

# CODEX COMMITTEE ON RESIDUES OF VETERINARY DRUGS IN FOODS

17<sup>th</sup> Session

Breckenridge, Colorado, USA, 3 -7 September 2007

## PROVISIONAL ANNOTATED AGENDA

### Division of Competence

**between the European Community and its Member States according to  
Rule of procedure II paragraph 5 of the Codex Alimentarius Commission**

Agenda Item	Subject Matter	Document Reference
1.	Adoption of the Agenda  <i>Member States competence. Member States vote.<sup>1</sup></i>	CX/RVDF 07/17/1
2.	Matters Referred by the Codex Alimentarius Commission and Other Codex Committees and Task Forces  <u>Part A:</u> <i>European Community Competence. European Community Vote.</i>  <u>Part B:</u> <i>Member States Competence. Member States Vote.</i>  <u>Part C:</u> <i>Mixed Competence. European Community Vote.</i>	CX/RVDF 07/17/2
3.	Matters of Interest Arising from FAO/WHO  <i>Mixed Competence. Member States Vote.</i>  a) 66th Meeting of the Joint FAO/WHO Expert Committee on Food Additives (JECFA) ( <i>in English only</i> )	CX/RVDF 07/17/3     <a href="http://whqlibdoc.who.int/">http://whqlibdoc.who.int/</a>

<sup>1</sup> This is without prejudice to the substantive competence that the Community has for individual items on the Agenda.

	<p><b>European Community Competence.</b> <b>European Community Vote.</b></p>	<p>publications/2006/ 9241209179_eng.pdf</p>
4.	<p><b>Report of the OIE activities, including the Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products</b></p> <p><b>European Community Competence.</b> <b>European Community Vote.</b></p>	<p><b>CX/RVDF 07/17/4</b></p>
5.	<p><b><u>Consideration of Maximum Residue Limits (MRLs) for Veterinary Drugs</u></b></p> <p><b>a) Draft MRLs for Veterinary Drugs (at Step 7)</b></p> <ul style="list-style-type: none"> <li>- Information on registered use of Flumequine in Black tiger shrimp and in shrimps (CL 2006/14-RVDF, part C)</li> </ul> <p><b>European Community Competence.</b> <b>European Community Vote.</b></p> <p><b>b) Draft MRLs for Veterinary Drugs (at Step 6)</b></p> <ul style="list-style-type: none"> <li>- Comments at Step 6 (CL 2006/35-RVDF)</li> </ul> <p><b>European Community Competence.</b> <b>European Community Vote.</b></p> <p><b>c) Proposed Draft MRLs for Veterinary Drugs (at Step 4)</b></p> <p><b>European Community Competence.</b> <b>European Community Vote.</b></p> <p><b>d) Proposed Draft MRLs for Veterinary Drugs (at Step 3)</b></p> <ul style="list-style-type: none"> <li>- Comments at Step 3 (CL 2006/14-RVDF, part C)</li> </ul> <p><b>European Community Competence.</b> <b>European Community Vote.</b></p>	<p><b>CX/RVDF 07/17/5</b></p> <p><b>ALINORM 06/29/31, App. III</b></p> <p><b>CX/RVDF 07/17/6</b></p> <p><b>ALINORM 06/29/31, App. IV</b></p> <p><b>CX/RVDF 07/17/7</b></p> <p><b>ALINORM 06/29/31, App. V</b></p> <p><b>ALINORM 06/29/31, App. VI</b></p> <p><b>CX/RVDF 07/17/8</b></p>

6.	<p><b>Draft Guidelines for the Design and Implementation of National Regulatory Food Safety Assurance Programmes Associated with the Use of Veterinary Drugs in Food Producing Animals</b></p> <ul style="list-style-type: none"> <li>- Comments at Step 6 (CL 2006/35-RVDF)</li> </ul> <p><b>European Community Competence. European Community Vote.</b></p>	<p><b>ALINORM 06/29/31, App. VII</b></p> <p><b>CX/RVDF 07/17/9 CX/RVDF 07/17/9-Add. 1</b></p>
7.	<p><b>Methods of Analysis for Residues of Veterinary Drugs in Foods</b></p> <ul style="list-style-type: none"> <li>- Comments (CL 2007/04-RVDF)</li> <li>- Report of the <i>ad hoc</i> Working Group on Methods of Analysis and Sampling</li> </ul> <p><b>Mixed Competence. European Community Vote.</b></p>	<p><b>CL 2007/04-RVDF</b></p> <p><b>CX/RVDF 07/17/10 CX/RVDF 07/17/10-Add. 1</b></p> <p><b>CRD 1</b></p>
8.	<p><b>Priority List of Veterinary Drugs Requiring Evaluation or Reevaluation</b></p> <ul style="list-style-type: none"> <li>- Comments (CL 2006/52-RVDF)</li> <li>- Report of the <i>ad hoc</i> Working Group on Priority</li> </ul> <p><b>European Community Competence. European Community Vote.</b></p>	<p><b>CX/RVDF 07/17/11</b></p> <p><b>CRD 2</b></p>
9.	<p><b>Report of the physical Working Group on Residues of Veterinary Drugs without ADI/MRL</b></p> <ul style="list-style-type: none"> <li>- Comments</li> </ul> <p><b>European Community Competence. European Community Vote.</b></p>	<p><b>CX/RVDF 07/17/12</b></p> <p><b>CX/RVDF 07/17/12-Add. 1</b></p>

10.	<p><b>Discussion Paper on Risk Management Topics and Options for the CCRVDF</b></p> <p>- Comments</p> <p><b>Mixed Competence.</b> <b>European Community Vote.</b></p>	<p><b>CX/RVDF 07/17/13</b></p> <p><b>CX/RVDF 07/17/13</b> <b>-Add.1</b></p>
11.	<p><b>Other Business and Future Work</b></p> <p><i>Competence and right to vote to be determined in the light of the issues to be dealt with (document not yet available).</i></p>	
12.	<p><b>Date and Place of the Next Session</b></p> <p><b>Member States competence.</b> <b>Member States vote.</b></p>	
13.	<p><b>Adoption of the Report</b></p> <p><b>Member States Competence.</b> <b>Member States Vote.<sup>2</sup></b></p>	

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<sup>2</sup> This is without prejudice to the substantive competence that the Community has for individual items in the Report.