup>th</sup> Session based on a draft provided by the Secretariat.