

JOINT FAO/WHO FOOD STANDARDS PROGRAMME

CODEX ALIMENTARIUS COMMISSION

Thirtieth Session

FAO Headquarters, Rome, Italy, 2–7 July 2007

Report

EXECUTIVE SUMMARY

The Commission:

- a) Adopted amendments to the Rules of Procedure and other amendments to the Procedural Manual;
- b) Adopted 44 new or revised Codex standards or related texts;
- c) Approved a number of new work proposals and proposals for discontinuation of work;
- d) Noted the Codex budget proposals for 2008-09 and expressed its strong desire that FAO and WHO allocate, to the Codex programme, a budget level that would allow the Secretariat to conserve, at least, the same purchasing power as in the 2006-07 biennium; and made recommendations on the use of Russian and Portuguese in selected Codex meetings, subject to certain conditions;
- e) Adopted the Strategic Plan 2008-2013 of the Codex Alimentarius Commission;
- f) Noted with satisfaction the progress made in implementing the proposals based on the recommendations from the Codex Evaluation (2002); agreed to some principles to streamline the Codex committee structure and session planning, while requesting the Executive Committee to follow up on other pending issues;
- g) Agreed to re-activate the Codex Committee on Natural Mineral Waters to update the provisions on health-related substances in the current standard;
- h) Supported continued cooperation and coordination with international governmental and non-governmental organizations; recommended that FAO and WHO study the possibility of reviewing or updating FAO and WHO Agreements with OIE, as might be required; and requested the Codex Secretariat to identify, in cooperation with the Legal Offices of FAO and WHO, any practical problems affecting the cooperation between Codex and OIE that might need to be addressed in a pragmatic manner;
- i) Expressed its appreciation to FAO and WHO and to the donors making financial contribution to the Codex Trust Fund;
- j) Expressed its appreciation to FAO and WHO for their ongoing activities in support of Codex, namely provision of scientific advice and capacity building in food safety and quality;
- k) Elected the following Officers of the Commission and other members of the Executive Committee:
 - **Chairperson:** Mr Claude J.S. Mosha (United Republic of Tanzania),
 - **Vice-Chairpersons:** Ms Karen Hulebak (USA), Ms. Noraini Mohd. Othman (Malaysia) and Mr Wim van Eck (the Netherlands)
 - **Regional Coordinators:** Ghana, Indonesia, Switzerland, Mexico, Tunisia and Tonga
 - **Other Members:** Mali, Japan, United Kingdom, Argentina, Jordan, Canada, New Zealand
- l) Designated/confirmed the host governments of Codex subsidiary bodies, including Malaysia as new host country for the Committee on Fats and Oils

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INTRODUCTION

1. The Codex Alimentarius Commission held its Thirtieth Session at FAO Headquarters, Rome, Italy, from 2-7 July 2007. Dr Claude J. S. Mosha (Tanzania), Chairperson of the Commission presided over the Session, assisted by the Vice-Chairpersons, Ms Noraini Mohd. Othman (Malaysia), Dr Karen Hulebak (United States of America) and Dr Wim van Eck (Netherlands). The Session was attended by 516 delegates from 122 Member countries and 1 Member Organisation, 40 international governmental and non-governmental organizations including UN agencies. A list of participants, including the Secretariat, is given in Appendix I to this report.

2. The Session was opened by Mr J. M. Sumpsi, Assistant Director-General, Agriculture and Consumer Protection Department, FAO and Dr G. Moy, Scientist, Department of Food Safety, Zoonoses and Foodborne Disease, WHO on behalf of Ms Weber-Mosdorf, Assistant Director-General, Sustainable Development and Healthy Environments, WHO, respectively.

3. A minute's silence was observed in memory of Mr Graham Kermodé, Joint Secretary of the Commission from 1964 to 1983, who had played an important role in the setting up and early running of the Codex Alimentarius Commission.

Division of Competence

4. The Commission noted the division of competence between the European Community and its Member States, according to paragraph 5, Rule II of the Procedure of the Codex Alimentarius Commission, as presented in document CAC/30 LIM/2.

ADOPTION OF THE AGENDA (Agenda Item 1)¹

5. The Commission adopted the Provisional Agenda as its Agenda for the Session.

6. At the request of the Delegation of Colombia, the Commission agreed to discuss, under Agenda Item 20, "Other Business", if time allowed, the necessity of circulating Codex documents simultaneously in all working languages.

7. The Commission, upon the proposal of the Delegation of Switzerland, the host country for the Codex Committee on Natural Mineral Waters and with a view to facilitating the Commission's discussion on the subject matter under Agenda Item 13, agreed to convene an in-session working group under the chairmanship of Switzerland, opened to all interested members and observers and working in English only, to review the comments submitted in response to the Circular Letter (CL) 2006/13-NMW and to consider whether an amendment to the Codex Standard for Natural Mineral Waters was necessary and, if affirmative, how to proceed.

REPORT BY THE CHAIRPERSON ON THE 59th SESSION OF THE EXECUTIVE COMMITTEE (Agenda item 2)²

8. In accordance with Rule V.7 of the Rules of Procedure, the Chairperson reported to the Commission on the outcome of the 59th Session of the Executive Committee, as follows.

9. The Chairperson recalled that the Executive Committee had not met since the 29th Session of the Commission and therefore had considered both the Critical Review of standards and related texts submitted to the Commission for adoption and the monitoring of standards development. It had recommended adoption of all the texts proposed for adoption and made some recommendations of a general nature, especially as regards food additives. As regards new work, some general recommendations had also been made considering the preparation of project documents and specific comments on individual proposals.

10. As regards the structure and mandate of Codex Committees the Executive Committee had considered the Proposal related to commodity work of regional committees (Proposal 8 of CL 2006/29-CAC) and

¹ ALINORM 07/30/1 and ALINORM 07/30/1A Rev.1

² ALINORM 07/30/3

recommended a revised text for this Proposal, which had been used in the consideration of the Proposed Draft Standards and proposals for new work originating from the Coordinating Committee for Asia.

11. The Executive Committee had considered in detail financial and budgetary matters and had proposed a revised version of the Draft Strategic Plan 2008-2013, taking into account the comments from Coordinating Committees and further discussion during the session.

12. In accordance with Rule IX.6, the Executive Committee had considered the application of five international non-governmental organizations and had recommended to the Directors-General of FAO and WHO to grant them Observer status with the Codex Alimentarius Commission.

13. The Commission noted that this report was presented for information with the understanding that the recommendations made by the Executive Committee on specific questions would be considered under the relevant Agenda items.

REPORTS OF FAO/WHO REGIONAL COORDINATING COMMITTEES (Agenda Item 3)³

14. The Commission noted the reports of the Coordinating Committees, as presented by their respective Coordinators. The Coordinators expressed their gratitude for the capacity building activities of FAO and WHO and the Codex Trust Fund, while expressing the wish that such activities would continue in order to facilitate the participation of countries in Codex work. The Coordinating Committees had discussed food legislation and food control systems, consumer participation, the use of Codex standards, the Draft Strategic Plan, and had expressed their views on issues of interest to their regions.

FAO/WHO Coordinating Committee for Africa

15. The Delegation of Morocco informed the Commission that the Coordinating Committee for Africa had endorsed a Strategic Plan to enhance participation of countries of the region in Codex and was developing a regional plan of action for the CCAFRICA together with a set of indicators for evaluation and a mechanism for monitoring its progress. A website with a discussion forum had been developed and was operational within the region. Several countries in the region were harmonising national legislation with Codex standards, although some difficulties still existed in the process and countries in the region still faced difficulties in accessing international markets due to the fact that importing countries adopted more stringent standards than Codex standards.

FAO/WHO Coordinating Committee for Asia

16. The Delegation of the Republic of Korea reported the progress of the four commodity standards under elaboration by the Coordinating Committee for Asia and noted that the Proposed Draft Standard for Gochujang and the Proposed Draft Standard for Ginseng Product had been forwarded to the Commission for adoption at Step 5, with the recommendation by the Committee to finalize these standards in relevant Commodity Committees. The Committee further agreed to request the Commission for approval of new work on two commodity standards. The Committee had initiated the consideration of the Strategic Plan for CCASIA, with the goal of strengthening the food safety infrastructure of the countries of the region.

FAO/WHO Coordinating Committee for Latin America and the Caribbean

17. The Delegation of Argentina informed the Commission about different Codex related activities held in conjunction with the Coordinating Committee, including the development of a project platform to enhance food quality and safety in the region as a follow-up to the recommendations of the FAO/WHO Regional Conference on Food Safety in the Americas. Other activities aimed at improving communication within the region included the enhancement of the CCLAC webpage (www.cclac.org); the establishment of electronic fora to discuss Codex matters of interest to the region and the development of e-learning tools, and presently the second online course to increase knowledge about Codex had been initiated. In addition, the Delegation highlighted some of the matters of interest to the region such as the establishment of and compliance with Codex maximum residue limits for pesticides, and the matters associated with the availability of data to set Codex MRLs for pesticides and the criteria for the establishment of such MRLs.

³ ALINORM 07/30/28, ALINORM 07/30/15, ALINORM 07/30/36, ALINORM 07/30/19, ALINORM 07/30/40, ALINORM 07/30/32

FAO/WHO Coordinating Committee for Europe

18. The Delegation of Switzerland indicated that the Coordinating Committee had recognised the need to further encourage participation from countries outside the European Union; had supported Codex work on the implementation of the Global Strategy on Diet, Physical Activity and Health; had stressed the importance of risk analysis for food control; and had urged FAO/WHO to allocate sufficient resources to allow Codex to fulfil its mandate, especially as regards Coordinating Committees. A website to foster intra-regional cooperation had been created (www.codexeurope.ch) and three regional training courses had been organized. The Delegation noted that food legislation was well harmonized within the European Community and the European Economic Area and that it was a long standing practice in Europe to involve consumers in food legislation, food control and Codex matters.

FAO/WHO Coordinating Committee for the Near East

19. The Delegation of Jordan informed the Commission that the Coordinating Committee had advanced three draft regional standards for final adoption and was working on the proposed draft Regional Guidelines for Street-Vended Foods. The Committee had endorsed the recommendations of the 28th Session of the Commission regarding the role and activities of Regional Coordinating Committees and recommended that the Committee on General Principles should seek to define the respective roles of the Coordinators and members of the Executive Committee elected on a geographical basis. It had supported the adoption of the draft Strategic Plan 2008-2013 of the Commission; had expressed its appreciation to FAO and WHO for their work in the area of scientific advice; and supported various coordinating activities in the region.

FAO/WHO Coordinating Committee for North America and the South West Pacific

20. The Delegation of New Zealand, on behalf of the Delegation of Samoa, absent, presented the report and indicated that the Coordinating Committee had made several specific comments on the draft Strategic Plan 2008-2013 and on the review of the Structure and Mandate of Codex Committees and Task Forces. The Committee also discussed the status of implementation of the Strategic Plan for the CCNASWP, the objectives of which included the improvement of coordination and communication of regional activities on Codex and the promotion of participation of all members countries of the region in the activities of the Committee. The Delegation congratulated Samoa for the excellent work carried out as Coordinator and wished to the delegate of Samoa a very prompt recovery.

AMENDMENTS TO THE PROCEDURAL MANUAL OF THE CODEX ALIMENTARIUS COMMISSION (Agenda Item 4)⁴

Proposed Amendments to the Rules of Procedure

Amendments concerning the role of coordinators and members elected on a geographical basis

21. The Commission determined that the quorum specified in Rule VI.7 for the amendment of the Rules of Procedure was constituted⁵. The Commission noted that there was general support for the amendments to Rules IV. Coordinators, Rule V. Executive Committee and Rule XI. Subsidiary Bodies, as proposed by the Committee on General Principles. In accordance with Rule VIII.7 and XV.1 of the Commission's Rules of Procedure and Rule XII.7 of the General Rules of FAO, the Commission agreed to proceed to a single roll call vote for all amendments, as they all concerned the role of coordinators and members elected on a geographical basis, with the following results.

Votes in favour: Algeria, Angola, Antigua and Barbuda, Argentina, Armenia, Australia, Austria, Belgium, Benin, Bhutan, Botswana, Brazil, Bulgaria, Burkina Faso, Burundi, Cameroon, Canada, Chile, China, Colombia, Costa Rica, Cote d'Ivoire, Croatia, Cuba, Cyprus, Czech Republic, Democratic Republic of the Congo, Denmark, Dominican Republic, Egypt, El Salvador, Eritrea, Estonia, Ethiopia, Fiji, Finland, France, Gabon, Germany, Ghana, Greece, Guinea, Honduras, Hungary, India, Indonesia, Islamic Republic of Iran, Ireland, Jamaica, Japan, Jordan, Kenya,

⁴ ALINORM 07/30/4, ALINORM 07/30/4A (comments of Islamic Republic of Iran, CIAA), ALINORM 07/30/4A-Add.1 (comments of Brazil), CAC/30 LIM/9 (comments of Ghana)

⁵ The number of Codex Member Countries [174] / 2 + 1 = 88

Kuwait, Lao People's Democratic Republic, Latvia, Lithuania, Malawi, Malaysia, Mali, Malta, Mexico, Morocco, Mozambique, Netherlands, New Zealand, Niger, Norway, Panama, Papua New Guinea, Paraguay, Peru, Poland, Portugal, Republic of Korea, Saudi Arabia, Senegal, Serbia, Singapore, Slovakia, Slovenia, South Africa, Spain, Sri Lanka, Sudan, Sweden, Switzerland, Syrian Arab Republic, Tanzania, Thailand, Tunisia, Turkey, Uganda, United Arab Emirates, United Kingdom, United States of America, Vanuatu, Viet Nam, Zambia, Zimbabwe.

Votes against:	None
Abstaining:	Bolivia, Nigeria
Tally:	99 votes cast, 99 in favour, 0 against, 2 abstentions (two-thirds majority required: 66)
Result:	The amendments were adopted

22. The Commission noted that the amendments to the Rules of Procedure would enter into force only after their approval by the Directors-General of FAO and WHO (Rule XVI). The amendments to the Rules as adopted by the Commission are presented in Appendix II to the present report.

23. The Delegation of Italy subsequently indicated that it was in agreement with the amendment proposed.

Proposals to Amend Other Sections of the Procedural Manual

Amendments to the Procedures for the Elaboration of Codex Standards and Related texts - *Guide to the Procedure for the Revision and Amendment of Codex Standards and Arrangements for the Amendments of Codex Standards Elaborated by Codex Committees which have been adjourned sine die*

24. The Delegation of Colombia, supported by other delegations, expressed the view that the provisions for methods of analysis and sampling should be considered as revisions rather than amendments as they were of a substantial nature and could create technical barriers to trade.

25. The Delegation indicated that the current text in the Procedural Manual in the Spanish version did not refer to "updating" but only to "finalization" of methods of analysis. The Commission noted that the current text in the English version was identical and that the changes to the proposed text resulted from the discussion in the Committee on General Principles.

26. The Secretariat recalled that methods of analysis and sampling were considered for adoption by the Commission as any other provision in Codex standards, and noted that the text under consideration specified that the Commission had the final authority to decide whether an amendment was of an editorial or substantive nature.

27. Some delegations pointed out that these questions had been discussed in detail in the Committee on General Principles and supported the current text. After some discussion, the Commission agreed to adopt the text as proposed, recalling that it would have the authority to decide on any proposed amendment or revision.

Amendments to the *General Principles of the Codex Alimentarius*

28. The Delegation of Argentina expressed its reservation on paragraph 3 on the Nature of Codex Standards, as, in their view, it undermined the importance of Codex standards as an international reference in the framework of the WTO.

Amendment to the *Principles Concerning the Participation of International Non-Governmental Organizations in the Work of the Codex Alimentarius Commission*

Proposed amendments to the *Format for Commodity Standards*

29. The Commission adopted the above amendments as proposed.

Draft Risk Analysis Principles Applied by the Committee on Pesticide Residues

30. The Delegation of Argentina was of the view that the proposed procedures for the periodic review and criteria for deletion of the Maximum Residue Limits (MRLs) were not fully based on science and did not take sufficiently into consideration the concerns and situation of developing countries. The Delegation

pointed out that the issue to be addressed related to the procedures applied by the Committee on Pesticide Residues and should not be confused with the trade related problems that had also been discussed in the Committee and were also under consideration in the WTO Committee on Sanitary and Phytosanitary Measures, and proposed to return the document for further consideration by the Committee on Pesticide Residues. This view was supported by several delegations. Some delegations also pointed out that the revocation of MRLs in the framework of Codex should be based on the relevant new scientific evidence and not on commercial considerations or on a pre-established revision period, especially as Codex standards were an international reference in the framework of the WTO SPS Agreement.

31. Several other delegations pointed out that all aspects of the document had been discussed at length in the Committee on Pesticide Residues and clearly described the procedures applied in the Committee, as well as its relationship with the JMPR. These delegations stressed the need to adopt the Draft Risk Analysis Principles in order to document the risk analysis policies and procedures applied to the establishment of MRLs, in accordance with the earlier decision of the Commission.

32. After some discussion, the Commission adopted the document as proposed, with the understanding that, in accordance with the Strategic Plan, this matter could be further considered when the Committee on General Principles reviewed all relevant texts on risk analysis policies applied by Codex Committees as a whole, in order to ensure consistency throughout Codex.

33. The Delegations of Argentina, Brazil, Chile, Colombia, Costa Rica, Dominican Republic, Ecuador, Guatemala, India, Indonesia, Jamaica, Panama, Paraguay, and Peru expressed their reservation on this decision, not only because what is mentioned above but also because in their view consensus had not been reached.

Proposed Draft Risk Management Methodologies, including Risk Assessment Policies in the Committee on Residues of Veterinary Drugs in Foods

34. The Delegation of Australia, while supporting the adoption of the text, expressed the view that when reviewing the risk analysis policies applicable to the Codex Committees concerned, the Committee on General Principles should review the provisions on data protection and confidentiality in order to ensure consistency throughout Codex. The Commission adopted the document as proposed.

Proposed Amendment of the Principles for the Establishment or Selection of Codex Sampling Procedures

Proposed Procedure for Consideration of the Entry and Review of Food Additive Provisions in the General Standard for Food Additives

Proposed amendments to harmonise the text concerning the membership of the Coordinating Committee for Europe with that of the other Coordinating Committees

Proposed Amendments to the Terms of Reference for the Committee on Food Additives

Proposed Amendments to the Terms of Reference for the Committee on Contaminants in Foods

Proposed Amendments to the Risk Analysis Principles Applied by the Codex Committee on Food Additives and Contaminants

Proposed Amendments to the CCFAC Policy for Exposure Assessment of Contaminants and Toxins in Foods or Food Groups

Proposed Definition for Codex Maximum Level for a Contaminant in a Food or Feed Commodity

35. The Commission adopted the above texts as proposed.

Proposed Amendments to the Format for Codex Commodity Standards – Food Additives and Relations between Commodity Committees and General Committees

36. The Commission recalled that the proposed changes to the *Relations between Commodity Committees and General Committees* applied only to additives and that, if they were adopted, the current provisions applicable to contaminants would be lost, and noted that the Executive Committee had recommended to defer the adoption of the amendment proposed, with the understanding that the Secretariat would review the current provisions and prepare a revised proposed amendment addressing both additives and contaminants.

37. Some delegations expressed the view that it would be preferable to adopt the proposed text at the present session as it was necessary in order to clarify the relations between Commodity Committees and the Committee on Food Additives, although it would result in the deletion of the provisions applicable to contaminants in the immediate future. It was however recognised that similar provision for contaminants should be developed for inclusion in the *Relations between Commodity Committees and General Committees*. The Commission therefore adopted the provisions applicable to additives as proposed, and recommended that the Committee on Contaminants in Foods develop necessary provisions for contaminants, on the basis of a draft to be prepared by the Secretariat.

38. The amendments to the Procedural Manual as adopted by the Commission are presented in Appendix III to the present report.

DRAFT STANDARDS AND RELATED TEXTS AT STEP 8 OF THE PROCEDURE (Agenda Item 5)⁶

39. The Commission **adopted** all of the Draft Standards and Related Texts submitted by its subsidiary bodies at Step 8 (including those submitted at Step 5 with a recommendation to omit Steps 6 and 7 and those submitted at Step 5 of the Accelerated Procedure) as presented in Appendix IV to this report.

40. The following paragraphs provide additional information on the comments made and the decisions taken on certain items.

Contaminants in Foods

Draft Maximum Levels for Tin in Canned Foods (other than beverages) and in Canned Beverages⁷

41. The Committee **adopted** the draft Maximum Levels and agreed to include them in Schedule I of the General Standard for Contaminants and Toxins in Foods (GSCTF), with the understanding that the existing maximum levels for tin in certain canned foods included in Schedule I of the GSCTF would be replaced by the adopted maximum levels.

42. The Delegation of the European Community maintained its reservation expressed at the First Session of the Codex Committee on Contaminants in Foods, stating that the proposed maximum levels for tin might lead to the PTWI set by JECFA being exceeded in certain vulnerable groups, that the maximum levels for tin should be set as low as reasonably achievable and that the technological need did not justify the proposed levels.

Proposed Draft Code of Practice for the Prevention and Reduction of Ochratoxin A Contamination in Wine (N05-2006)⁸

43. The Committee **adopted** the proposed draft Code of Practice at Step 8, omitting Steps 6 and 7, with some editorial changes in the English text.

⁶ ALINORM 07/30/5; ALINORM 07/30/3; ALINORM 07/30/5A (comments of Argentina, Australian, Bolivia, Brazil, European Community, Ecuador, Guatemala, Iran, Lebanon, Mexico, Peru, United States of America, CIAA, IADSA, ISDI and WPTC); ALINORM 07/30/5A Corr. (comments of Argentina); CAC/30 LIM/3 (comments of France, Malaysia, Peru, South Africa and Sri Lanka), CAC/30 LIM/7 (comments of European Community, Guatemala, Indonesia, Philippines and Switzerland); CAC/30 LIM/12 (comments of France); CAC/30 LIM/15 (comments of Kenya); CAC/30 LIM/17 (comments of the United States of America); CAC/30 LIM/18 (comments of Honduras)

⁷ ALINORM 07/30/41, Appendix IX

⁸ ALINORM 07/30/41, Appendix VIII

Food Additives

*Draft and Proposed Draft Food Additive Provisions of the General Standard for Food Additives (GSFA)*⁹

44. The Commission noting the recommendations of the 59th Session of the Executive Committee, within the framework of the Critical Review, **agreed** that: (i) no consequential changes should be made to commodity standards at this stage when adopting food additive provisions in the GSFA, recognizing that inconsistencies would exist between the GSFA and commodity standards until the General Standard would be finalized; (ii) the food additive provisions in Annex 1 of CX/EXEC 09/59/2 be forwarded by the Codex Committee on Food Additive (CCFA) to active commodity committees as appropriate; and (iii) the CCFA give the highest priority to the completion of the GSFA.¹⁰

45. In view of the above decision and of the difficulty of the CCFA to simultaneously work on the completion of the GSFA and address the food additive provisions in the commodity standards, the Commission agreed to suspend the following decision taken at its 29th Session:

*When provision for additives for inclusion into the GSFA result in amendments to additive provisions in Codex standards, consequential amendments should be made to the relevant standards and that the report of CCFAC should include a table showing the existing food additive provisions in Codex standards.*¹¹

46. The Commission **adopted** the draft and proposed draft food additive provisions of the GSFA as proposed by the 39th Session of the CCFA, noting the reservation of the Delegation of Cuba with regard to the provisions for alitame (INS 956) in food category 01.1.2 “Dairy-based drinks, flavoured and/or fermented (e.g. chocolate milk, cocoa, egnog, drinking yoghurt, whey-based drink)”.

*Proposed Draft Amendments to the International Numbering (INS) System for Food Additives*¹²

47. The Commission noted that a number of amendments to the INS system adopted at the current session should be reflected in the food additive provisions of existing commodity standards and requested the Codex Secretariat to revise the INS numbers in commodity standards accordingly.

Fish and Fishery Products

*Draft Code of Practice for Fish and Fishery Products (Sections on Quick Frozen Coated Products, Salted Fish and relevant Definitions)*¹³

48. The Commission **adopted** the Draft Sections with the amendments proposed by the Committee on Food Hygiene to Sections 10.4 and 10.5 Processing Operations for Molluscan Shellfish and Coated Shrimp, respectively, and to Section 11.4.4 Dry Salting and Section 11.5.3 Weighing, Wrapping and Packaging.

Fresh Fruits and Vegetables

*Draft Standard for Table Grapes and its proposed draft Sections 2.1.2 - Maturity Requirements and 3.1 - Minimum Bunch Weight*¹⁴

49. The Delegation of the United States of America, while not opposing the adoption of the Standard, expressed its reservation on the provisions for maturity requirements, stating that the uniform approach applied to Section 2.1.2 and the values therein did not reflect maturity requirements for all table grape varieties and producing regions in the world.

⁹ ALINORM 07/30/12 Rev, Appendix VII

¹⁰ ALINORM 07/30/3, para. 8

¹¹ ALINORM 06/29/41, para. 42, point (ii)

¹² ALINORM 07/30/12 Rev, Appendix XIII

¹³ ALINORM 07/30/18, Appendix II

¹⁴ ALINORM 07/30/35, Appendices IV and V

50. The Commission **adopted** the draft Standard for Table Grapes at Step 8 and the proposed draft Sections 2.1.2 - Maturity Requirements and 3.1- Minimum Bunch Weight at Step 5/8, with the omission of Steps 6 and 7, for inclusion in the Standard.

Food Hygiene

Draft Guidelines on the Application of General Principles of Food Hygiene to the Control of Listeria monocytogenes in Foods¹⁵

51. The Commission **adopted** the Draft Guidelines on the Application of General Principles of Food Hygiene to the Control of *Listeria monocytogenes* in Foods with amendments, as proposed by Brazil and Guatemala in their written comments, to reinsert “ready-to-eat foods” in the title and to insert a reference to the *General Standard for the Labelling of Prepackaged Foods* in Section 9.3.

Food Import and Export Inspection and Certification Systems

Proposed Draft Revision to the Guidelines for Generic Official Certificate Formats and Design, Production, Issuance and Use of Certificates¹⁶

52. The Delegation of Australia, speaking as the Chairperson of the Codex Committee on Food Import and Export Inspection and Certification Systems, stated that, after a careful review of the written comments submitted, the following amendments could be taken on board with a view to improving the document:

- Amend the title to read: “Guidelines for Design, Production, Issuance and Use of Generic Official Certificates”, in order to better reflect the content of the guidelines.

- Amend the first sentence of paragraph 18 as follows to avoid translation issues with the term “trade sample”:

“A consignment consisting of a food sample intended for evaluation, testing or research in the importing country ~~may be described using a term such as “trade sample”~~ should be clearly identified according to its intended use.”

- Amend the beginning of paragraph 34 for clarification as follows:

“Where, in exceptional cases justified by ~~immediate~~ a documented public health problem ~~concern~~, [...]”.

53. The Commission noted that additionally several editorial corrections to the Spanish version would be made in the final version of the text.

54. The Commission **adopted** the Proposed Draft Revision to the Guidelines at Steps 5/8, with the omission of Steps 6 and 7, with the amendments indicated above.

Fats and Oils

Draft Standard for Fat Spreads and Blended Spreads¹⁷

55. The Commission **adopted** the Draft Standard with the amendments made in the endorsement process by the Committee on Food Additives and the Committee on Methods of Analysis and Sampling. In particular, the Commission recommended that the Committee on Fats and Oils reconsider the levels of annatto extracts in fat spreads and other relevant products in existing standards to take into account the new ADIs established by JECFA.

¹⁵ ALINORM 07/30/13, Appendix III

¹⁶ ALINORM 07/30/30, para. 39 and Appendix II

¹⁷ ALINORM 07/30/17, Appendix II

General Principles

*Proposed Draft Working Principles for Risk Analysis for Food Safety for Application by Governments*¹⁸

56. The Commission briefly recalled the history of the proposed draft document which had begun as a general document on the application of risk analysis principles and had then been split into a document on *Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius*, which was subsequently adopted by the Commission at its 26th session (2003), and the proposed draft principles intended for application by governments, which had been discussed for several sessions in the Committee on General Principles (CCGP).

57. Several delegations stated that while not being opposed to the adoption of the text at Step 5, they had reservations on adopting it at Steps 5/8. They felt that while significant progress had been made in a physical working group and in the last session of the CCGP, not all comments had been taken into account and there was still room for improving the text. Some delegations, referring to the 59th Session of the Executive Committee,¹⁹ felt that the document should be circulated for comments at Step 6 to allow for more time before its finalization.

58. The Delegations of Chile, Mexico, Paraguay and South Africa expressed their reservation with respect to the procedure adopted in the CCGP, whereby comments on the proposals from the working group were not accepted, which was not consistent with the Guidelines on Physical Working Groups.

59. Many delegations and one Observer supported adoption of the Proposed Draft Principles at Steps 5/8. They expressed the view that the text was the fruit of long-lasting work in the CCGP and contained important guidance for governments to be made available to Codex members without delay. Some delegations noted that Codex was the only organisation among the “three sisters” under the WTO SPS agreement that had not as yet adopted such principles.

60. The Commission **adopted** the Proposed Draft Principles at Steps 5/8, with the omission of Steps 6 and 7. The Commission noted the reservations of the delegations of Costa Rica, Mexico, Paraguay and Thailand, who requested another round of discussion on this document before final adoption, on the decision to omit Steps 6 and 7, as well as the reservations of Argentina, Costa Rica, Paraguay and Thailand with the respect to the inclusion of the first sentence of paragraph 12 of the Proposed Draft Principles.

Coordinating Committee for Near East

*Draft Regional Standards for Canned Humus with Tehena, for Foul Medames and for Tehena*²⁰

61. The Commission **adopted** all three Standards at Step 8, with the amendments proposed by Lebanon improving the clarity of the texts, especially in Arabic, with the understanding that their sections on methods of analysis and sampling would be considered by the Commission after their endorsement by the Committee on Methods of Analysis and Sampling.

Nutrition and Foods for Special Dietary Uses

*Draft Revised Standard for Infant Formula and Formula for Special Medical Purposes Intended for Infants*²¹

62. The Delegation of Côte d'Ivoire indicated that the reference to good hygienic practice and other provisions in Section 9.5 “Information for Use” did not provide sufficiently detailed guidance for safe preparation and use of powdered infant formula and proposed to reference the recently published WHO/FAO *Guidelines on the Safe Preparation, Storage and Handling of Powdered Infant Formula*, which provided useful information for customers on safe handling and use of powdered infant formula and which had not

¹⁸ ALINORM 07/30/33 para. 89 and Appendix VIII

¹⁹ ALINORM 07/30/3, para. 13

²⁰ ALINORM 07/30/40, Appendices II, III and IV

²¹ ALINORM 07/30/26, Appendix II; ALINORM 07/30/3 para. 15

been available during finalization of the Standard by the Committee on Nutrition and Foods for Special Dietary Uses. The Representative of WHO proposed, instead, to make a reference, in the Section on Hygiene, to the WHO/FAO *Guidelines* in order to better protect infants. These proposals were supported by the Delegation of Singapore and several other delegations and observers.

63. Several other delegations and one observer pointed out that the standard had been under revision during a very long period of time, that the current text had been agreed by consensus at the last session of the CCNFSDU and that introduction of the reference to the WHO/FAO Guidelines might open some technical issues which should be dealt with by the Committee. It was noted that the Committee on Food Hygiene has been revising the Code of Hygienic Practice for Powdered Infant Formula for Infants and Young Children and that the WHO/FAO Guidelines could be fully taken into account in the revision of the Code.

64. After some discussion, the Commission agreed to **adopt** the Standard at Step 8 as proposed by the CCNFSDU with some editorial changes in Spanish and deletion of the reference to arginine in Annex I containing the list of essential amino acids, with the understanding that the section on methods of analysis would be reviewed by the next session of the CCNFSDU with a view to its submission to CCMAS for endorsement.

65. The Commission also **agreed** to request the Committee on Food Hygiene to take into account the WHO/FAO *Guidelines on the Safe Preparation, Storage and Handling of Powdered Infant Formula* when revising the Code of Hygienic Practice for Powdered Infant formula for Infants and Young Children and to invite the CCNFSDU to review the Section on Hygiene of the adopted Standard once the above Code had been finalized by the CCFH.

Processed Fruits and Vegetables

*Draft Standard for Processed Tomato Concentrates*²²

66. The Delegation of Cuba expressed concern on the technological justification for the inclusion of citric acid as an acidity regulator for this commodity. The Delegation indicated that the current Standard for Processed Tomato Concentrates (CODEX STAN 57-1981) did not contemplate the use of this additive whose addition might create a technical barrier to trade. The Delegation of the United States, speaking on behalf of the Chairperson of the Committee on Processed Fruits and Vegetables, informed the Commission that food additive provisions, including citric acid in processed tomato concentrates, had been discussed and agreed upon by the Committee and subsequently endorsed by the Committee on Food Additives.

67. The Commission **adopted** the draft Standard for Processed Tomato Concentrates at Step 8 as proposed by the Committee. The Delegations of Cuba and Egypt reserved their position on this decision of the Commission.

Pesticide Residues

*Draft and Proposed Draft Maximum Residue Limits for Pesticides*²³

68. In reply to the strong opposition expressed by the European Community and Norway on Indoxacarb (216), the WHO Secretariat of the FAO/WHO Joint Meetings on Pesticide Residues (JMPR) clarified that their concern had been considered by the JMPR in 2006. After detailed review of the relevant toxicological studies, the JMPR confirmed its previous opinion and a detailed analysis of this matter was presented at the 39th Session of the Committee, which accepted the opinion and assessment of the JMPR and recommended the advancement of the draft MRLs for adoption at Step 8.

69. The Commission **adopted** the MRLs as proposed in Appendices II and III of ALINORM 07/30/24 with the addition of the explanatory note for exclusion for the MRLs for Boscalid (221), which had inadvertently been omitted from the report of the Committee, and noted the reservation expressed by the European Community and Norway on MRLs for Endosulfan (32), Pirimicarb (101), Propamocarb (148), Fenpropathrin (185) and Pyraclostrobin (210) as presented in CAC/30 LIM/7.

²² ALINORM 07/30/27, Appendix III

²³ ALINORM 07/30/24, Appendices II and III

Methods of Analysis and Sampling

*Methods of Analysis in Codex Standards at different steps*²⁴

70. The Commission adopted the methods as proposed by the Committee on Methods of Analysis and Sampling. The Delegation of Brazil expressed its reservation on the methods of analysis for inclusion in the four Standards developed by the Committee on Processed Fruits and Vegetables and adopted at the current session and on the update of the methods in current standards for fats and oils, as mentioned in their written comments in ALINORM 07/30/5A.

Standards and related texts held at the Commission at Step 8

*Draft MRLs for Bovine Somatotropin*²⁵

71. Some delegations noted that the draft MRLs for bovine somatotropin (BST) had been held at Step 8 since the 23rd Session of the Commission in 1999 and expressed the view that the Commission needed to take action to finalise this matter in the near future. The Delegation of Chile requested that the status and procedures be defined for standards retained at Step 8.

72. The Commission noted that no request had been received to change the status of the draft MRLs for BST and therefore agreed to retain them at Step 8.

*Draft Revised Standards for: Cheddar (C-1); Danbo (C-3); Edam (C-4); Gouda (C-5); Havarti (C-6); Samsø (C-7); Tilsiter (C-11); Saint-Paulin (C-13); Provolone (C-15); Cottage Cheese (C-16); Coulommiers (C-18); Cream Cheese (C-31); Camembert (C-33); Brie (C-34); and draft Standard for Mozzarella*²⁶

73. The Commission recalled that its 29th Session had agreed to hold 16 individual cheese standards at Step 8 due to non-endorsement of their labelling provisions²⁷ and that the 35th Session of the Committee on Food Labelling had subsequently endorsed all provisions in section 7.2 “Country of Origin” as originally proposed by the Committee on Milk and Milk Products.²⁸ The Commission further recalled that the 38th Session of the Committee on Food Additives and Contaminants had not endorsed the food additive provisions for annatto extracts (INS 160b) and gluconic acids (INS 574).²⁹

74. The Commission agreed to adopt the 15 draft standards above mentioned with the amendments made in the endorsement process by the Committee on Food Additives and Contaminants. Furthermore, the Commission recommended that the Committee on Milk and Milk Products reconsider the levels of annatto extracts in individual cheese standards and in existing standards for milk products to take into account the new ADIs established by JECFA, as recommended by the Executive Committee in the framework of the Critical Review³⁰.

*Draft Revised Standard for Emmental (C-9)*³¹

75. The Delegation of Switzerland, referring to its written comments in CAC/30 LIM/7, recalled that it had always been opposed to the deletion of the reference to Switzerland as the historical country of origin of Emmental cheese in the Draft Revised Standard. The Delegation stressed that the omission of the reference to Switzerland as the historical country of origin would mislead or deceive the consumers especially since Switzerland had been recognised as the historical country of origin in the current Codex Emmental Standard since 1967. In addition, the Delegation of Switzerland opposed the establishment of a link to the question of

²⁴ ALINORM 07/30/23, Appendix III

²⁵ ALINORM 95/31, Appendix II

²⁶ ALINORM 06/29/11, para. 85 and Appendices VI, VII, IX-XII, XIV-XXII

²⁷ ALINORM 06/29/41, para. 88 and Appendix VI

²⁸ ALINORM 07/30/22, para. 16

²⁹ ALINORM 06/29/12, paras 40 and 44 and Appendix IV

³⁰ ALINORM 07/30/3, para. 16

³¹ ALINORM 06/29/11, para. 85 and Appendix XIII

unascertained goods and reiterated the opinion that individual cheese standards should be replaced by more general health-based group standards in accordance with the recommendations of the Codex *Evaluation* and the Strategic Framework. The Delegation expressed its strong opposition to the adoption of the Draft Revised Standard, unless a footnote in section 7.2 was added to indicate the country (Switzerland) in which the name “Emmental” had historically originated and the labelling provisions were dealt with separately from questions regarding unascertained goods and that no mention with regard to the nature of the names of the cheeses concerned was made or accepted.

76. Other delegations supported the adoption of the draft revised standard as submitted by the Committee on Milk and Milk Products and noted that Emmental cheese was produced all over the world and its name had become generic. It was also noted that the standard had been considered by the Committee on Milk and Milk Products on the basis of the same criteria used for the other standards for individual cheeses.

77. On the basis of the above debate, the Chairperson closed the discussion and concluded that the revised Standard for Emmental had been adopted, with the amendments made in the endorsement process by the Committee on Food Additives and Contaminants, and the strong opposition of the Delegation of Switzerland recorded in the report.

78. The Delegation of Switzerland challenged the Chairperson’s ruling and, in accordance with the General Rules of FAO, for instance Rules IX.4 and XII.16, they submitted an alternative proposal, seconded by the Delegation of Jamaica, to hold the Draft Revised Standard at Step 8 and to refer its labelling provisions in section 7.2 “Country of Origin” back to the Committee on Food Labelling for further discussion. According to the above mentioned Rules a vote was conducted on the alternative proposal made by Switzerland. The motion was put to a roll-call vote, as opposed to a show of hands, at the request of the Delegation of Switzerland, with the following results.

Votes in favour: Algeria, Angola, Armenia, Benin, Bhutan, Bulgaria, Croatia, Cyprus, Islamic Republic of Iran, Iraq, Jamaica, Lao People’s Democratic Republic, Niger, Papua New Guinea, Qatar, Romania, Rwanda, Senegal, Serbia, Sudan, Switzerland, Syrian Arab Republic, Turkey

Votes against: Argentina, Australia, Austria, Belgium, Botswana, Brazil, Burundi, Cameroon, Canada, Chile, China, Colombia, Costa Rica, Côte d’Ivoire, Cuba, Democratic Republic of Congo, Denmark, Dominican Republic, Ecuador, Egypt, Estonia, Ethiopia, Fiji, Finland, France, Germany, Greece, Guatemala, Guinea, Hungary, India, Indonesia, Japan, Kenya, Kuwait, Latvia, Lithuania, Malawi, Malaysia, Mali, Malta, Mexico, Netherlands, New Zealand, Nigeria, Norway, Oman, Paraguay, Peru, Poland, Portugal, Republic of Korea, Saudi Arabia, Singapore, Slovakia, Slovenia, South Africa, Spain, Sweden, Tanzania, Tunisia, Uganda, United Arab Emirates, United Kingdom, United States of America, Vanuatu, Venezuela, Viet Nam, Zambia, Zimbabwe

Abstaining: Antigua and Barbuda, Czech Republic, Ghana, Iceland, Ireland, Jordan, Lebanon, Lesotho, Morocco, Panama, Thailand

Tally: 93 votes cast, 23 in favour, 70 against, 11 abstentions (majority required 47)

Result: The motion by Switzerland was not approved.

79. The ruling of the Chairperson that the Commission had adopted the revised Standard for Emmental, as per above, stood.

PROPOSED DRAFT STANDARDS AND RELATED TEXTS AT STEP 5 (Agenda Item 6)³²

80. The Commission **adopted** the Proposed Draft Standards and Related Texts at Step 5 submitted by its subsidiary bodies as presented in Appendix V to this report and advanced them to Step 6. The Commission noted that technical comments raised during the session would be referred to the relevant Committees for their consideration. The Commission encouraged members and observers that have submitted comments in writing or orally at the session to submit these comments at Step 6 of the Procedure.

81. The following paragraphs provide additional information on the comments made and the decisions taken on certain items.

Food Additives***Proposed Draft Guidelines for the Use of Flavourings (N03-2006) (with the exception of Section 4 and Annexes A and B)***³³

82. The Commission **adopted** the proposed draft Guidelines at Step 5 as proposed by the Committee and advanced them to Step 6, with the understanding that several translation issues in Spanish would be addressed by the next session of the Committee on Food Additives.

Coordinating Committee for Asia***Proposed Draft Standards for Gochujang and Ginseng Product***³⁴

83. The Commission agreed to defer the discussion on the proposed draft Standards for Gochujang and Ginseng Product until it had discussed the issues on the role of Coordinating Committees in developing regional standards and the conversion of regional standards into worldwide standards, as a general issue, under Agenda Item 12 (b). The Delegation of the Republic of Korea proposed to apply Proposal 8, as amended by the Executive Committee, to new work rather than to ongoing work.

84. Taking into account the outcome of the discussion under Agenda Item 12 (b), based on the recommendation of the 59th Session of the Executive Committee³⁵, the Commission **adopted** the proposed draft Standards at Step 5 as Draft Regional Standards for further elaboration by the CCASIA with a view to finalization as regional standards. The Delegation of the United States, speaking as the Vice-Chair of the Commission, reminded the Commission of the Executive Committee's commitment to develop policies concerning regional standards including their conversion into world-wide standards. The Delegation of the Republic of Korea stated that the conversion of these regional standards into world-wide standards should be actively considered after adoption at Step 8.

³² ALINORM 07/30/6, ALINORM 06/29/6A (comments of Japan (CCFA), International Alliance of Dietary/Food Supplement Associations (CCASIA), European Community, Peru (CCFFP), Argentina, Australia, Mexico, New Zealand, United States of America, International Special Dietary Food Industry (CCNFSDU), Argentina (CCFFV), International Hydrolyzed Protein Council (CCCF), Brazil (CCPFV), CAC/30 LIM-4 (comments of Guatemala, Malaysia (CCFL), South Africa (CCNFSDU), Malaysia, Peru (CCPFV), Malaysia (CCASIA), CAC/30 LIM-8 (comments of Indonesia (CCFA), Indonesia (CCASIA), Indonesia, Peru, Philippines (CCFL), European Community, Indonesia (CCPR), Indonesia (CCNFSDU, CCPFV, CCFFV), Indonesia, Philippines (CCCF), CAC/30 LIM-18 (comments of Honduras)

³³ ALINORM 07/30/12 Rev, Appendix XI

³⁴ ALINORM 07/30/15, Appendices II and III

³⁵ ALINORM 07/30/3 para. 114

Fish and Fishery Products

*Proposed Draft Standard for Live and Raw Bivalve Molluscs*³⁶

85. The Commission endorsed the recommendation of the Executive Committee that the Committee on Fish and Fishery Products consider the questions from the Committee on Food Hygiene in the development of the standard and consider the need for further scientific advice on biotoxins and **adopted** the draft Standard at Step 5 and advanced it to Step 6.

Committee on Food Labelling

*Proposed Draft Amendment to the Guidelines for Organically Produced Foods (Ethylene)*³⁷

86. Some delegations, while not objecting in principle to the use of ethylene, expressed the view that this substance should be used only in accordance with good agricultural practices and that all required conditions for use should be met in order to ensure the safety and quality of the products treated with ethylene.

87. The Commission agreed that these comments should be taken into account in the finalisation of the amendment on the inclusion of ethylene.

*Proposed Draft Amendment to the General Standard for the Labelling of Prepackaged Foods (Quantitative Declaration of Ingredients)*³⁸

88. The Delegation of Norway, while not objecting to the adoption of the text, expressed concern with the deletion of the provisions for added sugars in view of their importance in the perspective of the WHO Global Strategy for Diet, Physical Activity and Health and proposed that the purpose of these provisions should not be lost and should be considered further by the Committee in relation with the implementation of the Global Strategy.

*Proposed Draft Definition of Advertising in relation to health and nutrition claims*³⁹

89. The Delegation of China expressed the view that advertising should not be defined in the framework of Codex but should be left to national authorities.

90. The Commission recommended that the Committee on Food Labelling clarify in which text the definition should be included when finalised.

Pesticide Residues

*Proposed Draft Maximum Residue Limits for Pesticides*⁴⁰

91. The Commission **adopted** the draft MRLs as proposed in Appendices IV of ALINORM 07/30/24 at Step 5 and advanced them to Step 6, noting the reservations expressed by the European Community and Norway on the MRLs for Endosulfan (32). The Commission noted that the reference to “marine mammals” appearing under Thiabendazole (65) was an editorial error and should be deleted.

Fresh Fruits and Vegetables

*Proposed Draft Standard for Bitter Cassava*⁴¹

92. The Commission concurred with the recommendation of the Executive Committee to adopt the Standard at Step 5 and that, as a separate issue, the Committee on Contaminants in Foods should consider the

³⁶ ALINORM 07/30/18 Appendix V, ALINORM 07/30/3 paras 25 – 27, ALINORM 07/30/13 paras 219 – 223

³⁷ ALINORM 07/30/22 Appendix IV

³⁸ ALINORM 07/30/22 Appendix V

³⁹ ALINORM 07/30/22 Appendix VI

⁴⁰ ALINORM 03/30/24, Appendix IV

⁴¹ ALINORM 07/30/ 35 Appendix VI, ALINORM 07/30/3, paras 28 - 30

safety levels of hydrogen cyanide proposed in the Standard, with a view to a re-evaluation of cyanogenic glycosides by JECFA.

Proposed Draft Guidelines for the Inspection and Certification of Fresh Fruits and Vegetables for Conformity to Quality Standards⁴²

93. In adopting the proposed draft Standard at Step 5, the Commission recognized that the Guidelines addressed inspection, certification and sampling matters specific to fresh fruits and vegetables and therefore, agreed to forward the document to the Committees on Food Import and Export Inspection and Certification Systems as well as on Methods of Analysis and Sampling to provide observations from a horizontal perspective on certification, inspection and sampling provisions in order to ensure consistency in the approach followed on these matters within Codex.

Contaminants in Foods

Proposed Draft Maximum Levels for 3-MCPD in Liquid Condiments containing Acid-Hydrolyzed Vegetable Proteins (excluding Naturally Fermented Soy Sauce) (N08-2004)⁴³

94. The Delegations of the European Community and Norway expressed their reservation on the decision of the Commission to adopt the proposed draft Maximum Levels of 0.4 mg/kg for 3-MCPD at Step 5, stating that the maximum level for 3-MCPD should be set as low as reasonably achievable in view of possible public health concerns, that the levels lower than 0.4 mg /kg were achievable by implementing good manufacturing practices and that lower maximum levels should be considered in the light of the finalization and implementation of the Code of Practice for the Reduction of 3-MCPD during the Production of Acid-Hydrolyzed Vegetable Proteins (Acid-HVPs) and Products that contains Acid-HVPs.

REVOCATION OF EXISTING CODEX STANDARDS AND RELATED TEXTS (Agenda Item 7)⁴⁴

95. The Commission **approved** the revocation from the *Codex Alimentarius* of previously adopted texts as summarized in Appendix VI to this report.

PROPOSALS FOR THE ELABORATION OF NEW STANDARDS AND RELATED TEXTS AND FOR THE DISCONTINUATION OF WORK (Agenda Item 8)⁴⁵

ELABORATION OF NEW STANDARDS AND RELATED TEXTS

96. The Commission **approved** the elaboration of new standards and related texts as summarized in Appendix VII to this report. The following paragraphs provide additional information on the comments made and the decisions taken on certain items.

97. The Commission noted that project documents submitted to the 59th Session of the Executive Committee contained information that, while respecting the overall format as set out in the Procedural Manual, varied significantly in terms of quantity and quality, and therefore, endorsed the recommendation of the Executive Committee to encourage Codex committees, task forces and Codex Members to prepare future project documents according to the format set out in the current revision of the Procedural Manual and provide sufficiently detailed, relevant information with particular regard to the evidence-based assessment against each of all the Criteria for the Establishment of Work Priorities⁴⁶.

⁴² ALINORM 07/30/35 Appendix VII

⁴³ ALINORM 07/30/41 Appendix X

⁴⁴ ALINORM 07/30/7; CAC/30 LIM/10 (comments of Sri Lanka)

⁴⁵ ALINORM 07/30/8, CAC/30 LIM/6 (comments of Ghana, Malaysia and Philippines), CAC/30 LIM/10 (comments of Japan and Malaysia), CAC/30 LIM/13 (comments of Indonesia) and CAC/30 LIM/15 (comments of Kenya)

⁴⁶ ALINORM 07/30/3, para 46

Fish and Fishery Products

Revision of the Procedure for the Inclusion of Additional Species in Standards for Fish and Fishery Products

98. The Delegation of Morocco, supported by the Delegation of Chile, noting the status of the document which was for internal use by the Committee on Fish and Fishery Products, stated that there was a need for more transparency with regard to the procedures used by this Committee.

99. In view of the above, the Commission recommended that the Committee consider, upon finalisation of the document, its inclusion in the Procedural Manual and subsequent publication on the Codex website in order to enhance transparency of the processes used by the Committee.

Standard for Fresh/Live and Frozen Abalone (*Haliotis* spp.)⁴⁷

100. The Commission, while approving the proposal for new work on the revision of the Standard for Fresh/Live and Frozen Abalone (*Haliotis* spp) endorsed the recommendation of the Executive Committee that the Committee on Fish and Fishery Products consider broadening the scope of the standard to include other gastropods.

Processed Fruits and Vegetables

Sampling Plan Including Methodological Provisions for Controlling Minimum Drained Weight of Canned Fruits and Vegetables⁴⁸

101. The Commission agreed with the recommendation of the Executive Committee to amend the title by referring to canned fruits and vegetables “in packing media” as there are other canned fruits and vegetables not requiring provisions for minimum drained weight.

102. The Delegation of the United States, while not opposing to the elaboration of the proposed document, questioned the need for developing a separate sampling plan for minimum drained weight as the current, simple provisions in the relevant standards for processed fruits and vegetables had not created problems in international trade and the proposed work did not substantially advance the Codex objectives in terms of the protection of consumers’ health and fair trade practices. The Delegation stressed the need to consider carefully priorities amongst proposals for new work to be undertaken by subsidiary bodies of the Commission in view of their workload and resource limitations.

Coordinating Committee for Asia⁴⁹

Standard for Chili Sauce

103. Many delegations from Asia, referring to the recommendation of the 59th Session of the Executive Committee, supported the elaboration of a standard for chili sauce with a view to its finalization as a regional standard by the CCASIA. Its conversion into a worldwide standard could be considered after adoption at Step 8. Some other delegations from the region and from outside the region did not support the proposal because in their countries chili sauce referred to tomato-based products, not to chili pepper-based products only as suggested in the Project Document, and stated that the establishment of a Codex Standard for Chili Sauce, whether regional or worldwide, could have adverse impact on fair trade of the related products.

104. Some delegations from outside the region expressed the view that, depending on the scope to be defined for this standard, it could refer to products which were widely traded in other regions and therefore they could be interested in participating in this work. The Delegation of Mexico informed the Commission of its intention to propose new work on chili peppers at the next meeting of the Committee on Fresh Fruits and Vegetables.

⁴⁷ ALINORM 07/30/3 para. 34

⁴⁸ ALINORM 07/30/3, para 35

⁴⁹ ALINORM 07/30/3, paras 40-41

105. Given the support by the majority of the Members from Asia, the Commission **approved** the proposal for new work to elaborate a Regional Standard for Chili Sauce by the CCASIA and agreed to encourage the Committee, in undertaking the work, to take into account the comments made at the present session and seek comments and information from members belonging to other regions. The Commission also recommended that the Committee on Processed Fruits and Vegetables be informed of the status of work in CCASIA at its next session and be invited to provide its views on the need for an international standard for chili sauce. The Delegation of the Republic of Korea reserved its position on this decision of the Commission, stating that the standard should be developed as a worldwide standard.

Standard for Edible Sago Flour

106. The Delegation of Japan, while not opposing the proposal for new work, suggested that the scope of the standard should exclude sago starch, whose manufacturing process and quality factors were quite different from sago flour.

107. The Commission **approved** the elaboration of a Regional Standard for Edible Sago Flour by CCASIA.

Foods Derived from Biotechnology

Annex to the Guideline for the Conduct of Food Safety Assessment of Foods Derived From Recombinant-DNA Plants on Low-level Presence of Recombinant-DNA Plant Material

108. The Delegation of the European Community, supporting the new work, stressed that the proposed annex must be developed in parallel with the establishment of a mechanism for data sharing and information exchange to support the actions of food control authorities in situations of low-level presence of unauthorized recombinant-DNA plant materials and urged FAO to take a lead to coordinate with other relevant international organizations and stakeholders with a view to facilitating construction of such mechanism.

109. The Representative of FAO, speaking on behalf of FAO and WHO, informed the Commission of the ongoing work for the establishment of a database to be developed within FAO as part of the International Portal on Food Safety, Animal and Plant Health, and stated that the Codex Task Force on Food Derived from Biotechnology would be kept informed of the progress made on this matter.

Food Hygiene

Guidelines for the Control of Campylobacter and Salmonella spp. in Broiler (young bird) Chicken Meat⁵⁰

110. The Commission agreed to the recommendation of the Executive Committee that the scope of the new work be expanded to cover chicken meat in general, thereby deleting reference to “broiler (young bird)” in the title, and that the Committee for Food Hygiene be invited to consider re-scoping the document, as appropriate, taking into account all relevant factors including the availability of risk assessments.

111. The Commission noted that this decision would impact on the proposed work plan for the new work and might require a longer time-frame for the completion of the guidelines since the guidelines would follow a novel farm-to-fork approach based on quantitative risk assessment to the widest extent practicable; that there existed considerable scientific data and a risk assessment by JEMRA for broiler chickens but not for non-broiler chickens with different risk profiles, production and processing conditions; that a new worldwide call for scientific data for the latter category of chicken meat might be necessary before a risk assessment be conducted by JEMRA.

112. The Commission further noted that the OIE would undertake work on *Salmonella* and *Campylobacter* in broiler chickens that could contribute to this new work in the Committee on Food Hygiene.

⁵⁰ ALINORM 07/30/3 paras 43-45

DISCONTINUATION OF WORK

Committee on Food Labelling⁵¹

Guidelines for the Production, Processing, Labelling and Marketing of Organically produced Foods Proposed Draft Revised Annex 2: Table 1 (Natural Sodium Nitrate)

113. The Delegation of Chile expressed the view that the consideration of this issue in the Committee on Food Labelling had not been consistent with the criteria applicable to substances for inclusion in the Guidelines as, although all relevant scientific justification had been provided, the use of Natural Sodium Nitrate had not been accepted, although other member countries did not provide written counter arguments. The Delegation also pointed out that other substances which were not based on similar scientific data had been included in the list of permitted substances.

FINANCIAL AND BUDGETARY MATTERS (Agenda Item 9)

114. The Commission noted the extensive discussions held at the 59th Session of the Executive Committee on the Codex Budget 2006-07 and the estimates of expenditure for 2008-09⁵², on alternative funding mechanisms and on the FAO/WHO budgets for scientific advice.

Codex Budget 2006-07 and estimated budget for 2008-09⁵³

115. The Commission noted the introduction by the Secretariat of document ALINORM 07/30/9 on the budget for the current biennium (2006-07) (Table 1) and estimates of expenditure for the biennium 2008-09 (Tables 2A and 2B). The estimates were based on the assumption of two sessions of the Commission, three sessions of the Executive Committee, one session of each Coordinating Committee, and maintaining the current cost saving measures concerning publication/document distribution. Different budgetary scenarios were envisaged pending the introduction of additional languages (Russian in CCEURO and CAC; Portuguese in CCAFRICA).

116. One delegation stated that the Commission should await the outcome of the FAO Conference in November 2007 before pronouncing its view on the budget 2008-09. The Commission noted the opinion of one delegation that four sessions of the Executive Committee should be held during the next biennium in order to allow it to adequately fulfil its standards management function.

117. The Commission recalled that FAO and WHO had significantly increased their contributions to the Codex programme between the 2002-03 and 2004-05 biennia but noted further that the main expenditure of the Codex Secretariat was in Euro and Swiss Francs, against which the US dollar had weakened significantly in the same period of time, resulting in loss of purchasing power. The Commission further noted that WHO had adopted its overall budget level for 2008-09 in May 2007 whereas FAO would discuss its budget proposals only at the FAO Conference in November 2007. The Commission welcomed the commitment of the parent organisations to secure the Codex budget. In reply to requests for WHO to increase its share in the Codex budget, the Representative of WHO stated that any requests for additional resources from WHO should be well justified by a business plan based on the objectives of the Strategic Plan. The Representative further stated that the increase of one budget (e.g. Codex) might lead to a decrease in another (e.g. scientific advice) and stressed the need for a clear prioritisation of Codex work as well as further investigation of relevant cost-saving measures.

118. The Commission noted that the overall cost of the Codex undertakings was significantly higher than the figures given in document ALINORM 07/30/9 due to the fact that host governments of Codex Committees and Task Forces also provided substantial contributions through the provision of resources such as venue, translation and interpretation services.

119. The Commission noted requests from delegations for increased transparency in the expenditure of the Codex programme and welcomed the information given by the Secretariat to the 59th Session of the

⁵¹ ALINORM 07/30/22, para. 92

⁵² ALINORM 07/29/3, paras 88-106

⁵³ ALINORM 07/30/9

Executive Committee on the staff structure of the Codex Secretariat and the breakdown of chargebacks⁵⁴. The Commission further noted that the capacity of the Codex Secretariat was stretched to its limits due to the increased workload related to the annual sessions of the Commission and the newly assigned functions of the Executive Committee such as the critical review and the assessment of the applications for observer status. The Commission also noted that the need for continuously monitoring and reporting the progress made in the implementation of the Strategic Plan 2008-2013 would lead to additional work.

120. The Commission noted proposals from delegations on the possibility to make savings by outsourcing translation and printing but recalled that as the Joint FAO/WHO Food Standards Programme was administered by FAO on behalf of FAO and WHO, the Codex Secretariat had to follow FAO rules and practices in the purchase of services and goods. Any proposal to change the current arrangement would have to be reviewed as an organization-wide management issue. The Representative of FAO stressed the need to identify further cost-saving measures, such as producing outcome-based, significantly shorter reports of Codex sessions. The Commission however noted concerns of some members that this might impact negatively on the transparency of the Codex standards setting process, especially for developing countries which could not participate in all Codex sessions. The Representative of WHO informed the Commission that WHO had now outsourced many administrative tasks including printing.

121. The Commission further noted concerns of some delegations on the late availability of Codex documentation in other languages than English and problems with the quality of translations. The Commission noted the information from the Secretariat that delays in translation were often due to the complexity and tightness of the Codex session schedule especially with annual Commission sessions, limited human resources of the Codex Secretariat and late submission of government comments.

122. Some members strongly supported the use of Portuguese in CCAFRICA as a language of interpretation but not for documentation.

123. In conclusion the Commission:

- **Noted** the Codex budget proposals for 2008-09 and **expressed** its strong desire that FAO and WHO allocate, to the Codex programme, a budget level that would allow the Secretariat to conserve, at least, the same purchasing power as in the 2006-07 biennium, calling upon Codex members to voice their support to Codex in the governing bodies of FAO and WHO;
- **Requested** FAO and WHO, pending the adoption of Russian by the FAO Conference as an FAO language, to increase the Codex budget to allow the use of Russian in the Commission and in the FAO/WHO Coordinating Committee for Europe; and
- **Requested** FAO and WHO to study the possibility to add Portuguese as a language of interpretation in the FAO/WHO Coordinating Committee for Africa in line with the policy applied at the FAO Regional Conference for Africa and the WHO Regional Committee for Africa.

Consideration of alternative funding mechanisms⁵⁵

124. The Commission recalled that at its 29th Session it had requested the Secretariat to prepare a discussion paper, in collaboration with FAO and WHO, on the possibilities for more sustainable funding including through other funding sources and alternative ways of achieving it⁵⁶.

125. The Commission noted the discussion held at the 59th Session of the Executive Committee on this matter⁵⁷ and endorsed the recommendation of the Executive Committee to consider the document at its 60th Session in December 2007.

FAO/WHO budgets for scientific advice⁵⁸

126. The Commission noted the information provided by FAO and WHO, including the strengthening of their scientific activity in the area of nutrition. The Commission **expressed** its desire that FAO and WHO

⁵⁴ ALINORM 07/30/3, para. 95, CRD 9 at the 59th Session of the Executive Committee

⁵⁵ ALINORM 07/30/9-Add.1

⁵⁶ ALINORM 06/29/41, para. 150

⁵⁷ ALINORM 07/29/3, paras 88-99

⁵⁸ CAC/30 INF/3

maintain an adequate budgetary level for the provision of scientific advice and welcomed the initiative of FAO and WHO to develop adequate funding strategies for these activities supporting Codex. The Commission was informed that the Global Initiative for Food Related Scientific Advice (GIFSA), was launched at a side-event of this Commission session, allowing FAO and WHO to collect extrabudgetary contributions from members and civil society.

PROPOSED SCHEDULE OF CODEX SESSIONS (Agenda Item 10)⁵⁹

127. The Commission considered the tentative schedule that had been prepared on the basis of the information provided by host countries of Codex Committees and Task Forces. The Commission noted the changes in the dates and venues proposed for some sessions.

128. The Delegation of Mexico proposed to retain the current five-day meeting of the Coordinating Committee for Latin America and the Caribbean instead of the four-day meeting mentioned in the proposed Schedule. The Secretariat pointed out that this duration was consistent with the schedule of other Coordinating Committees, and drew the attention of the Commission to the financial implications of longer sessions in the case of Coordinating Committees as interpretation and translation were funded by the Codex budget.

129. The Delegation of Argentina stressed the importance of Coordinating Committees in order to allow developing countries to participate in Codex and therefore expressed the view that the current five days meeting should be retained so as to allow the Committee to consider all issues of importance to countries in the region.

130. The Commission noted that the final dates and venues of Codex sessions were determined by the Directors-General of FAO and WHO, in consultation with the host country, and would be indicated in formal invitations. The Commission endorsed the Proposed Schedule in principle with the above mentioned changes and noted that some further changes might be made to the schedule in the future.

STRATEGIC PLANNING OF THE CODEX ALIMENTARIUS COMMISSION⁶⁰ (Agenda Item 11)

131. The Commission considered the revised draft Strategic Plan 2008 – 2013 and the relevant discussion in the report of the 59th Session of the Executive Committee (ALINORM 07/30/3), which conducted a final revision on the basis of comments forwarded by the six FAO/WHO Coordinating Committees as presented in document ALINORM 07/30/9B. The Commission noted the observations made, and agreed to some amendments as follows.

132. The Commission agreed that the term “risk-based” be replaced with “based on risk” in the first bullet point of paragraph 6, Part 1 and in Activity 1.1 of Part 2 in the English text, on the basis of a proposal from Brazil, in order to ensure consistency with the language that was already used in the Codex Alimentarius.

133. The Commission agreed to delete Activity 2.6 in Part 2 and in Table 1, Part 3 regarding the elaboration of working principles for risk analysis for food safety for application by governments, taking note of the adoption of the “Working Principles for Risk Analysis for Food Safety for Application by Governments” at its current Session (see Appendix III). The Commission, while noting the proposal from the Delegation of Cameroon to replace activity 2.6 with another activity to develop indicators to monitor the implementation of the Working Principles, agreed with the proposal of the Delegation of New Zealand to replace activity 2.6 with an alternative activity aimed at facilitating the implementation and application of the Working Principles at the national level. The Commission therefore agreed to add a new Activity 2.6 in Part 2 and in Table 1, Part 3, on the basis of a proposal by the Delegation of New Zealand (LIM 21) with a minor modification, noting the important role of FAO, WHO and Codex Members in the provision of technical assistance. The Commission noted the proposal of the delegation of Chile to include under activity 4.4. of the Strategic Plan the activities on cooperation with the WTO, as they were included in the Goals but not in the activities, however decided not to make changes in this regard.

⁵⁹ ALINORM 07/30/9A

⁶⁰ ALINORM 07/30/3 paras 72 - 87 and Appendix II, ALINORM 07/30/9B, CAC/30 LIM/21 (Proposed text from New Zealand)

134. The Delegation of Australia, supporting the adoption of the draft Strategic Plan, suggested that the Executive Committee could consider, probably at its next session, the elaboration of a biennial business plan based on the priority of work identified in the Strategic Plan, along with corresponding estimates of funding and expenditure to cover the activities foreseen in 2010 - 2011. This business plan could also be used as a basis for future requests for increased funding. The Delegation noted that this approach was already used by the International Plant Protection Convention (IPPC).

135. In regard to the concern expressed by the Delegation of Mexico, on the need to strengthen scientific advisory bodies particularly in the area of pesticide residues, the Commission, recalling the extensive discussion at the 59th Session of the Executive Committee on this issue, agreed that there was no need to amend the text of the draft Strategic Plan, noted that it was necessary to explore possible solutions to mitigate or eliminate concerns of developing countries on this matter and therefore suggested that FAO and WHO organize a tailor-made action-oriented workshop to address some of these concerns of developing countries on pesticide residue issues. Such a workshop would facilitate the understanding of the impact of the current processes followed by JMPR and CCPR in developing MRLs for pesticides and the identification of the need of developing countries.

136. The Representative of FAO stated that FAO, WHO and IAEA continued to provide capacity building activities to assist member countries in generating data as well as controlling and monitoring pesticide residues to ensure food safety. The Representative expressed their willingness to continue efforts to respond to greater needs of developing countries on this issue.

137. The Delegation of Argentina, supporting the view expressed by Mexico, suggested that an international FAO/WHO conference with all interested parties, that would take into account the concerns of developing countries, be held to explore an alternative mechanism for data generation and collection with a view to facilitating the provision of scientific advice on pesticide residues to Codex.

Status of the Draft Strategic Plan 2008-2013

138. The Commission adopted its Strategic Plan 2008 – 2013 as contained in Appendix IX to this report.

IMPLEMENTATION OF THE JOINT FAO/WHO EVALUATION OF THE CODEX ALIMENTARIUS AND OTHER FAO AND WHO WORK ON FOOD STANDARDS (Agenda Item 12)

GENERAL IMPLEMENTATION STATUS (Agenda Item 12a)⁶¹

139. The Commission noted with satisfaction the status of implementation of the proposals as presented in Tables 1 and 2 of document ALINORM 07/30/9C. Five years after the Joint FAO/WHO Evaluation of the Codex Alimentarius and other FAO and WHO Work on Food Standards, nearly all proposals originating from the recommendations of the Joint Evaluation and endorsed by the 26th Session of the Commission had been implemented and no further action required at this juncture, except the two recommendations below which were being experimented at the committee level:

- Proposal No. 19 “Use of facilitators”; and
- Proposals No. 32 “Co-chairmanship”

140. With regard to Proposal No. 12 “Participation of observers in the Executive Committee”, the Commission noted that recent sessions of the Executive Committee had been audio-recorded and the audio-recording posted on the Codex website on an experimental basis. In view of the positive outcome of this arrangement which allowed publicizing of the proceedings of the Executive Committee, the Commission recommended that the current arrangements for audio-recording and web-posting be implemented on an ongoing basis.

141. Regarding Proposal No. 34 “Definition of Consensus”, the Delegations of Chile and Colombia expressed their reservation on the way the matter had been handled in the Committee on General Principles and expressed the wish that this matter be addressed as a high priority.

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142. The Commission acknowledged the remarkable work done by the Committee on General Principles hosted by the Government of France as well as the support so far provided by FAO and WHO in this regard. The Chairperson observed that now the Commission was fully capacitated to efficiently develop international food standards for a decade to come and stand up to new challenges.

143. The Commission noted that some work was still necessary, especially on the review of the structure and mandates of the Committees and Task Forces (see Item 12b). One delegation stated that the Commission should consider monitoring the effectiveness of the new measures taken and developing an implementation plan for any additional actions that would contribute to the goal of the Commission.

REVIEW OF CODEX COMMITTEE STRUCTURE AND MANDATES OF CODEX COMMITTEES AND TASK FORCES (Agenda Item 12 b)⁶²

144. The Commission recalled that the 29th Session of the Commission had considered proposals put forward by the Secretariat on the structure and mandates of Codex Committees and Task Forces, had requested government comments through a Circular Letter (CL 2006/29-CAC) and had also invited FAO/WHO Coordinating Committees to provide their views on these proposals. The Commission noted that, due to time constraints, the 59th Session of the Executive Committee did not discuss in detail individual proposals as presented in CL 2006/29-CAC, with the exception of Proposal 8.

145. The Commission took note of the general comments of several members urging to further focus and expedite the work of Codex and to enhance the Committee management and standards management function of the Executive Committee. The matter was referred to the next session of the Executive Committee.

Proposal 1 (number of meetings)

146. The Delegation of Cameroon, while agreeing to the setting of upper limits, stated that they should be adopted on a temporary basis pending the improvement of the strategic planning process.

147. The Commission **agreed** to set an indicative upper limit on the number of Codex sessions planned per biennium (forty) and an indicative upper limit on the number of Codex sessions planned in one calendar year (twenty), in order to achieve a balanced session schedule within a biennium, with the understanding that these upper limits were based on the current, quite full Codex meeting schedule, but that they should be considered as indicative targets to allow for some flexibility. They would serve as a management tool to call the attention of the Executive Committee and the Commission where the number of meetings exceeded these targets.

148. The Commission recognized that an increased number of Codex meetings would have negative impact on the overall management of Codex standards setting work and on effective participation of Codex members.

149. Some delegations pointed out that the increasing number of physical working group meetings posed serious concerns and therefore suggested that the number of physical working group meetings should be closely monitored, with a view to better management of Codex work.

Proposal 2 (number of subsidiary bodies)

150. The Commission **agreed** to set an indicative target upper-limit (eighteen, excluding coordinating committees) on the number of active subsidiary bodies that could co-exist at one time, in order to avoid the increase of Codex sessions beyond a manageable level. The Commission also agreed to consider, before proposing to establish a new subsidiary body, dissolving or adjourning others, in accordance with work priorities identified by the Commission and in particular the Commission's six-year Strategic Plan.

⁶² CL 2006/29-CAC, ALINORM 07/30/9C Part II, ALINORM 07/30/9C Add.1, ALINORM 07/30/3 paras 109-118, LIM 11 (comments of Brazil, Malaysia and South Africa), LIM 14 (comments of Indonesia), LIM 16 (comments of China), LIM 20 (comments of Republic of Korea)

Proposal 3 (interval of meetings)

151. The Commission **agreed** to invite Codex committees to consider adopting a longer inter-session interval with the understanding that a structured, effective inter-session working mechanism should then be put in place in accordance with the Guidelines on Physical Working Groups and on Electronic Working Groups.

152. The Commission, referring to the recommendation under Agenda Item 10 on CCRVDF and CCFICS⁶³, noted that the intervals of meetings should be decided depending on the length of agenda as well as the use of working groups in the committees and that decision on the meeting intervals should be made on a committee-by-committee basis.

153. With regard to inter-session physical working group meetings, several delegations suggested that the number of physical working group meetings should be limited and that physical working groups should only undertake work on non-controversial issues and should not impose a limit to discussion at the committee level, and expressed concern that many physical working groups were held with a single working language. It was also proposed that physical working groups be held in conjunction with Codex subsidiary body meetings as much as possible, to save travel costs and enhance participation.

154. The Commission recalled that a wider coverage of languages was desirable in working groups and that the Guideline for Working Groups, in the Procedural Manual, clearly stated that working groups could not take any decisions on behalf of the committee that established them.

Proposal 4 (duration of meetings)

155. The Commission **agreed** that the duration of a Codex session should be kept within seven days, including the pre-session meetings of working groups, if any, in order to keep its proceedings well focused, ensure transparency, and facilitate effective participation of the members, with the understanding that a certain margin of flexibility should be allowed, depending on the workload of each subsidiary bodies.

Proposal 5 (use of ad hoc task forces)**Proposal 6 (consideration of merging or dissolving existing committees)****Proposal 7 (next comprehensive review)**

156. Due to time constraints, the Commission agreed to request the 60th Session of the Executive Committee to further consider the three proposals above.

Proposal 8 (conversion of regional standards into world-wide standards)

157. The Commission, taking into account the extensive discussion held and recommendations made at the 59th Session of the Executive Committee on Proposal 8, **endorsed** the amended Proposal 8, on a temporary basis, as follows:

“a) The commodity work of coordinating committees should concentrate on the development of regional standards, in compliance with their terms of reference. Conversion of a regional standard into a worldwide standard should, in principle, be considered after its adoption at Step 8, at the request of Codex members or a coordinating committee or at the recommendation of the commodity committee concerned, substantiated by a project document to be reviewed by the Executive Committee in the framework of the critical review, taking into account the programme of work of commodity committees concerned.

b) The proposal for new work for commodities having international trade potential should preferably be submitted through a worldwide commodity committee if such committee exists and is active, or in other cases, to the Commission through the Executive Committee.”

158. The Commission further noted that the 60th Session of the Executive Committee (December 2007) would review the outcome of a study to be undertaken by the bureau of the Commission to identify a set of draft procedures and criteria for use by the Executive Committee in its critical review process and eventually

⁶³ ALINORM 07/30/9A para. 6

by the Commission which would, amongst others, assist the Commission in streamlining its work on development of regional standards as opposed to worldwide standards and their conversion into worldwide standards.

159. The Delegation of Portugal, speaking on behalf of the Member States of the European Community, expressed the view that regional standards should not routinely be converted into world-wide standards after adoption at Step 8 and that such decision should be made on a case-by-case basis.

160. The Delegation of Cameroon stated that the principles adopted on a temporary basis should not constitute unnecessary obstacles to conversion of regional standards into world-wide standards and stressed that the standards development process by developing countries should be facilitated and encouraged as a way of enhancing their participation in the work of Codex.

Proposal 9 (relation between committees)

Proposal 10 (tasks related to nutrition)

Proposal 11 (role of private standards)

161. Due to time constraints, the Commission agreed to request the 60th Session of the Executive Committee to further consider the three proposals above.

MATTERS ARISING FROM REPORTS OF THE COMMISSION, CODEX COMMITTEES AND TASK FORCES (Agenda Item 13)⁶⁴

162. The Commission noted several matters arising from the reports of Codex Committees, including those matters arising from the previous session of the Commission, as contained in working documents ALINORM 07/30/9D and ALINORM 07/30/9D-Add.1.

163. The following paragraphs provide additional information on the comments made and decisions taken on certain items.

29th Session of the Codex Alimentarius Commission

Revision of WHO Guidelines for Drinking Water Quality⁶⁵

164. The Commission recalled that the completion of the revision of the WHO Guidelines for Drinking Water Quality (Third Edition, 2004) resulted in discrepancies of the values of certain health related substances between the Codex Standard for Natural Mineral Waters (CODEX STAN 108-1981) and the above WHO Guidelines. The Commission also recalled that an in-session working group chaired by Switzerland, acting as host country for the Committee on Natural Mineral Waters, had met during its present session to review the written comments received and provide recommendations to the Commission on whether the amendment to the Codex Standard on Natural Mineral Waters was necessary and, in the affirmative, how to proceed.

165. The Commission **endorsed** the conclusions of the working group presented in document CAC/30 LIM/19 that:

- the health-related limits for certain substances in the Codex Standard on Natural Mineral Waters should be reviewed and amended as necessary;
- this review and amendment should consider substances listed in the Annex to the Codex Circular Letter CL 2006/13-NMW on a case-by-case basis; and

⁶⁴ ALINORM 07/30/9D; ALINORM 07/30/9D-Add.1; CAC/30 LIM/06 (comments of Ghana); CAC/30 LIM/10 (comments of Malaysia); CAC/30 LIM/13 (comments of Brazil, European Community and Vietnam); CAC/30 LIM/18 (comments of Honduras)

⁶⁵ CL 2006/13-NMW; ALINORM 07/30/9D (comments of Australia, Brazil, Canada, Costa Rica, European Community, Norway, Paraguay, Peru, United States, Vietnam, ICBA, ICBWA); ALINORM 07/30/9D-Add.1 (comments of Mexico); LIM 19 (Report of an intrasessional working group of the Codex Alimentarius Commission on the Need to for Amendment of Health Related Substances in the Standard for Natural Mineral Waters)

- due to the complexity of issues involved, it would be difficult to reach agreement on the alignment of certain health related substances in the Codex Standard for Natural Mineral Waters working by an electronic working group or by correspondence.

166. The Commission therefore **agreed** that:

- work should be initiated through a Circular Letter, seeking further comments on each of the substances listed in the Annex of CL 2006/13-NMW including new discrepancies between the health-related limits for certain substances in the Codex Standard and the current version of the WHO guideline values for chemicals of health significance in drinking water;
- the Committee on Natural Mineral Waters (CCNMW) should be reactivated with a scope necessary to the review and amendment outlined above, in the light of comments already received and comments to be received in reply to the new Circular Letter; and
- the CCNMW should complete this task in no more than two sessions, and should propose a revised Section 3.2, "Health-related limits for certain substances" of the Codex Standard for Natural Mineral Waters for final adoption by the Commission at its Session in 2009.

167. The Delegation of Switzerland informed the Commission that a session of the CCNMW could be convened in Switzerland, in February 2008, with the exact dates and venue to be determined by the Host Government and the Codex Secretariat in due course.

Committee on Food Hygiene

The Use of the Lactoperoxidase System for Milk and Milk Products in International Trade⁶⁶

168. The Commission recalled that, during the adoption of the Code of Practice for Milk and Milk Products at its 27th Session, it had added as footnote 9 that the use of the lactoperoxidase system for milk and milk products would be examined by the Committee on Food Hygiene following the completion of an expert review by FAO and WHO of available data and considering the FAO Lactoperoxidase Expert Group Report about potential risks and benefits of lactoperoxidase.

169. The Commission was informed that the Committee on Food Hygiene had examined the issue as requested, based on the conclusions and recommendations of the FAO/WHO Technical Meeting on the Benefits and Potential Risks of the Lactoperoxidase System of Raw Milk Preservation (Rome, Italy, 28 November – 2 December 2005), but that it could not reach consensus because of the divergent views held by members on the recommendation for the removal of the restriction on the use of the lactoperoxidase system in milk and milk products intended for international trade as agreed to by the 19th Session of the Commission.

170. The Delegation of Cuba, supported by several other delegations, requested the removal of the restriction since the FAO/WHO Technical Meeting on the Benefits and Potential Risks of the Lactoperoxidase System of Raw Milk Preservation mandated by the Commission had taken place and had concluded that the lactoperoxidase system was safe if used in accordance with the *Guidelines for the Preservation of Raw Milk by Use of the Lactoperoxidase System* (CAC/GL 13-1991) and had recommended that this restriction be lifted.

171. The Delegation of the United States of America opposed the removal of the restriction and raised their concern that the technical meeting had mainly based their decisions on toxicological data, but had not considered literature that suggested that reduced rates of acid production might result in the possibility of outgrowth of pathogens, particularly acid-resistant strains of pathogens.

172. The Delegation of Singapore questioned the addition of chemicals to raw milk as this could constitute an adulteration of milk and questioned the safety of the chemicals used to activate the lactoperoxidase system in milk.

173. The Representatives of WHO and FAO emphasized the recommendations of the technical meeting, based on assessment of all available data, that the lactoperoxidase system was safe for use if used in accordance with CAC/GL 13-1991 and that there was a scientific basis for Codex to consider the removal of

⁶⁶ ALINORM 07/30/13, paras 29-32 and 188-195

the restriction on the international trade of milk and milk products treated with the lactoperoxidase system. It was further explained that the technical meeting had considered the issue of the lactoperoxidase system not only from a human health and nutrition perspective, but also from microbiological, processing and economic perspectives.

174. The FAO JECFA Secretariat further clarified that the 29th and 35th Sessions of JECFA had evaluated hydrogen peroxide and thiocyanate used to activate the lactoperoxidase system and had concluded that at the levels used, these chemicals did not pose a hazard to human health, but that thiocyanate could have a toxicological effect if iodine intake was not adequate and that the reports of these sessions of JECFA had been taken into account by the FAO/WHO technical meeting.

175. In view of the diversity of views and lack of consensus, the Commission **agreed** to refer the matter back to the Committee on Food Hygiene and to request, by Circular Letter, government comments that would facilitate the identification of additional information regarding the potential risks in respect of the lactoperoxidase system, for consideration by the Committee on Food Hygiene. The Committee on Food Hygiene would then evaluate all available evidence regarding the safety of the lactoperoxidase system when used in accordance with the *Guidelines for the Preservation of Raw Milk by Use of the Lactoperoxidase System* (CAC/GL13). The Committee should take into account the report of the FAO/WHO technical meeting and all other information submitted in response to the Circular Letter. The Commission agreed to remove footnote 9 from the Code of Hygienic Practice in Milk and Milk Products in view of the discussions that had taken place in the Committee on Food Hygiene.

176. The Delegations of Cuba, Colombia, Costa Rica and Burundi expressed their reservations on the decision to refer the matter back to the CCFH, observing that there was sufficient scientific evidence allowing the Commission to take a decision on the removal of the restriction.

177. The Commission also noted the concern of the Delegation of Cuba that the Delegation had not been able to attend the last session of the Committee on Food Hygiene held in the United States of America and that this matter was being addressed through channels outside Codex.

Committee on Sugars

Codex Standard for Sugars: Consideration of Method for determination of Colour in Plantation and Mill White Sugar⁶⁷

178. The Commission recalled that, while adopting a change of the method for determination of colour in the Codex Standard for Sugars (CODEX STAN 212-1999), the 24th Session of the Commission had requested the Committee on Sugars to examine whether a change in the method of analysis for the determination of colour might require a change in the specification of colour for plantation and mill white sugar. The Commission also recalled that, following the ICUMSA review of the methodology for colour determination and subsequent analysis of comments received in response to the CL 2006/32-CCS, the Committee on Sugars proposed to change the method of determination of colour for all sugars including plantation or mill white sugar without making a change to the colour specification for plantation and mill white sugar.

179. The Delegation of Brazil referred to its comments in CAC/30 LIM/13 and reiterated its concern with the recommendation of the Committee on Sugars to exclude the method GS2/3-9 currently extensively applied in Brazil.

180. After some discussion, the Commission **agreed** to forward the recommendations of the Committee on Sugars contained in document ALINORM 07/30/9D and the written comments submitted on this matter in CAC/30 LIM/13 to the Committee on Methods of Analysis and Sampling for their consideration and endorsement of methods of colour determination for all sugars, with a view to adopting it by the 31st Session of the Commission in 2008.

⁶⁷ ALINORM 07/30/9D, paras 9-14; CL 2006/32-CCS; CAC/30 LIM/13 (comments of Brazil and European Community)

Committees on Nutrition and Foods for Special Dietary Uses and on Food Labelling

*WHO Global Strategy on Diet, Physical Activity and Health: actions that could be taken by Codex*⁶⁸

181. The Commission recalled that the background to the consideration of the Global Strategy was as follows. WHA Resolution 57.17 endorsing the Global Strategy requested the Codex Alimentarius Commission “to continue to give full consideration within the framework of its operational mandate, to measures which it might take to contribute towards the improvement of health standards of foods consistent with the aims and objectives of the Global Strategy.”

182. The 28th Session of the Commission agreed to ask WHO, in cooperation with FAO, to prepare a document focused on actions that could be taken by Codex including specific proposals for new work for consideration by the Committee on Nutrition and Foods for Special Dietary Uses (CCNFSDU) and the Committee on Food Labelling (CCFL).

183. The 29th Session of the Commission considered a progress report on the consideration of the Global Strategy and agreed that WHO and FAO would complete a document containing concrete proposals for possible actions by Codex that would be circulated for comments and consideration by the CCNFSDU and CCFL.

184. The Representative of WHO recalled that the document on the implementation of the Global Strategy contained a series of proposed actions related to nutrition labelling, certain nutrition claims, quantitative declaration of ingredients, modification of standardized foods and production and processing standards regarding nutritional quality and safety of foods. The Representative thanked the members who had provided comments and participated in the work of the two Committees concerned for their active contribution and indicated that WHO and FAO planned to strengthen their interaction with the Committees concerned as regards the implementation of the Global Strategy. The Representative indicated that WHO and FAO would make every effort to inform these Committees of the scientific advice currently being produced by WHO and FAO and of any plans for relevant scientific work which may affect the implementation of the Global Strategy, including the planned FAO/WHO Expert Consultation on Fats and Oils in Human Nutrition.

185. The Representative of FAO expressed his appreciation for the work carried out so far and looked forward to further consideration of the Global Strategy in the relevant Codex Committees, while recalling the ongoing cooperation with WHO in this area.

186. The Delegation of Germany, speaking as Chair of the CCNFSDU, stressed the importance of the Global Strategy to address public health issues related to non communicable diseases and indicated that the Committee had agreed to proceed with the consideration of the revision of the Nutrient Reference Values (NRVs) for vitamins and minerals and to ask the advice of the Committee on Food Labelling concerning the revision and extension of the list of NRVs in the Guidelines for Nutrition Labelling to other nutrients associated with increased and decreased risk of non communicable diseases. The Committee had agreed that if this reply was positive it would consider new work on the revision and extension of the list to relevant nutrients at its next session. The Delegation noted that there was no support in the Committee to initiate work on claims for trans fatty acids and restrictions on saturated and trans fatty acids in the conditions for comparative claims.

187. The Delegation of Canada, speaking as Chair of the Committee on Food Labelling, recalled that the Committee had discussed extensively the proposals for action related to labelling issues, with the following result: there was no support for the amendment of the Purpose of the *Guidelines on Nutrition Labelling*, no conclusion on the need to amend the *Guidelines* to require mandatory nutrient declaration, on the revision of the current list of nutrients that should always be declared, and on the development of additional criteria for nutrient presentation. There was no support to undertake new work on nutrition claims for trans fatty acids. The Committee had agreed with the proposal of the CCNFSDU to revise the list of vitamins and minerals but had not reached a conclusion on the extension of the list to other nutrients. The Delegation further advised that a working group would be held immediately prior to the next session of the Committee in May 2008 to consider all issues relevant to the draft action plan.

⁶⁸ ALINORM 07/30/26, paras. 144-147, ALINORM 07/30/22, paras. 20-64, CAC/30 INF/13 (Progress Report on the Implementation of the WHO Global Strategy on Diet, Physical Activity and Health as related to Codex)

188. The Commission **agreed** that it would defer further consideration of the implementation of the Global Strategy to its next session in order to consider the outcome of the Committee on Food Labelling to be held in May 2008.

Committee on Methods of Analysis and Sampling

Reference to the IUPAC/ISO/AOAC Protocols and Guidelines⁶⁹

189. The Commission noted the reply of the Committee on Methods of Analysis to the request for clarification at its 29th Session as to whether the above Protocols and Guidelines should be identified separately. The Commission therefore **agreed** to replace the current single reference to *Food Control Laboratory Management Recommendations* (CAC/GL 28-1995) with individual references to the following texts:

- International Harmonised Protocol for the Proficiency Testing of (Chemical) Analytical Laboratories (1995, revised 2006)
- Protocol for the Design, Conduct and Interpretation of Method Performance Studies (1997)
- Harmonised Guidelines for Internal Quality Control in Analytical Chemistry Laboratories (1997)

Committee on Fats and Oils

Linolenic Acid Level for the Standard for Olive Oils and Olive Pomace Oils⁷⁰

190. The Commission recalled that its 26th Session had adopted the Standard for Olive Oils and Olive Pomace Oils without a level for linolenic acid and with a footnote stating “Pending the result of the IOOC survey and further consideration by the Committee on Fats and Oils, national limits may remain in place”. The Commission was informed that the Committee had considered the above survey and agreed to circulate a Proposed Draft Linolenic Acid Level in Section 3.9 of the Standard with a footnote, as a proposed draft amendment to the Standard at Step 3. The Commission noted the Committee had resumed its work on the Standard for Olive Oils and Olive Pomace Oils and encouraged the Committee to solve the issue of the level of linolenic acid.

Committee on Contaminants in Foods

Amendments to Schedule I of the General Standards for Contaminants and Toxins in Foods⁷¹

191. The Commission **adopted** the following amendments to Schedule I: i) deletion of references ‘CS 248-2005’ and their replacement by the adoption year ‘2005’; ii) reorganization of contaminants into the following four categories: metals, mycotoxins, other chemicals and radionuclides, as proposed by the Committee.

Guideline levels for Methylmercury in Fish

192. The Commission recalled that its 29th Session⁷² had requested FAO and WHO for scientific advice on the health risks associated with methylmercury and dioxins and dioxin-like PCBs in fish and the health benefits of fish consumption. The Representative of FAO, speaking on behalf of FAO and WHO, informed the Commission that a step-wise preparatory process was being taken, given the complex nature of the issue and the need for innovative principles and methodology. The Representative indicated that, possibly at a first stage, FAO and WHO would consider conducting qualitative risk-benefit assessment of fish consumption, specifically addressing issues related to the impact of methylmercury exposure on women of child-bearing age and at a later stage, conducting quantitative assessment including the intake of dioxin and dioxin-like PCBs, taking into account consumption of fatty fish, considered as a significant source of beneficial fatty acids.

⁶⁹ ALINORM 07/30/23, paras 16-17, ALINORM 06/29/41, paras 197-198

⁷⁰ ALINORM 03/30/17, para. 107, Appendix VII

⁷¹ ALINORM 07/30/41, para. 46

⁷² ALINORM 06/29/41 para.195

Committee on General Principles

*Proposed Draft Code of Ethics for International Trade in Food*⁷³

193. The Commission recalled that during the ongoing revision work of the existing Code, the Committee on General Principles (CCGP) had requested the Committee on Food Import and Export Inspection and Certification Systems (CCFICS) to evaluate if part of the elements of the present Code were covered by existing texts in the Codex Alimentarius developed by the CCFICS or if these texts could be expanded accordingly. The CCFICS had found that some of the elements were covered by the existing texts and had additionally forwarded three recommendations to the CCGP. When deciding to circulate the Proposed Draft Revised Code of Ethics for International Trade in Food for comments at Step 3, the CCGP at its last session also decided to forward the recommendations from the CCFICS to the Commission for endorsement.

194. The Commission, bearing in mind the difficulties encountered by member countries with insufficient capacity for food import and export control, **decided** to:

- encourage member countries to further implement the provisions in existing Codex guidelines on food import and export inspection and certification systems related to the subsequent export of food, whether imported or produced domestically, that had been found to be unsafe or unsuitable;
- encourage FAO, WHO and other international organizations to give priority to providing technical assistance to member countries with insufficient capacity for establishing and implementing food import and export control systems; and
- encourage those member countries with insufficient control systems to give priority in their capacity building/technical assistance needs assessments to the issue of import control systems.

195. The Commission noted an intervention from the Representative of WHO that bilateral donor agencies should also be encouraged to provide technical assistance for establishing and implementing food import and export control systems.

*Procedures for the Elaboration of Codex Standards and Related Texts*⁷⁴

196. The Commission recalled that at its 27th Session it had referred a number of comments from India on the *Procedures for Elaboration of Codex Standards and Related Texts* to the CCGP. The CCGP at its 22nd Session agreed that the Delegation of India would prepare a discussion paper providing the objectives and rationale for the proposed changes to the Elaboration Procedure. At its 23rd Session, the CCGP briefly discussed the paper but considered that it was yet premature to request approval of the Commission to initiate new work on these subjects. The 24th Session of the Committee held an in-depth discussion on the basis of document CX/GP 06/23/6 Part-I and agreed to forward the content of the discussion on the Procedure for the Elaboration of Codex Standards and Related Texts to the Commission for further advice on how this work could be pursued and in which forum.

197. The Commission considered the proposals contained in the above-mentioned document one by one, as follows.

Reference to decisions taken by consensus in the Elaboration Procedure, including a definition of that term

198. The Commission noted that the definition of consensus and how the concept was handled in practice in Codex was considered an important issue by many members, to be further discussed as a matter of priority within the CCGP.

199. After some discussion as to how best to prepare for discussions at the forthcoming session of the CCGP in 2009 with due input from the Chairpersons of Codex subsidiary bodies, while ensuring transparency and inclusiveness, the Commission **agreed** that:

⁷³ ALINORM 07/30/33, paras 106-107

⁷⁴ ALINORM 07/30/33, paras 116-130

- The issue be discussed by the 60th Session of the Executive Committee (December 2007), including how to request chairpersons of Codex subsidiary bodies to provide their input on the matter and especially their experiences with the application of the *Measures to Facilitate Consensus*;
- The Secretariat compile the replies from the chairpersons into a discussion paper and circulate it to members and observers at the earliest possible time in the second half of 2008, in order to allow for ample time for reflection and the preparation of comments and proposals by Codex members for discussion of the issue at the 25th Session of the CCGP in April 2009.

200. The Commission noted that this process would not prevent the matter from being discussed at the 61st Session of the Executive Committee and the 31st Session of the Commission if the members so wished.

Elaboration of provisions on how to take into account the situation of developing countries within the Critical Review

201. The Commission confirmed that the special needs of developing countries had already been taken in the current Elaboration Procedure, particularly its Part 2 “Critical Review” and in the *Criteria for the Establishment of Work Priorities* and that no new work was necessary on this matter.

Scope of the Critical Review including the basis of the decision to entrust work to a Committee other than the one to which it had originally been entrusted

202. The Commission confirmed that as per the explanation of the Secretariat provided at the 24th Session of the CCGP⁷⁵, no further discussion on this matter was necessary.

Committee on Food Additives

Codex General Standard for Food Additives – Food Category 02.2.1.2

203. The Commission noted that the new work on the revision of the Food Category System (FCS) of the Codex General Standard for Food Additives (see Appendix VII) addressed the need to ensure better correspondence of the FCS and the products covered by the adopted Standard for Fat Spreads and Blended Spreads (see Appendix IV). In view of the revocation of the Codex Standard for Margarine (see Appendix VI), which had covered the products that were now included in the newly adopted Standard, the Commission agreed to delete food category 02.2.1.2 “Margarine and similar products” from the Annex to Table 3 of the General Standard for Food Additives.

Codex General Standard for Food Additives – Food Category 02.1.1

204. The Commission agreed to adopt the amendment to food category 2.1.1 “Butter oil, anhydrous milkfat and ghee” of the Codex General Standard for Food Additives, as proposed by the Committee on Food Additives.

Committee on Pesticide Residues

Enforcement of Codex MRLs at National Level⁷⁶

205. The Commission recalled that the issue of enforcement of Codex MRLs at national level had been discussed by the Committee on Pesticide Residues where many member governments expressed their concerns that some countries were imposing stricter MRLs than those in the Codex Alimentarius without sufficient scientific justification and that this impeded trade for developing countries.

206. The Secretariat confirmed that the enforcement of Codex standards including MRLs was an issue outside the mandate of the Commission and indicated that possible venues for considering this issue were the FAO/WHO Regional Coordinating Committees, where this matter could be considered under the standing agenda item “Information on Use of Codex Standards at National and Regional Level”, or the WTO SPS Committee that regularly monitored the use or non use of international standards including Codex standards and related texts.

⁷⁵ ALINORM 07/30/33, para. 129

⁷⁶ ALINORM 07/30/24, paras 204-211; ALINORM 07/30/3 para.80; CAC/30 LIM/13 (comments of European Community)

207. The Observer from WTO, referring to the relevant provisions of the SPS Agreement in this area, in particular, Article 3 and Article 12, pointed out that the SPS Agreement strongly encouraged the use of international standards by WTO members and that the SPS Committee had developed a procedure to monitor the use of international standards, in accordance with Article 12.4.

208. Some delegations stressed the importance of further consideration of this matter, expressing the view that technical assistance was necessary for developing countries to overcome this problem.

209. The Commission noted that the lack of capacity to generate scientific data by developing countries, especially with regard to pesticide MRLs, had been discussed at the 59th Session of the Executive Committee which had noted a proposal made by the Member for Latin America and the Caribbean to include an additional activity which would address the strengthening of scientific advisory groups, to improve their efficiency as well as to equip them with necessary resources, particularly in the area of pesticide residues. The Commission noted that the concerns expressed in relation to this proposal were related amongst others to the lack of capacity to generate scientific data in developing countries and that some of these concerns might best be addressed outside Codex, for instance, through international FAO/WHO workshops on pesticide residues. The Commission further noted that this matter had been addressed in Activity 1.7 under Goal 1 of the Strategic Plan 2008 -2013.

RELATIONS BETWEEN THE CODEX ALIMENTARIUS COMMISSION AND OTHER INTERNATIONAL ORGANIZATIONS (Agenda Item 14)

RELATIONS BETWEEN THE CODEX ALIMENTARIUS COMMISSION AND OTHER INTERNATIONAL INTERGOVERNMENTAL ORGANIZATIONS (Agenda Item 14a)⁷⁷

Relation between Codex and the World Organisation for Animal Health (OIE)

210. The Commission was reminded that at its 28th Session it had endorsed the recommendations of the 55th Session of the Executive Committee related to the collaboration between Codex and OIE and that the effectiveness of cooperative arrangements between Codex and OIE in accordance with the recommendations should be reviewed by the 30th Session of the Commission with a view to considering if further arrangements would be necessary or desirable, including those provisions mentioned in paragraph 13 of the *Guidelines on Cooperation with International Intergovernmental Organizations*.⁷⁸

211. The Observer from OIE, referring to the written submission, drew the attention of the Commission to two main points: the framework for cooperation between OIE and Codex, and the organisation of an international Conference addressing both OIE and Codex standards on traceability and their implementation at the national level.

212. With regard to the current framework for cooperation between OIE and Codex, he explained that OIE had been involved in consultations with Codex, FAO and WHO since 2001 with a view to improving the coordination of standards setting activities. He highlighted that the cooperation between OIE and Codex had produced over the present years positive results through the exchange of information and the cross-referencing between the respective international standards. He indicated that good examples of collaboration in the development of standards included texts produced by the two organizations regarding meat inspection and animal/product identification and that there was still room for further improvement.

213. The Observer stated that it would be appropriate to formalize the relationship between OIE and Codex by strengthening the legal basis for the production of international standards, including the development of joint OIE-Codex standards, where appropriate. He encouraged the Commission to recommend that OIE, FAO and WHO legal services work together with a view to eventually updating their existing mutual cooperation agreements to enable the establishment of a formal agreement between OIE and Codex.

214. On the second point, the Observer informed the Commission of the OIE's plan to hold an international Conference in 2009 in Argentina on the implementation of OIE standards on identification and traceability of live animals. He invited the Commission to consider broadening the scope of the Conference by including

⁷⁷ ALINORM 07/30/9E; CAC/30 INF/4 Rev.1 (OIE); CAC/30 INF/5 (WTO); CAC/30 INF/6 (IAEA); CAC/30 INF/7 (OIV)

⁷⁸ ALINORM 05/28/41, paras 201-203

Codex standards thus covering the entire food chain and invited FAO and WHO to work together with OIE to organise the event.

215. The Representative of the Legal Counsel of FAO, in presenting the common view of the Legal Offices of FAO and WHO, stated that Codex, despite its functional autonomy, was a statutory body of the parent organizations FAO and WHO and could only act through its parent organizations as regard its relations with external organizations. He informed the Commission that Codex had no legal capacity to conclude agreements and to be, in its own right, directly a party to any agreement or any contractual arrangement and that any agreement concerning Codex, if the need would arise, would necessarily be between FAO and WHO, on the one hand, and OIE, on the other hand. The Representative further stated that OIE was currently a party to agreements with both FAO and WHO and that these agreements provided a broad framework under which a wide range of cooperation activities could be carried out. He indicated the need to identify all practical issues that deserve special and specific treatment and clearly identify problems that hindered practical collaboration between Codex and OIE, with due consideration being paid to the status of FAO and WHO as organizations of the United Nations System.

216. Many delegations expressed their support for further strengthening the collaboration between Codex and OIE. Views expressed included: that there was a need for more consistency between Codex and OIE texts in order to ensure a coordinated approach to food safety throughout the food chain; that collaboration between veterinary and public health services needed to be strengthened to enhance food safety; that bilateral agreements between OIE and FAO/WHO currently in place needed to be modified to foster more systematic collaboration between Codex and OIE; that there were many areas of common interest for Codex and OIE including traceability, salmonellosis, certification, antimicrobial resistance; that there was a need to ensure more consistency in the decision-making process and standards between Codex and OIE; that collaboration between Codex and OIE needed to be strengthened also at national and regional level; and that the *Guidelines on Cooperation with International Intergovernmental Organizations* provided adequate guidance to ensure good collaboration between Codex and OIE and there was no need to review the current arrangements for cooperation.

217. In response to the statement made by the Representative of the Legal Counsel of FAO, the Observer from OIE indicated the lack of a specific reference to Codex in the WHO-OIE Agreement as one of the problems to be addressed to further strengthen the collaboration between Codex and OIE.

218. The Commission concluded its discussion by noting the ongoing substantial cooperation between Codex and OIE and recognised the need to further strengthen this collaboration on substantive matters.

219. The Commission **recommended** that FAO and WHO study the possibility of reviewing or updating FAO and WHO Agreements with OIE, as might be required. It also **requested** the Codex Secretariat to identify, in cooperation with the Legal Offices of FAO and WHO, any practical problems affecting the cooperation between Codex and OIE that might need to be addressed in a pragmatic manner, and taking into account all relevant circumstances.

220. With regard to the planned OIE Conference on traceability, the Commission noted that such event could best be organised in collaboration with FAO and WHO. It was also noted that the experts having Codex background could usefully participate in the planned Conference to keep informed the Commission and its subsidiary bodies on relevant development in order to facilitate the participation of those interested at this important event.

World Trade Organization (WTO)

221. In addition to the information provided in CAC/30 INF/5 on the work of the World Trade Organisation (WTO), the Observer from WTO informed the Commission of a number of other issues, especially those from the SPS Committee which had met just prior to this session of the Commission. The Observer highlighted four areas related to transparency, private standards, the second review of the implementation of the SPS Agreement, and technical assistance.

222. In particular, the Commission was informed that a workshop on transparency was scheduled for 15 October 2007 and that members were invited to submit proposals regarding possible modification to the Recommended Procedures for Implementing the Transparency Obligations of the SPS Agreement for consideration at the workshop and at subsequent Committee meetings and that one proposal already under discussion suggested that WTO members notify all new or modified sanitary and phytosanitary measures of trade significance, whether or not they conformed to international standards. In addition, the Commission noted that there were ongoing discussions on private standards in the SPS Committee, that an information

session had been held on 25 June 2007 on this matter and that the presentations were available on the WTO webpage (www.wto.org).

223. The Commission also noted that the SPS Committee, in the context of the second review of the implementation of the SPS Agreement, decided to give priority to the consideration of two issues: the use of *ad hoc* consultations and the relationship between the SPS Committee and Codex, OIE and IPPC.

224. With regard to technical assistance, the Commission noted the expression of thanks to the Codex Secretariat for their participation in and contribution to WTO regional workshops on the SPS and TBT Agreements as well as specialised courses on the SPS Agreement.

225. The Delegation of Chile emphasized the importance of the coordination between Codex and WTO in particular to cooperate in the monitoring use of international standards in relation to harmonization. The Delegation requested that Codex should continue its contacts with NGOs, considering that a number of private standards had a bearing on Codex standards.

International Atomic Energy Agency (IAEA)

226. The Commission noted the information provided in CAC/30 INF/6 and thanked the IAEA for the information.

The International Organisation of Vine and Wine (OIV)

227. The Observer from OIV, referring to the information provided in CAC/30 INF/7, highlighted the cooperation of OIV with Codex on a number of issues. These included work on the Code of Practice for the Prevention and Reduction of Ochratoxin A Contamination in Wine and the Standard for Table Grapes, both adopted at this session of the Commission.

228. The Commission also noted that the OIV at its General Assembly in June 2007 had taken into account different principles established by the Commission through the Committee on Food Import and Export Inspection and Certification Systems in the elaboration of the OIV draft resolution related to traceability guidelines in the vitiviculture sector.

229. The Commission **expressed** its thanks to the Observers of the intergovernmental organizations for the useful information provided in the present session and their continued cooperation with the Codex Alimentarius Commission.

RELATIONS BETWEEN THE CODEX ALIMENTARIUS COMMISSION AND INTERNATIONAL NON-GOVERNMENTAL ORGANIZATIONS (Agenda Item 14b)⁷⁹

230. In accordance with section 6, paragraph 4 of the *Principles Concerning the Participation of International Non-Governmental Organizations in the Work of the Codex Alimentarius Commission*, the Secretariat reported to the Commission on the relations between the Commission and international non-governmental organizations.

Relations with the International Organization for Standardization (ISO)

231. The Commission noted the detailed information provided in document CAC/30 INF/8 on the work of ISO relevant to Codex work as well as the ongoing contacts for information exchange between the Codex and ISO Secretariats.

232. The Observer from ISO expressed the hope that through ongoing coordination and cooperation, Codex and ISO standards could remain complementary. The Observer informed the Commission of the recent creation of ISO/TC 234, "Fisheries and Aquaculture" and the publication of a joint ISO/ITC handbook on ISO 22000 to assist small businesses, especially in developing countries and transition economies, in their effort to improve their market share of food and agricultural products in the global market.

233. Several delegations stressed the importance of maintaining and strengthening coordination and cooperation between Codex and ISO both at the secretariat level as well as between Codex Contact Points

⁷⁹ ALINORM 07/30/9E, CAC/30 INF/2 (International non-governmental organization in observer status with the Codex Alimentarius Commission); CAC/30 INF/8 (Communication from ISO – Report of Activities Relevant to Codex)

and national member bodies of ISO in order to ensure the complementarity of the work carried out and to avoid duplication of work or contradiction in the standards. One delegation suggested that the cooperation should not be limited to the technical committees but also apply to the policy committees of ISO such as the Committee on Conformity Assessment (CASCO).

234. The Commission **supported** continued cooperation and coordination with ISO and agreed that the Codex Secretariat should maintain its contacts with ISO and continue to report regularly to the Commission on ISO activities of relevance to Codex work. The Commission also **supported** increased coordination and cooperation between the focal points of Codex and ISO at the national level.

FAO/WHO PROJECT AND TRUST FUND FOR ENHANCED PARTICIPATION IN CODEX (Agenda Item 15)⁸⁰

235. The Representative of WHO, on behalf of FAO and WHO, expressed their appreciation to the generous contribution from donor countries to the Trust Fund and drew the attention of the Commission to the additional study carried out in 2007 focusing on the impact of the Trust Fund on enhanced trade opportunity and the strengthening of national food safety institutions. The Representative noted that the study also contained several recommendations to improve the effectiveness of the Trust Fund, including the more support to activities at national level, which could lead to enhanced participation in Codex meetings and better implementation of Codex standards and related texts at national level.

236. Many delegations stated that the increased participation in Codex meetings achieved thanks to the Trust Fund had had various positive impacts in their countries, which could be further amplified if paralleled with capacity building activities to strengthen food safety legislation and enforcement. In this regard, the Commission noted that capacity building activities provided through the Trust Fund were restricted to those for the purpose of better participation in Codex meetings and that capacity building in the area of food safety in general would better be sought by other means, including through bilateral technical assistance and the Standards and Trade Development Facility.

237. The Delegation of Cameroon suggested that a midterm review of the Trust Fund should be considered to follow up on the matters identified in the first biennial review and that two-thirds of the funds should be targeted at the capacity building.

238. Some delegations expressed concern on the current criteria for the classification of eligible countries because they believed that the current classification primarily based on economic indicators did not correctly reflect the need and capacity of each country. The Representative of WHO responded that the selection criteria were under review while the basic principle would continue to be giving higher support to countries with larger difficulties in participating in the Codex process.

239. The Commission expressed its appreciations to the effort being made by FAO and WHO and to the donors making financial contribution. The Commission encouraged current donors to continue to provide funds to the Trust Fund and invited other countries to consider contributing to the Fund in order to ensure its sustainability, welcoming the move of Brazil and Malaysia to make contributions to the Fund.

OTHER MATTERS ARISING FROM FAO AND WHO (Agenda Item 16)⁸¹

Part I: Outcomes of Recent FAO/WHO Expert Meetings

240. The Representative of FAO, on behalf of FAO and WHO, informed the Commission of the major outcomes of the FAO and WHO expert meetings and related activities carried out since the last Session of the Commission and future meetings to be held in 2007, including those of JECFA, JMPR and JEMRA. The Representative noted that in view of increasing workload and requests for scientific advice, alternative ways were being sought to mobilize the necessary resources to provide such advice.

241. The Delegations of Paraguay and Brazil requested information on the outcomes of the recently held 68th meeting of JECFA, especially with regard to evaluation of steviol glycosides. The FAO Secretariat of JECFA informed the Commission that the JECFA meeting considered that the newly examined data did not

⁸⁰ ALINORM 07/30/9F Rev

⁸¹ ALINORM 07/30/9G, ALINORM 07/30/9G Add.1

raise additional safety concerns, but that the results of some ongoing clinical studies, which had been specifically requested at the sixty-third meeting of JECFA, needed to be provided to complete full evaluation. Therefore JECFA had agreed to maintain the temporary ADI, expressed as steviol, pending submission of the results of ongoing studies. The Commission further noted that this matter would be considered by the next session of JECFA in June 2008, that a summary report of the 68th meeting would be available on the website in two weeks time, and that a full report would be published by WHO early 2008. The Delegation of Paraguay, while expressing the great interest for steviol glycosides for their country, expressed its concern with the results of the evaluation and the decision taken by JECFA.

Part II: Report of Conclusions of the FAO/WHO Consultative Process on Provision of Scientific Advice to Codex and Member Countries

242. The Representative of FAO, speaking on behalf of FAO and WHO, informed the Commission of the final conclusions to the FAO/WHO Consultative Process initiated at the request of the 24th Session of the Commission (2001) and highlighted some of the main points, as follows.

- The FAO/WHO Framework on the Provision of Scientific Advice was developed to document the principles, practices and procedures currently applied by FAO and WHO for the provision of scientific advice in order to continue strengthening the independence, transparency and quality of scientific advice.
- FAO and WHO jointly continued to prioritise Codex requests for scientific advice, taking in consideration the criteria proposed by Codex (ALINORM 05/28/3, para. 75) as well as the requests of advice from member countries and the availability of resources. The status of requests for scientific advice was presented annually to the Commission.
- The FAO/WHO meeting on “*Enhancing Developing Country Participation in FAO/WHO Scientific Advice Activities*” (Belgrade, 12-15 December 2005) provided a range of recommendations to FAO and WHO in three main areas: greater inclusion of data from developing countries; enhancement of the potential for experts from developing countries; and means to enhance the enabling environment at national, regional and international levels.

Part III: Status of Requests for FAO/WHO Scientific Advice

243. The Representative of WHO, speaking on behalf of FAO and WHO, highlighted that recent increase in requests from Codex subsidiary bodies and member states of FAO and WHO for scientific advice related to food safety required more resources to be mobilized so as to facilitate the provision of scientific evidence in a timely and appropriate manner, particularly among others, in the area of risk assessment of microbial hazards in fresh foods including viruses. The Representative stated that priority had also been given to emerging issues in food production related to human health (e.g. antimicrobial resistance, biotechnology and nanotechnology) as well as to areas requiring innovative approach such as risk-benefit assessment (e.g. active chlorine, consumption of fish associated with methylmercury) .

Part IV: Other Matters from FAO and WHO

244. The Representative of WHO informed the Commission that the International Health Regulations (IHR (2005)), which were legally binding to Member States of WHO and covered serious health hazards in foods in international trade, had come into force in June 2007. The Representative also indicated that the International Food Safety Authorities Network (INFOSAN) was an effective tool to communicate emergency information related to food safety and manage food safety related events which required notification under the IHR, and urged that relevant food safety authorities register their contact point for INFOSAN.

Part V: FAO/WHO Capacity Building Activities in Food Safety and Quality

245. The Commission was informed of the FAO/WHO activities in capacity building in the field of food safety and quality as provided in ALINORM 07/30/9G Add.1. The Commission was informed that capacity building activities were conducted at national, regional and global level and took the form of field projects, training activities and development of tools such as manuals and training packages. Several key activities were highlighted: the programme for improving the quality and safety of fresh fruits and vegetables; strengthening the capacity of national food control systems: guidelines to assess capacity building needs; risk analysis and support to national Codex work, amongst others.

246. The Commission noted that FAO and WHO were trying to keep up with member country demands and were improving effectiveness with new training modalities and approaches such as the e-learning courses and the use of indicators for the impact evaluation of capacity building activities.

APPOINTMENT OF REGIONAL COORDINATORS (Agenda Item 17)⁸²

247. In accordance with Rule IV.2 of the Commission's Rules of Procedure, and on the basis of the nominations made by the Coordinating Committees, the following Members of the Commission were **appointed** as Coordinators to hold office from the end of the Thirtieth Session of the Commission until the end of the regular session of the Commission held in 2009.

Africa: Ghana
 Asia: Indonesia
 Europe: Switzerland
 Latin America & the Caribbean: Mexico
 Near East: Tunisia
 North America & South-West Pacific: Tonga

ELECTION OF OFFICERS OF THE COMMISSION AND ELECTION OF MEMBERS OF THE EXECUTIVE COMMITTEE (Agenda Item 18)⁸³

248. The Commission **elected**, by general consent, the following persons to hold office as Chairperson and Vice-Chairpersons of the Codex Alimentarius Commission from the end of its present Session to the end of the next regular session of the Commission.

Chairperson: Dr Claude J.S. MOSHA (United Republic of Tanzania)

Vice-Chairpersons: Dr Karen HULEBAK (United States of America)

Ms NORAINI Mohd. Othman (Malaysia)

Dr Wim VAN ECK (The Netherlands)

249. The following Members of the Executive Committee were **elected** on a geographic basis for the period from the end of the current session to the end of the second succeeding regular session of the Commission:

Africa: Mali
 Asia: Japan
 Europe: United Kingdom
 Latin America & the Caribbean: Argentina
 Near East: Jordan
 North America: Canada
 South-West Pacific: New Zealand

⁸² ALINORM 07/30/2, paras 33–34; ALINORM 07/30/15, paras 142–145; ALINORM 07/30/19, paras 54–55; ALINORM 07/30/28, paras 61–62; ALINORM 07/30/32, para. 78; ALINORM 07/30/36, para. 124; ALINORM 07/30/40, para. 82

⁸³ ALINORM 07/30/2

DESIGNATION OF COUNTRIES RESPONSIBLE FOR APPOINTING THE CHAIRPERSON OF CODEX COMMITTEES AND AD HOC TASK FORCES (Agenda Item 19)⁸⁴

250. The Commission **confirmed** the designation of the Host Governments as listed in the Appendix X to this report.

251. In arriving at its decision, the Commission noted that United Kingdom no longer sought to host the Committee on Fats and Oils and noted the willingness of Malaysia and Argentina to serve as host government for this Committee. The Commission proceeded with a secret ballot and designated Malaysia as the host government for the Committee.

252. The Delegation of Malaysia indicated its commitment to ensure effective operation of this Committee. The Delegation of Argentina congratulated Malaysia on their designation and wished them every success.

OTHER BUSINESS (Agenda Item 20)

253. Due to time constraints, the Commission did not discuss the necessity of circulating Codex documents simultaneously in all working languages, which had been proposed for discussion under this agenda item by the Delegation of Columbia (*see* para. 6).

254. The Commission noted that its 31st Session would be held in Geneva, Switzerland, from 30 June to 5 July 2008, subject to further confirmation.

APPENDIX I

Chairperson:
Président:
Presidente:

Dr Claude J.S. MOSHA
Chief Standards Officer (Food Safety & Quality)
Head, Agriculture and Food Section
Tanzania Bureau of Standards
P.O. Box 9524
Dar-es-Salaam
Tel.: +255 22 245 0206 (mobile: 255 713 324495)
Fax: +255 22 245 0959
Email: claudem.mosha@tbstz.org; codex@tbstz.org;
info@tbstz.org

LIST OF PARTICIPANTS
LISTE DES PARTICIPANTS
LISTA DE PARTICIPANTES

ALGERIA - ALGÉRIE - ARGELIA

M Abdelhalim ACHELI
Directeur
Direction de la qualité et de la consommation
Ministère du commerce
Boulevard Mohamed V
Alger
Phone: +213 21 63 1355/213 21 63 2255
Email: cacqe@cacqe.org

Mme Fouzia DOUHI
Secrétaire du Comité National du Codex
Alimentarius
Chef de service Méthodes d'Analyses
Centre Algérien du contrôle de la qualité et de
l'emballage
Route nationale N° 5 Bab Ezzouar BP 206
Bordj El Kiffen
Alger
Phone: +213 (0) 71953003
Fax: +213 (0) 21 243011
Email: douhi_fouzia@yahoo.fr

Mme Baya CHETTOUF
Sous Directeur de la réglementation des produits
alimentaires
Ministère du commerce
Boulevard Mohamed V
Alger
Phone: +213 21 631280
Email: cacqe@cacqe.org

M Abderrahman HAMIDAOU
Ministre plénipotentiaire
Représentant permanent adjoint auprès de la FAO
Ambassade de la République algérienne
démocratique et populaire
Via Bartolomeo Eustachio, 12
00161 Rome

Mme Habiba TALEB
Secrétaire diplomatique
Ambassade de la République algérienne
démocratique et populaire
Via Bartolomeo Eustachio, 12
00161 Rome

ANGOLA

M Manuel Pedro PACAVIRA
Ambassadeur
Représentant permanent auprès de la FAO
Ambassade de la République d'Angola
Via Druso, 39
00184 Rome

M Kiala KIA MATEVA
Conseiller
Représentant permanent adjoint auprès de la FAO
Ambassade de la République d'Angola
Via Druso, 39
Rome
Email: nsengalu@hotmail.com
kiala2002@libero.it

Sra. Rosa JACINTO PEDRO JOÃO PRATA
Técnica do Secretariado do Codex - Angola
Ministério da Agricultura e do Desenvolvimento
Rural
7ª Andar Rua Comandante Gika
Luanda
Phone: +002442 323724
Fax: +002442 323724
Email: secretariado.codex@yahoo.com.br

Mme Teodora LOURENÇO SILVA
Directrice Nationale
Institut Nationale des Normes et Qualité
Ministère de l'Industrie
Rua Cerqueira Lukoki
N° 25, 7A
Luanda
Angola
Email: ianorq@netangola.com

M Carlos Alberto AMARAL
Conseiller
Représentant permanent suppléant auprès de la FAO
Ambassade de la République d'Angola
Via Druso, 39
00184 Rome

Ms Maria PACAVIRA
Ambassade de la République d'Angola
Via Druso, 39
00184 Rome

**ANTIGUA AND BARBUDA –
ANTIGUA-ET-BARBUDA –
ANTIGUA Y BARBUDA**

Mrs Solange A. BAPTISTE
Manager Information Services
Antigua and Barbuda Bureau of Standards
Redcliffe Street and Corn Alley
P.O. Box 1550
St. John's
Phone: +268 562 4011
Fax: +268 462 1625
Email: codex@antigua.gov.ag
abbs@antigua.gov.ag

ARGENTINA - ARGENTINE

Sra. Gabriela Alejandra CATALANI
Coordinadora del Punto Focal del CODEX
Secretaría de Agricultura, Ganadería, Pesca y
Alimentos
Ministerio de Economía y Producción
Paseo Colón 922
B.P. of 29
Buenos Aires
Phone: +54 11 4349 2549/2747
Fax: +54 11 4349 2549
Email: gcatal@mecon.gov.ar
codex@mecon.gov.ar

Sra. María del Carmen SQUEFF
Consejero
Representante Permanente Alternativa ante la FAO
Embajada de la República Argentina
Piazza dell'Esquilino 2
00185 Roma

Sr Luis Eduardo ECHANIZ
Responsable del Área de Internacionales
Servicio Nacional de Sanidad y Calidad
Agroalimentaria
Buenos Aires
Phone: +54 11 41215334
Fax: +54 11 41215360
Email: eechaniz@senasa.gov.ar

Sr Carlos E. CAPPARELLI
Director Ejecutivo
Cámara de la Industria Argentina de Fertilizantes y
Agroquímicos
(CIAFA)
Rivadavia 1367 - 7° "B"
(C1033 AAD) Buenos Aires
Phone: +5411 4381 2742
Fax: +5411 4383 1562
Email: carloscapparelli@ciafa.org.ar

ARMENIA - ARMÉNIE

Ambassador Zohrab V. MALEK
Permanent Representative to FAO
Permanent Representation of the Republic of
Armenia to FAO
Via Camillo Sabatini 102
00100 Rome

Mrs Gayane NASOYAN
Codex Contact Point
Ministry of Agriculture
Government Building #3
Republic Square
375010 Yerevan
Phone: +374 10 565871/525453
Fax: +374 10 524610/565871
Email: gayane@agrounit.am
agrounit@xter.net

AUSTRALIA - AUSTRALIE

Mr Steve MCCUTCHEON
Executive Manager
Product Integrity, Animal and Plant Health
Australian Government Department of Agriculture,
Fisheries and Forestry
GPO Box 858
CANBERRA, ACT, 2601
Australia
Phone: +61 2 6272 4316
Fax: +61 2 6272 5697
Email: steve.mccutcheon@daff.gov.au

Mr Greg READ
Executive Manager, Exports
Australian Quarantine and Inspection Service
Australian Government Department of Agriculture,
Fisheries and Forestry
GPO Box 858
CANBERRA ACT 2601
Australia
Phone: +61 2 6272 3594
Fax: +61 2 6272 4112
Email: gregory.read@daff.gov.au

Dr Paul BRENT
Acting Chief Scientist
Food Standards Australia New Zealand
P.O. Box 7186
CANBERRA BC ACT 2610
Australia
Phone: +61 2 6271 2222
Fax: +61 2 6271 2261
Email: Paul.brent@foodstandards.gov.au

Ms Ann BACKHOUSE
 Manager, Codex Australia
 Product Integrity, Animal and Plant Health
 Australian Government Department of Agriculture,
 Fisheries and Forestry
 GPO Box 858
 CANBERRA ACT 2601
 Australia
 Phone: +61 2 6272 5692
 Fax: +61 2 6272 3103
 Email: ann.backhouse@daff.gov.au

Ms Judy BARFIELD
 Counsellor Agricultural Affairs
 Alternate Permanent Representative to FAO
 Embassy of Australia
 Via Antonio Bosio, 5
 00161 Rome
 Phone: +39 06 85272376
 Fax: +39 06 85272346
 Email: judy.barfield@dfat.gov.au

Mr Richard FARLEIGH
 Administrator
 Australian Government Department of Agriculture,
 Fisheries and Forestry
 GPO Box 858
 CANBERRA ACT 2601
 Australia
 Phone: +61 2 6272 4802
 Fax: +61 2 6272 3307
 Email: richard.farleigh@daff.gov.au

AUSTRIA - AUTRICHE

Mr Dieter JENEWEIN
 Ministry of Health, Family and Youth
 Radetzkystr.2
 A2020 Vienna, Austria
 Phone: +43 6648398030
 Email: dieter.jenewein@ages.at

Dr Erhard HÖBAUS
 Head of Division "Nutrition and Quality Assurance"
 Federal Ministry of Agriculture, Forestry,
 Environment and Water Management
 A-1012 Vienna, Stubenring 12
 Phone: + 431 71100-2855
 Fax: +431 71100-2901
 Email: erhard.hoebaus@lebensministerium.at

BELGIUM - BELGIQUE - BÉLGICA

M Charles CRÉMER
 Directeur
 Service Public Fédéral, Santé publique
 Sécurité de la chaîne alimentaire et Environnement
 DG Animaux, Végétaux et Alimentation
 Place Victor Horta, 40 Boîte 10
 Bloc II -7^e étage
 1060 Bruxelles
 Phone: +32 (0) 2 5247371
 Fax: +32 (0) 2 5247399
 Email: codex.be@health.fgov.be

M Marc CORNELIS
 Conseiller général
 Agence fédérale pour la Sécurité de la Chaîne
 alimentaire
 DG Politique de contrôle - Relations internationales
 WTC III - 19^eme étage
 Boulevard Simon Bolivar, 30
 1000 Bruxelles
 Phone: +32 2 2083834
 Fax: +32 2 2083823
 Email: marc.cornelis@favv.be

M Johan HALLAERT
 Directeur Politique Alimentaire
 Fédération belge des Industries alimentaires (FEVIA)
 Avenue des Arts, 43
 1040 Bruxelles
 Phone: +32 2 5501760
 Fax: +32 2 5501754
 Email: jh@fevia.be

M Guido KAYAERT
 Vice-President
 Relations with the European Institutions
 Nestle Coordination Center
 Rue de Birmingham, 221
 1070 Bruxelles
 Phone: +32 2 5295330
 Fax: +32 2 5295667
 Email: guido.kayaert@be.nestle.com

M Luc OGIERS
 Directeur
 SPF Economie
 City Atrium C
 Rue de Progrès 50
 1210 Bruxelles
 Phone: 0032 2 2777481
 Fax: 0032 2 277 5304
 Email: luc.ogiers@economie.fgov.be

BENIN - BÉNIN

Dr Ahoukpo Denis MIKODE
 Directeur et Secrétaire Permanent
 Direction, de l'alimentation et de la nutrition
 Appliquée
 Secrétariat permanent du comité national du Codex
 Alimentarius
 Ministère de l'agriculture, de l'élevage, et de la pêche
 Boîte postale 03-2900
 Cotonou
 Phone: +229 20212670
 Fax: +229 20213963
 Email: admikode@yahoo.fr

BHUTAN - BHOUTAN - BHUTÁN

Mr Thuji TSHERING
 Chief Regulatory and Quarantine Officer
 Bhutan Agriculture and Food Regulatory Authority
 (BAFRA)
 Ministry of Agriculture
 P.O. Box #1071
 Royal Government of Bhutan
 Thimphu
 Phone: +9752 327031
 Fax: +9752 327032
 Email: t_tshering@moa.gov.bt

Mr Kinga WANGDI
 Chief Human Resource Officer
 Human Resource Management Division
 Ministry of Agriculture
 P.O. Box #1071
 Royal Government of Bhutan
 Thimphu
 Phone: +975 2 335647
 Fax: +975 2 323841
 Email: kin_wangdi@moa.gov.bt

BOLIVIA - BOLIVIE

Sra. Isabel CADIMA PAZ
 Consejera
 Representante Permanente Adjunto ante la FAO
 Embajada de la República de Bolivia
 Via Brenta 2a - Int. 28
 00198 Roma

BOTSWANA

Dr Bernard BULAWAYO
 Principal Research Scientist (Biotechnology)
 National Food Technology Research Centre
 Private Bag 008
 Kanye
 Botswana
 Phone: +267 5440441
 Fax: +267 5440713
 Email: bernard@naftec.org
 bernadtbml@yahoo.co.uk

BRAZIL - BRÉSIL - BRASIL

Ms Ligia Maria SCHERER
 Minister Counsellor
 Deputy Permanent Representative to FAO
 Permanent Representation of the Federative Republic
 of Brazil to FAO
 Via di Santa Maria dell'Anima 32
 00186 Rome
 Phone: + 39 06 68307576
 Fax: +39 06 68398802

Mr Felipe Haddock Lobo GOULART
 Secretary
 Alternate Permanent Representative to FAO
 Permanent Representation of the Federative Republic
 of Brazil to FAO
 Via di Santa Maria dell'Anima 32
 00186 Rome
 Phone: +39 06 678 9353/68307576
 Fax: +39 06 6839 8802

Mrs Maria Aparecida MARTINELLI
 Coordinator of Brazilian Codex Committee
 National Institute for Metrology, Standardization and
 Industrial Quality - INMETRO
 Ministry of Development, Industry and Trade
 SEPN 511, edificio Bittar III, Bloco B, 4º Andar
 70 750-527 Brasília, DF
 Brazil
 Phone: +55 61 3340 2211
 Fax: +55 6161 3347 3284
 Email: codexbrasil@inmetro.gov.br

Mrs Antonia Maria AQUINO
 Manager of Especial Products
 Agência Nacional de Vigilância Sanitária (ANVISA)
 SEPN Av. W3 Norte QD
 511 Bloco A Edifício Bittar II
 Brasília DF
 Brasil
 CEP 70750 541
 Phone: +55 61 448 6289
 Fax: +55 61 448 6472
 Email: antonia.maria@anvisa.gov.br

Ms Renata Campos MOTTA
 Specialist in Regulation and Health Surveillance
 Agência Nacional de Vigilância Sanitária (ANVISA)
 SEPN 515 Ed. Ômega Bloco B
 70770502 Brasília DF
 Brasil
 Phone: +55 61 3448 1078
 Fax: +55 61 3448 1089
 Email: renata.motta@anvisa.gov.br
 rel@anvisa.gov.br

Mrs Denise de Oliveira RESENDE
 National Health Surveillance Agency
 SEPN 511, BL A, Ed. Ômega
 70770-541 Brasília DF
 Brazil
 Phone: + 55 61 3448 6091
 Fax: +55 61 3448 6274
 Email: denise.resende@anvisa.gov.br

Mr Francisco Frederico Sparenberg OLIVEIRA
 Assessor da Diretoria Executiva
 Brazilian Association of Technical Standards-ABNT
 Rua Minas Gerais, 190 Higienópolis
 01244-010 São Paulo
 Brazil
 Phone: +55 11 30173655
 Fax: +55 11 30173633
 Email: ffsoliveira@abnt.org.br

Dr Antonio Marcos PUPIN
 ABIA Brazilian Association of Food Industries
 Av. Brigadeiro Faria Lima
 1478 11° andar
 São Paulo SP brazil
 CEP 01451 008
 Phone: +41 21 924 4277
 Fax: +41 21 924 4547
 Email: marcos.pupin@nestle.com

BULGARIA - BULGARIE

Mr Krassimir KOSTOV
 Minister Plenipotentiary
 Permanent Representative to FAO
 Permanent Representation of the Republic of
 Bulgaria to FAO
 Via Pietro Paolo Rubens, 21
 00197 Rome
 Phone: +39 06 321 3986
 Fax: +39 06 322 6122
 Email: kikostov@yahoo.com

BURKINA FASO

Boubakar Cisse
 Conseiller économique
 Représentant permanent adjoint auprès de la FAO
 Ambassade du Burkina Faso
 Via XX Settembre, 86
 00187 Rome

BURUNDI

M Nestor BIKORIMANA
 Directeur
 Bureau Burundais de Normalisation et Contrôle de la
 Qualité
 Point de contact du codex alimentarius
 21, Boulevard de l'Indépendance
 Bujumbura
 Phone: +257 22221815/257 77730585
 Fax: +257 22225595
 Email: bbn@onatel.bi
 bikonest@yahoo.fr

CAMEROON - CAMEROUN - CAMERÚN

M Michael TABONG KIMA
 Ambassadeur
 Représentant permanent auprès de la FAO
 Ambassade de la République du Cameroun
 Via Siracusa 4-6
 00161 Rome
 Phone: +39 06 44291285
 Fax: +39 06 44291323

M MOUNGUI MÉDI
 Conseiller
 Représentant Permanent Adjoint auprès de la FAO
 Ambassade de la République du Cameroun
 Via Siracusa 4-6
 00161 Rome
 Phone: +39 06 44291285
 Fax: +39 06 44291323
 Email: medimoungui@virgilio.it

Mr Boniface Charles BOOTO À NGON
 Chef de Division de la normalisation et de la qualité
 Ministère de l'Industrie, des Mines et du
 Développement Technologique
 Yaoundé
 Phone: +00237 993 7621
 Fax: +00237 222 6496
 Email: bootangon@yahoo.fr

M Roger TCHAPET
 Métrologue
 Inspecteur Principal des Prix, Poids et Mesures
 Chef de Cellule de la Normalisation et de la
 Protection du Consommateur
 Ministère du Commerce
 B.P. 501
 Yaoundé
 Phone: 222 04 79
 Email: tchapiro06@yahoo.fr

M Philippe FOU DA TSILLA
 Chef de Service des Institutions Spécialisées de
 l'ONU à Caractère Économique et Technique
 Direction des Nations Unies et de la Coopération
 Décentralisée
 Ministère des relations extérieures
 B.P.15 855
 Yaoundé
 Phone: +237 99785978
 Email: philfouda2002@yahoo.fr

CANADA - CANADÁ

Ms Debra BRYANTON
 Executive Director
 Food Safety Directorate
 Canadian Food Inspection Agency
 159 Cleopatra Drive
 Ottawa, Ontario, K1A 0Y9
 Phone: +613 221-7155
 Fax: +613 221-7295
 Email: dbryanton@inspection.gc.ca

Ms Janet BEAUVAIS
 Director-General, Food Directorate
 Health Canada
 251 Frederick Banting Driveway
 (Banting Building #22)
 Room E237, Address Locator 2202E
 Ottawa, Ontario, K1A 0L2
 Phone: +613 957-1821
 Fax: +613 957-1784
 Email: janet_beauvais@hc-sc.gc.ca

Mr Ron BURKE
 Codex Contact Point for Canada
 Director, Bureau of Food Regulatory, International
 and Interagency Affairs
 Food Directorate, Health Canada
 200 Tunney's Pasture Driveway
 Room 2395 (0702C1)
 Ottawa, Ontario, K1A 0L2
 Phone: +613 957-1748
 Fax: +613 941-3537
 Email: ronald_burke@hc-sc.gc.ca

Dr Anne MACKENZIE
Science Advisor
Canadian Food Inspection Agency
159 Cleopatra Drive, Room 113
Ottawa, Ontario, K1A 0Y9
Phone: +613 221-7084
Fax: +613 221-7010
Email: amackenzie@inspection.gc.ca

Mr Allan MCCARVILLE
Senior Adviser, Codex
Bureau of Food Regulatory, International and
Interagency Affairs
Food Directorate, Health Canada
200 Tunney's Pasture Driveway
Room 2394 (0702C1)
Ottawa, Ontario, K1A 0L2
Phone: +613 957-0189
Fax: +613 941-3537
Email: allan_mccarville@hc-sc.gc.ca

Mr Bertrand GAGNON
Manager, Intergovernmental and International
Coordination
Canadian Food Inspection Agency
159 Cleopatra Drive
Ottawa, Ontario, K1A 0Y9
Phone: +613 221-7161
Fax: +613 221-7295
Email: bgagnon@inspection.gc.ca

Ms Angela BEHBOODI
Senior Trade Policy Officer
Department of Foreign Affairs and International
Trade
125 Sussex Drive
Ottawa, Ontario, K1A 0G2
Phone: +613 944-2100
Fax: +613 943-0346
Email: angela.behboodi@international.gc.ca

Mr Karl DUPUIS
Deputy Director
Multilateral Technical Trade Issues Division
Agriculture and Agri-Food Canada
Sir John Carling Building
930 Carling Ave., 10 th Floor, Room 10109
Ottawa, Ontario, K1A 0C5
Phone: + 613 759-7660
Fax: + 613 759 7503
Email: dupuisk@agr.gc.ca

Mr Réjean BOUCHARD
Assistant Director
Policy and Dairy Production
Dairy Farmers of Canada
75 Albert Street, Suite 1101
Ottawa, Ontario K1P 5E7
Phone: +613 236 9997
Fax: +613 236 0905
Email: rejeanb@dfc-plc.ca

CAPE VERDE - CAP-VERT - CABO VERDE

M. José Eduardo DANTAS FERREIRA BARBOSA
Ambassadeur
Représentant permanent auprès de la FAO
Ambassade de la République du Cap-Vert
Via Giosué Carducci 4 - Int. 3
00187 Rome

Mme Maria Goretti SANTOS LIMA
Conseiller
Représentant permanent adjoint auprès de la FAO
Ambassade de la République du Cap-Vert
Via Giosué Carducci 4 - Int. 3
00187 Rome

CENTRAL AFRICAN REPUBLIC – RÉPUBLIQUE CENTRAFRICAINE – REPÚBLICA CENTROAFRICANA

M Denis SAPOUA
Chargé de Mission en Matière d'élevage
Ministère du développement rural
B.P. 786
Bangui
Phone: +236 050106
Fax: +236 61197/236 611997
Email: dsapoua@yahoo.com

CHILE - CHILI

Sra. Antonieta URRUTIA ANABALON
Ing. Agrónomo
Servicio Agrícola y Ganadero
Ministerio de Agricultura
Av. Bulnes 140
Santiago
Chile
Phone: +56 2 345 1585
Fax: +56 2 345 1578
Email: antonieta.urrutia@sag.gob.cl

Sr Gonzalo RÍOS KANTOROWITZ
Ing Agrónomo
Servicio Agrícola y Ganadero
Ministerio de Agricultura
Av. Bulnes 140
Santiago
Chile
Phone: + 56 2 345 1581
Fax: +56 2 345 1578
Email: gonzalo.rios@sag.gob.cl

Sra. Gisela Angélica RODRÍGUEZ RIDEAU
Representante de la Empresa Privada
Santiago
Chile
Phone: +56 2 3384235
Fax: +56 2 3384257
Email: gisela.rodriiguez@nestle.com

CHINA - CHINE

Mr WANG Xuening
Deputy Director-General
Bureau of Health Inspection and Supervision
Ministry of Health
No 1., Nanlu Xizhimenwai
Beijing 100044
People's Republic of China
Phone: +86 10 68792188
Fax: +86 10 68792387
Email: xuening0325@sina.com

Mr DONG Hongyan
Deputy Director
Department of Market and Economy Information
Ministry of Agriculture
11 Nongzhanguan Nanli
Beijing 100026
People's Republic of China
Phone: +86 10 64193156
Fax: +86 10 64193315
Email: scszlc@agri.gov.cn

Mr ZHANG Jinjing
Deputy Director-General
Department of Food Safety Coordination
State Food and Drug Administration
A38 Beilishilu Beijing 100810
People's Republic of China
Phone: +010 88375679
Fax: +010 88375679
Email: jjzh26@yahoo.com

Mrs ZHAO Yanxia
Director
WTO Affairs Department
Ministry of Commerce
No.2 Dong Chang An Street
Beijing 100731
People's Republic of China
Phone: +0086 10 65197242
Fax: +0086 10 65197061
Email: zhaoyanxia@mofcom.gov.cn

Mr CHAN YUK TAK, Eddy
Director of Food and Environment Hygiene
Food and Environmental Hygiene Department
45/F, Queensway Government Offices
66 Queensway, Hong Kong
Phone: +852 2867 5333/9267 2283
Fax: +852 2524 1977
Email: eddy_yt_chan@fehd.gov.hk

Ms XU Jiao
Associate Professor
National Center of Health Inspection and Supervision
Bureau of Health Inspection and Supervision
Ministry of Health
No 1., Nanlu Xizhimenwai
Beijing 100044
People's Republic of China
Phone: +86 1068792404
Fax: +86 10 68792408
Email: xujiaobj@hotmail.com

Mr CHEN Junshi
Professor
National Institute of Nutrition and Food Safety
China CDC
Ministry of Health
29, Nanwei Road
Xuanwu District
Beijing 100050
People's Republic of China
Phone: +86 10 83132922
Fax: +86 10 83132922
Email: jshchen@ilsichina.org

Mrs LIU Xiumei
Chief Scientist
National Institute of Nutrition and Food Safety
China CDC
Ministry of Health
No.7, Panjiayuan
Beijing 100021
People's Republic of China
Phone: +86 10 67791259
Fax: +86 10 67711813
Email: xiumeiliu@ccfa.cc
xmliu01@yahoo.com.cn

Mrs TIAN Jing
Research Assistant
National Institute of Nutrition and Food Safety
China CDC
Ministry of Health
No. 7, Panjiayuan Nanli
Beijing 100021
People's Republic of China
Phone: +86 10 67791259
Fax: +86 10 67711813
Email: tianjing@ccfa.cc
tianjing960928@126.com

Mr YE Anping
Director
Department of International Cooperation
Ministry of Agriculture
11 Nongzhanguan Nanli
Beijing 100026
People's Republic of China
Phone: +86 10 64193339
Fax: +86 10 65003621
Email: yeanping@agri.gov.cn

Dr YE Zhihua
Director General
Institute of Quality Standards and Testing
Technology for Agri-Products
Chinese Academy of Agricultural Sciences
12 South Street
Zhongguancun
Haidan District
Beijing 100081
People's Republic of China
Phone: +86 10 68919419
Fax: +86 10 68975099
Email: zhihuaye@mail.caas.net.cn

Mr CHEN Zongmao
Professor
CCPR Chairman
Institute for the Control of Agrochemicals
Ministry of Agriculture
Room 902, Building 18
Maizidian Street,
Chaoyang District
Beijing 100026
People's Republic of China
Phone: +86 1064194246
Fax: +86 10 64194246
Email: zmchen2006@agri.gov.cn

Mr CUI Yehan
Director
Development Center of Science and Technology
Ministry of Agriculture
20 Maizidian Street
Chaoyang District
Beijing 100026
People's Republic of China
Phone: +86 10 64195082
Fax: +86 10 64194550
Email: cuiyehan@agri.gov.cn

Mr HE Yibing
Professor, Chief Secretary
CCPR Secretariat
Institute for the Control of Agrochemicals
Ministry of Agriculture
Room 902, Building 18
Maizidian Street
Chaoyang District
Beijing 100026
People's Republic of China
Phone: +86 10 64194253
Fax: +86 10 64194252
Email: heyibing@agri.gov.cn

Mrs ZANG Naping
Senior Engineer
Hubei Food and Drug Administration
Donghu Road 136
Wuhan City
Hubei Province
People's Republic of China 430071
Phone: +8627 87253865
Fax: +8627 87253865
Email: zhanglaping-
2004@tom.com/hubshipin@fda.com.cn

Mr NIE Dake
Section Chief
General Administration of Quality Supervision,
Inspection and Quarantine of China (AQSIQ)
No.9 Madian East Road
Haidian District
Beijing 100088
People's Republic of China
Phone: +86 10 82262212
Fax: +86 10 82260385
Email: niedk@aqsiq.gov.cn

Professor QIU Yueming
Deputy Director-General
Department of Agriculture and Food Standard
Standardization Administration of P.R. China
No.9, Madian Donglu
Haidian District
Beijing 100088
People's Republic of China
Phone: +86 10 8226913
Fax: +86 10 82260912
Email: qiuyem@sac.gov.cn

Mr XU Jianjun
Senior Engineer
China National Institute of Standardization (CNIS)
No.4, Zhichun Road
Haidian District
Beijing 100088
People's Republic of China
Phone: +86 10 58811648
Fax: +86 10 58811641
Email: xujj@cnis.gov.cn

Mr ZHOU Chong
Officer
Certification and Accreditation Administration of the
People's Republic of China
9# Madian East Road
Tower B
Haidian District
Beijing 100088
People's Republic of China
Phone: +086 010 82262760
Fax: +086 010 8226 9753
Email: zhouc@cnca.gov.cn

Ms CHAN HON YEE Constance
Assistant Director (Food Surveillance and Control)
Centre for Food Safety
Food and Environmental Hygiene Department
45/F Queensway Government Offices
66 Queensway, Hong Kong
Phone: +852 2867 5511/9032 7882
Fax: +852 2521 9527
Email: constancechan@fehd.gov.hk

Ms HUNG HOI YI Heidi
Administrative Officer (Centre for Food Safety)
Centre for Food Safety
Food and Environmental Hygiene Department
45/F, Queensway Government Offices
66 Queensway, Hong Kong
Phone: + 852 2867 5599/6795 6414
Fax: +852 2530 1368
Email: hhyhungi@fehd.gov.hk

Mr CHU YUEN Keung
Scientific Officer (Risk Assessment)
Centre for Food Safety
Food and Environmental Hygiene Department
43/F, Queensway Government Offices
66 Queensway, Hong Kong
Phone: +852 2867 5606
Fax: +852 2893 3547
Email: jykchu@fehd.gov.hk

COLOMBIA - COLOMBIE

Sra. Blanca Cristina OLARTE PINILLA
 Profesional Especializada
 Ministerio de Protección Social
 CARRERA 13 No.32-76 piso 14
 Phone: +57 1 3365000 ext.1428 1427
 Fax: +57 1 336 5000 ext.1428-330 5050
 Email: bolarte@minproteccionsocial.gov.co

Sr Javier MUÑOZ IBARRA
 Ingeniero Profesional Especializado
 Ministerio de Comercio, Industria y Turismo
 Calle 28 No. 13A-15
 Bogotá D.C.
 Phone: +571 6067676 ext 1205
 Email: jmunoz@mincomercio.gov.co

COSTA RICA

Sra. Isabel Cristina ARAYA BADILLA
 Directora de Mejora Regulatoria y Reglamentación
 Técnica y Coordinadora del Comité Nacional del
 Codex
 Ministerio de Economía, Industria y Comercio
 del Colegio Lincoln 200m oeste, 100m sur y 200m
 oeste
 Edificio IFAM
 Primer Piso
 Moravia, San José
 Phone: +(506) 236 2358
 Fax: +(506) 236-7192
 Email: iaraya@meic.go.cr infocodex@meic.go.cr

Sra Yolanda GAGO
 Representante Permanente Alterno ante la FAO
 Representación Permanente de Costa Rica
 Embajada ante la Santa Sede
 Via G.B. Benedetti, 3
 00197 Roma

Sra. Greta PREDELLA
 Asistente
 Representación Permanente de Costa Rica
 Embajada ante la Santa Sede
 Via G.B. Benedetti, 3
 00197 Roma

CÔTE D'IVOIRE

M Edmond N'DRI APIA
 Directeur des Productions Alimentaires et de la
 Diversification
 Ministère d'État et de l'Agriculture
 B.P. V84 Abidjan
 Phone: +00225 20212481
 Fax: +225 2021 1993
 Email: daq@aviso.ci

Dr Narcisse EHOUSSOU
 Président du Comité national du Codex Alimentarius
 Abidjan
 Phone: +00225 21 353349
 Email: narcehoussou@yahoo.fr

CROATIA - CROATIE - CROACIA

Ms Tea HAVRANEK
 Senior Advisor
 Croatian Codex Contact Point
 Croatian Standards Institute
 Ulica grada Vukovara 78
 10000 Zagreb
 Phone: +381 1 6106005
 Fax: +381 1 6109321
 Email: tea.havrane@hzn.hr

CUBA

Sra. Hortensia Nancy FERNÁNDEZ RODRÍGUEZ
 Directora General
 Oficina Nacional de Normalización (NC)
 Punto de Contacto de Cuba
 Calle E No. 261 esq.13
 Vedado, Plaza
 La Habana
 Cuba
 Phone: +537 830 0879
 Fax: +537 836 8048
 Email: nancy@ncnorma.cu

Don Enrique MORET ECHEVERRÍA
 Embajador
 Representante Permanente ante la FAO
 Misión Permanente de Cuba ante los Organismos de
 Naciones Unidas en Roma
 Via Licinia 13/A
 00153 Roma
 Phone: +39 06 5781123
 Fax: +39 06 5780614
 Email: faoprcub@miscuba.191.it

Dra. Marisa CHAILLOUX LAFFITA
 Directora
 Grupo Nacional Gestión de la Calidad
 Ministerio de Agricultura
 Ave. Independencia y Conill
 Plaza de la Revolución
 La Habana
 Phone: +53 7 884 5415
 Fax: +53 7 881 9464
 Email: lafita@minag.gov.cu

Ing. Gabriel LAHENS ESPINOSA
 Director
 Dirección de Regulaciones Técnicas y Calidad
 Ministerio del Comercio Exterior
 Calle Infanta 16, esq. 23
 Vedado
 La Habana

Ing. Arodis CABALLERO NÚÑEZ
 Director General
 Instituto Cubano de Investigaciones Azucareras
 (ICINAZ)
 Carretera a M. Martínez Prieto Km 2 ½
 Boyeros
 Ciudad de la Habana
 Phone: +537 2609549
 Fax: +537 2609549
 Email: arodisc@icinaz.co.cu

Ing. Pastor PONCE CEBALLO
 Director del CENLAC
 Centro Nacional de Sanidad Agropecuaria (CENSA)
 San Jose de las Lajas
 La Habana
 Phone: +53 47863145
 Email: pastor@censa.edu.cu

CYPRUS - CHYPRE - CHIPRE

Mr Gabriel ODYSSEOS
 Deputy Permanent Representative to FAO
 Permanent Representation of the Republic of Cyprus
 to FAO
 Piazza Farnese, 44
 00186 Rome

Mrs Elena ZACHARIOU
 Industrial Extension Officer
 Ministry of Commerce, Industry and Tourism
 Andrea Araouzou 13-15
 1421 Nicosia
 Phone: +00357 22409305
 Fax: +00357 722753104
 Email: ezachariou@gmail.com

**CZECH REPUBLIC –
 RÉPUBLIQUE TCHÈQUE –
 REPÚBLICA CHECA**

Dr Bibiana JANACKOVA
 Deputy Head of the Food Production and Legislation
 Department
 Ministry of Agriculture
 Tesnov 17
 11705 Praha 1
 Phone: +420 221 812 828
 Fax: +420 221 314 117
 Email: bibiana.janackova@mze.cz

**DEMOCRATIC PEOPLE'S REPUBLIC OF
 KOREA –
 RÉPUBLIQUE POPULAIRE DÉMOCRATIQUE
 DE CORÉE –
 REPÚBLICA POPULAR DEMOCRÁTICA DE
 COREA**

Mr PAK NYONG GIL Jong
 Embassy of the Democratic People's Republic of
 Korea
 Viale dell'Esperanto, 26
 00144 Rome

Mr SUN WON Jong
 Counsellor
 Embassy of the Democratic People's Republic of
 Korea
 Viale dell'Esperanto, 26
 00144 Rome
 Phone: +39 06 54220749
 Fax: +39 06 54210090
 Email: permrepun@hotmail.com

**DEMOCRATIC REPUBLIC OF THE CONGO –
 RÉPUBLIQUE DÉMOCRATIQUE DU CONGO –
 REPÚBLICA DEMOCRÁTICA DEL CONGO**

M Floribert DIBWE KALAMBA
 Chef de Division Industries Animales
 Direction de la production et santé animales
 Ministère de l'agriculture, pêche et élevage
 Croisement Blvd du 30 juin - Av. Batetela
 B.P. 8722 Kin 1
 Kinshasa-Gombe
 Phone: +23481 5043580
 Email: dikalambaflor@yahoo.fr

M Innocent MOKOSA MANDENDE
 Ministre Conseiller
 Représentant permanent adjoint auprès de la FAO
 Ambassade de la République démocratique du Congo
 Via Barberini, 3
 00187 Rome

DENMARK - DANEMARK - DINAMARCA

Mr Knud ØSTERGAARD
 Head of Division
 Danish Veterinary and Food Administration
 Mørkhøj Bygade 19
 DK-2860 Søborg
 Phone: +45 33956120
 Fax: +45 33 956001
 Email: koe@fvst.dk

Ms Jytte KJÆRGAARD
 Head of Section
 Danish Veterinary and Food Administration
 Mørkhøj Bygade 19
 DK-2860 Søborg
 Phone: +45 33 956233
 Fax: +45 33 956001
 Email: jk@fvst.dk

Søren SKAFTE
 Minister
 Deputy Permanent Representative to FAO
 Royal Danish Embassy
 Via dei Monti Parioli, 50
 00197 Rome

Mrs Linda JENSEN
 Chief Consultant, Food Scientist
 Danish Meat Association
 Axeltorv 3
 DK-1609 Copenhagen V
 Denmark
 Phone: +45 33 732500
 Fax: +45 33 931023
 Email: LMJ@danishmeat.dk

Mr Jørgen Hald CHRISTENSEN
 Head of Division - Quality and Food Safety
 Danish Dairy Board
 Frederiks Allé 22
 DK-8000 Århus C
 Denmark
 Phone: +45 873 12186
 Fax: +45 873 12001
 Email: jhc@mejeri.dk

**DOMINICAN REPUBLIC –
 RÉPUBLIQUE DOMINICAINE –
 REPÚBLICA DOMINICANA**

Sr Julio César SANTANA DE LEÓN
 Director Ejecutivo
 Dirección General de Normas y Sistemas de Calidad
 (DIGENOR)
 Secretaría de Estado de Industria y Comercio (SEIC)
 Av. México Esq. Leopoldo Navarro
 Santo Domingo
 Phone: +809 686 2205 ext.319
 Fax: +809 688 3843
 Email: digenor@gmail.com

Sra. Yanina GRATEREAUX
 Ministra Consejera
 Representante Permanente Alternante ante la FAO
 Representación Permanente de la República
 Dominicana ante la FAO
 Via Baldassarre Peruzzi, 10 int. 2
 00153 Roma
 Phone: +39 06 9761 3676
 Fax: +39 06 9725 6408
 Email: yaninag2000@hotmail.com

Sr Héctor Manuel MARTÍNEZ FERNÁNDEZ
 Ministro Consejero
 Representante Permanente Alternante ante la FAO
 Representación Permanente de la República
 Dominicana ante la FAO
 Via Baldassarre Peruzzi, 10 int. 2
 00153 Roma
 Phone: +39 06 7613676
 Email: hemafer09@hotmail.com

ECUADOR - ÉQUATEUR

Dr Francisco SALGADO
 Tercer Secretario
 Representante Permanente Alternante ante la FAO
 Embajada de la República del Ecuador
 Via Antonio Bertoloni, 8
 00197 Roma

Sra. Serena VIVIANI
 Embajada de la República del Ecuador
 Via Antonio Bertoloni, 8
 00197 Roma

EGYPT - ÉGYPTE - EGIPTO

Dr Nabih Abdel HAMID IBRAHIM
 Director
 Food Safety Information Center
 Egyptian Organization for Standardization and
 Quality (EOS)
 9 Gamaa St., Giza
 Egypt
 Phone: +02 5735375
 Fax: +02 5712049
 Email: Drnabih@internetegypt.com

Dr Salah EL-DIN HUSSEIN
 Professor of Food Industries
 Faculty of Agriculture
 Cairo University
 Phone: +202 333 75003

Mr Abdel Aziz Mohamed HOSNI ABDEL AZIZ
 Counsellor
 Deputy Permanent Representative to FAO
 Embassy of the Arab Republic of Egypt
 Via Salaria, 267 (Villa Savoia)
 00199 Rome
 Phone: +39 06 8548956
 Fax: +39 06 8542603
 Email: egypt@agrioffegypt.it

Mr Ahmed GABALLA
 Scientific and Regulatory Affairs Manager
 Atlantic Industries
 Free Zone
 Nasr City
 Cairo
 Phone: +202 2767138
 Fax: +202 2718826
 Email: agaballa@mena.ko.com

Eng. Kamal Darwish KAMAL
 Egyptian Organization for Standardization and
 Quality (EOS)
 16 Tadreeb El-Modarrebeen St.,
 Ameriya
 Cairo
 Fax: +202 22845504
 Email: moi@idsc.gov.eg

EL SALVADOR

Sra. María Eulalia JIMÉNEZ DE MOCHI ONORI
 Ministro Consejero
 Representante Permanente ante la FAO
 Embajada de la República de El Salvador
 Via Gualtiero Castellini, 13
 00197 Roma

ERITREA – ÉRYTHRÉE

Mr Yohannes TENSUE
 Alternate Permanent Representative to FAO
 Embassy of Eritrea
 Via Boncompagni, 16 - 3rd Floor
 Rome

ESTONIA - ESTONIE

Mrs Katrin LÕHMUS
Senior Officer
Ministry of Agriculture
Lai 39/41
Tallinn 15056
Estonia
Phone: +372 6256509
Fax: +372 6256210
Email: katrin.lohmus@agri.ee

ETHIOPIA - ÉTHIOPIE - ETIOPIÁ

Mr Aregahegn YIMER
Standards and Metrology Director
Quality and Standards Authority of Ethiopia
P.O. Box 2310
Addis Ababa
Ethiopia
Phone: +0251 1 116460567
Fax: +0251 1 11646088 81
Email: aregahegn@qsae.org

**EUROPEAN COMMUNITY (MEMBER ORGANIZATION) –
COMMUNAUTÉ EUROPÉENNE
(ORGANISATION MEMBRE) –
COMUNIDAD EUROPEA (ORGANIZACIÓN MIEMBRO)**

Mr Michael SCANNELL
Head of Unit
European Commission
Health and Consumer Protection Directorate-General
(SANCO)
Rue Froissart 101 (2/54)
B-1049 Brussels
Phone: +32 2 299 3364
Fax: +32 2 299 8566
Email: Michael.scannell@ec.europa.eu

Dr Jérôme LEPEINTRE
Principal Administrator
European Commission
Health and Consumer Protection Directorate-General
(SANCO)
Rue Froissart 101 (2/62)
B-1040 Bruxelles
Phone: +32 2 299 3701
Fax: +32 2 299 8566
Email: jerome.lepeintre@ec.europa.eu

Dr Eva ZAMORA ESCRIBANO
European Commission
Health and Consumer Protection Directorate-General
(SANCO)
B-1049 Brussels
Phone: +32 2 299 8682
Fax: +32 2 299 8566
Email:
eva-maria.zamora-escribano@ec.europa.eu

Mr Basil MATHIOUDAKIS
Head of Unit "Food Law, Nutrition and Labelling"
European Commission
Health and Consumer Protection Directorate-General
(SANCO)
B-1049 Brussels
Phone: +32 2 2959182
Email: basil.mathioudakis@ec.europa.eu

Dr Jean-François ROCHE
Administrator
European Commission
Health and Consumer Protection Directorate-General
(SANCO)
B-1049 Brussels
Phone: +32 2 29531 41
Email: jean-francois.roche@ec.europa.eu

Mr Miguel GRANERO ROSELL
European Commission
Health Consumer Protection Directorate-General
(SANCO)
B-1049 Brussels
Phone: +32 2 295 8110
Email:
miguel-angel.granerorosell@ec.europa.eu

FIJI – FIDJI

Mrs Miliakere NAWAIKULA
Principal Research Officer
Ministry of Agriculture, Fisheries and Forests
Koronivia Research Station
P.O. Box 77
Nausori
Phone: +679 3477044
Fax: +679 3400262
Email: miliakere.nawaikula@govnet.gov.fj

FINLAND - FINLANDE – FINLANDIA

Mr Matti AHO
Director-General
Ministry of Agriculture and Forestry
P.O. Box 30
00023 Government, FINLAND
Phone: +358-9 1605 3380
Fax: +358-9 1605 4777
Email: matti.aho@mmm.fi

Ms Anne HAIKONEN
Counsellor, Legal Affairs
Ministry of Trade and Industry
P.O. Box 32
00023 Government, FINLAND
Phone: +358-9 1606 3654
Fax: +358-9 1606 2670
Email: anne.haikonen@ktm.fi

FRANCE - FRANCIA

Mme Roseline LECOURT
 Chargée de Mission
 Ministère de l'Economie, des Finances et de
 l'Industrie
 DGCCRF - Télédéc 051
 59, boulevard Vincent Auriol
 75703 PARIS Cedex 13
 France
 Phone: +00 33 144 973470
 Fax: +00 33 144 973037
 Email: roseline.lecourt@dgccrf.finances.gouv.fr

Mme Catherine CHAPOUX
 Ministère de l'agriculture et de la pêche
 DGAL- Mission de coordination sanitaire
 internationale
 251, rue de Vaugirard
 75732 Paris Cedex 15
 Phone: +33 149 558486
 Fax: +33 149 554462
 Email: catherine.chapoux@agriculture.gouv.fr

M Pascal AUDEBERT
 Point de Contact du Codex alimentarius en France
 Premier Ministre -Secrétariat général aux affaires
 européennes (SGAE)
 Secteur AGRAP
 2, boulevard Diderot
 75572 Paris Cedex 12
 Phone: +33 144 871603
 Fax: +33 144 871604
 Email: sgae-codex-fr@sgae.gouv.fr
pascal.audebert@sgae.gouv.fr

Mme Dominique BUREL
 CNIEL
 42, rue de Châteaudun
 75314 Paris Cedex 09
 Phone: +33 149707105
 Fax: +33 142 806345
 Email: dburel-alf@cniel.com

M Jean-Claude ATTALE
 Cargill France SAS
 Director Regulatory Affairs
 18-20, rue des Gaudines
 78108 Saint-Germain -en Laye
 Phone: +33 1 30613741
 Fax: +33 1 396799
 Email: jean-claude_attale@cargill.com

Mme Annie LOC'H
 DANONE Directeur Affaires Réglementaires 17
 boulevard Haussmann
 75009 PARIS
 Phone: +33 1 44 352432
 Fax: +33 1 44 352469
 Email: annie.loch@danone.com

GABON – GABÓN

M Louis Stanislas CHARICAUTH
 Conseiller
 Représentant permanent suppléant auprès de la FAO
 Ambassade de la République gabonaise
 Via San Marino, 36-36A
 00198 Rome
 Email: lscharicauth@yahoo.fr

GERMANY - ALLEMAGNE – ALEMANIA

Mr Gerhard BIALONSKI
 Federal Ministry of Food, Agriculture and Consumer
 Protection
 Rochusstraße 1
 D-53123 Bonn
 Phone: +49(0)228-529 4651
 Fax: +49(0)228-529 4943
 Email: 314@bmvvel.bund.de

Mr Michael HAUCK
 Federal Ministry of Food, Agriculture and Consumer
 Protection
 Mauerstraße 29-32
 D-10117 Berlin
 Phone: +49 (0) 30 20063263
 Fax: +49 (0) 30 20063273
 Email: codex.germany@bmelv.bund.de

Dr Rolf GROSSKLAUS
 Director and Professor
 Federal Institute for Risk Assessment (BfR)
 Thielallee 88-92
 D-14195 Berlin
 Phone: +49 (03) 8412 3230
 Fax: +49 (03) 8412 3715
 Email: rolf.grossklaus@bfr.bund.de

Dr Jörg W. RIEKE
 Managing Director
 German Dairy Association
 Godesberger Allee 157
 D-53175 Bonn
 Phone: +49 (0) 228 9596923
 Fax: +49 (0) 228 371535
 Email: rieke@milchindustrie.de

Mr Thomas KÜTZEMAIER
 Managing Director
 German Dairy Association
 Meckenheimer Allee 137
 D-53115 Bonn
 Phone: +49 (0) 228 982430
 Fax: +49 (0) 228 982 4320
 Email: th.kuetzemeier@vdm-deutschland.de

Dr Michael PACKERT
 Südzucker AG Mannheim/Ochsenfurt
 Gottlieb-Daimler Str.10
 D-68165 Mannheim
 Phone: +49 (0) 621 421573 (571)
 Email: michael.packert@suedzucker.de

Mr Heiner THOFERN
 First Counsellor
 Deputy Permanent Representative to FAO
 Permanent Representation of the Federal Republic of
 Germany to FAO
 Via S. Martino della Battaglia, 4
 00185 Rome

GHANA

Mrs Charlotte Afi OHENE-MANU
 Deputy Executive Director
 Ghana Standards Board
 P.O. Box MB-245
 Accra
 Ghana
 Phone: +233 21 501939
 Fax: +233 21 500092
 Email: cohene-manu@ghanastandards.org

Mr Kwaku NICOL
 Minister Counsellor
 Alternate Permanent Representative to FAO
 Embassy of the Republic of Ghana
 Via Ostriana, 4
 00199 Rome

Mr Kwamina VAN-ESS
 Deputy Chief Executive (Food Division)
 Food and Drugs Board
 P.O. Box Ct 1873 Cantonments
 Accra
 Ghana
 Phone: +233 244 653167
 Fax: +233 21 660389
 Email: kwaminav@yahoo.com
 fdb@ghana.com

Mrs Genevieve BAAH
 Codex Contact Point Officer
 Ghana Standards Board
 P.O. Box MB-245
 Accra
 Ghana
 Phone: +233 021 501937
 Fax: +233 021 500092/500231
 Email: gsbnep@ghanastandards.org
 gsbnepmails@yahoo.com

Prof. Samuel K. SEFA-DEDEH
 Dean
 Faculty of Engineering Sciences
 University of Ghana
 Accra
 Ghana
 Phone: +233 277 533090
 Email: sefad@ug.edu.gh

Mrs Rubina Ahinei AHINEI AMARTEIFIO
 Board Member
 Food and Drugs Board
 P.O. CT 2783 Cantonments
 Accra
 Phone: +233 21 660489
 Fax: +233 21 660389
 Email: fdb@ghana.com

GREECE - GRÈCE – GRECIA

Dr Eleni PAPANTONIOU
 Head of Unit
 Unit of Quality Standards, National Codex Contact
 Point
 Directorate of Nutrition Policy and Research
 Ministry of Development - Hellenic Food Authority
 (EFET)
 124 Kifisias Av&2 Iatridou Str
 115 26 Athens
 Phone: +30 210 6971552
 Fax: +30 210 6971650
 Email: epapantoniou@efet.gr

Dr Danai PAPANASTASIOU
 Officer
 Unit of Quality Standards
 Directorate of Nutrition Policy and Research
 Ministry of Development - Hellenic Food Authority
 (EFET)
 124 Kifisias Av&2 Iatridou Str.
 115 26 Athens
 Phone: +30 210 6971660
 Fax: +30 210 6971650
 Email: dpapanastasiou@efet.gr

Mr Vasileios KONTOLAIMOS
 Legal Adviser
 Special Community Law Office
 Ministry of Rural Development and Food
 29 Acharnon
 10439 Athens
 Phone: +30 210 8250307
 Fax: +30 210 8254621
 Email: cohalka@otenet.gr

GUATEMALA

Sr. Francisco Eduardo BONIFAZ RODRÍGUEZ
 Embajador
 Representante Permanente ante la FAO
 Embajada de la República de Guatemala
 Via dei Colli della Farnesina, 128
 00194 Roma
 Phone: +39 06 36381143
 Fax: +39 06 3291639
 Email: embaguante.italia@tin.it

Sra. Ileana RIVERA DE ANGOTTI
 Ministro Consejero
 Representante Permanente Adjunto ante la FAO
 Embajada de la República de Guatemala
 Via dei Colli della Farnesina, 128
 Roma
 Phone: +39 06 36381143
 Fax: +39 06 3291639
 Email: embaguante.italia@tin.it

Sra. Barbara DEVIGUS
 Embajada de la República de Guatemala
 Via dei Colli della Farnesina, 128
 00194 Roma

Sr Rino FINAMORE
Embajada de la República de Guatemala
Via dei Colli della Farnesina, 128
00194 Roma

GUINEA - GUINÉE

Mme Hadja Minte CISSE
Directrice Générale
Institut National de Normalisation et de Métrologie
Ministère de l'Industrie, du Commerce, du Tourisme
et de l'Artisanat
BP 1639 Conakry
Phone: +224 60 299539/64325006
Fax: +224 30 413990
Email: abdoulayetraore05@yahoo.fr

M Abdoulaye TRAORE
Conseiller Economique
Ambassade de la République de Guinée
Via Adelaide Ristori, 9b/13
00197 Rome
Phone: +39 06 8078989
Fax: +39 06 8077588
Email: mintecisse

HAÏTI - HAÏTI - HAÏTÍ

M Carl Benny RAYMOND
Conseiller
Représentant permanent suppléant auprès de la FAO
Ambassade de la République d'Haïti
Via di Villa Patrizi 7 - 7A
00161 Rome

HONDURAS

Sra. Mayra REINA
Representante Permanente Adjunto ante la FAO
Embajada de la República de Honduras
Via Giambattista Vico 40, int. 8
00196 Roma

HUNGARY - HONGRIE - HUNGRÍA

Dr Károlyné SZERDAHELYI
Senior Counsellor
Ministry of Agriculture and Rural Development
Department of Food Industry
P.O. Box 1
1860 Budapest 55
Hungary
Phone: +36 1 301 4110
Fax: +36 1 301 4808
Email: tanya.szerdahelyi@fvm.hu

Dr Mária VÁRADI
Chairperson of Hungarian CC MAS Committee
Central Food Research Institute
Herman Otto út 15
H-1022 Budapest
Hungary
Phone: +36 1 355 8982
Fax: +36 1 292 9853
Email: m.varadi@cfri.hu

ICELAND - ISLANDE - ISLANDIA

Mr Thordur ASGEIRSSON
Director of Fisheries
Directorate of Fisheries
Dalshraun 1
210 Hafnarfjörður
Iceland
Phone: +354 569 7900
Fax: +354 569 7991
Email: thordur@fiskistofa.is

INDIA - INDE

Shri DEBASISH PANDA
Joint Secretary
Ministry of Health and Family Welfare
Government of India
Nirman Bhavan
New Delhi - 110011
India
Fax: +91 11 23061447
Email: debasish.panda@nic.in

Dr R.M. SHUKLA
Deputy-Director (Entomology)
Department of Agriculture and Cooperation
Directorate of Plant Protection, Quarantine and
Storage
Ministry of Agriculture
NH-IV Faridabad
Phone: +91 0129 2429067
Email: rmshukla1977@yahoo.co.in

Ms Indrani KAR
Senior Director and Head - Agriculture, Food
Processing & FMCG
Cofederation of Indian Industry
The Mantosh Sondhi Centre
23, Institutional Area
Lodhi Road, New Delhi: 110003
India
Phone: +91 11 2462 2228
Fax: +91 11 2461 5693
Email: indrani.kar@ciionline.org

INDONESIA - INDONÉSIE

Dr SUNARYA
Deputy Director-General
The National Standardization Agency of Indonesia
Manggala wanabakti Block IV Fl, 4
Jl. Jend gatot Subroto, Senayan
Jakarta 10270
Phone: +62 21 5747043
Fax: +62 21 574045
Email: sps-2@bsn.or.id
sunarya@bsn.or.id

Mr Erizal SODIKIN
Agricultural Attaché
Alternate Permanent Representative to FAO
Embassy of the Republic of Indonesia
Via Campania 55
00187 Rome

Mr Imam HARYONO
 Director of Beverage Industry and Tobacco
 Directorate General of Agro and Chemical Industry
 Ministry of Industry
 Jl. Gatot Subroto Kav.52-53, 17th Floor
 Jakarta 12950
 Indonesia
 Phone: +62 21 5252236
 Fax: +62 21 5252236
 Email: imam_haryono_dr@yahoo.com
 imam-haryono@dprin.go.id

Mrs Sri Irawati SUSALIT
 Director
 Food Product Standardization
 National Agency for Drugs and Food Control
 The National Agency for Drugs and Foods Control
 Jl Percetakan Negara No 23
 Jakarta 10560
 Indonesia
 Phone: +62 21 42875584
 Fax: +62 21 42875580
 Email: iras48@yahoo.com
 subdit_spo@yahoo.com

Dr Nyoman Oka TRIDJAJA
 Director
 Quality and Standardization
 Directorate General of Processing Marketing for
 Agricultural Products
 Ministry of Agriculture
 Jl. Harsono RMNo.3 Jakarta 12550
 Indonesia
 Phone: +62 21 781 5881
 Fax: +62 21 7811486
 Email: ntridjaja@yahoo.com

Mr Wicaksono BOEDIMAN
 Head of Section of Goods and Services
 Standardization
 Directorate of Trade, Industry, Investment and IPR's
 Ministry of Foreign Affairs
 Jl. taman Pejambon No. 6
 Jakarta
 Phone: +62 21 3812133
 Fax: +62 21 3859593
 Email: wicaksono@deplu.go.id

Mr Singgih HARJANTO
 Officer
 Codex Contact Point Secretariat
 The National Standardization Agency of Indonesia
 Manggala Wanabakti Block IV Fl.4
 Jl. jend Gatot Subroto, Senayan
 Jakarta 10270
 Indonesia
 Phone: +62 21 5747043
 Fax: +62 21 5747045
 Email: sps-2@bsn.or.id
 singgih@bsn.or.id

**IRAN (ISLAMIC REPUBLIC OF) –
 IRAN (RÉPUBLIQUE ISLAMIQUE D') –
 IRÁN (REPÚBLICA ISLÁMICA DEL)**

Mr Javad Shakhs TAVAKOLIAN
 Ambassador
 Permanent Representative to FAO
 Permanent Representation of the Islamic Republic of
 Iran to FAO
 Via Aventina, 8
 00153 Rome

IRAQ

Mr Akram H. AL-JAFF
 Ambassador
 Permanent Representative to FAO
 Permanent Representation of the Republic of Iraq to
 FAO
 Via della Camilluccia, 355
 00135 Rome

IRELAND - IRLANDE - IRLANDA

Mr Richard HOWELL
 Agricultural Inspector
 Department of Agriculture and Food
 Agriculture House
 7C Kildare Street
 Dublin 2
 Phone: + 00353 1 6072572
 Fax: +00353 1 6616263
 Email: Richard.howell@agriculture.gov.ie

Ms Joan REGAN
 Assistant Principal Officer
 Food Unit
 Department of Health and Children
 Hawkins House
 Hawkins Street
 Dublin 2
 Dublin 2
 Phone: +00353 1 6354247
 Fax: +00353 1 6354552
 Email: Joan_regan@health.irlgov.ie

Mr Padraic DEMPSEY
 First Secretary
 Deputy Permanent Representative to FAO
 Embassy of Ireland
 Piazza di Campitelli, 3
 00186 Rome

ITALY - ITALIE - ITALIA

Amb. Romualdo BETTINI
 Ambassadeur
 Représentant permanent auprès de la FAO
 Représentation permanente de l'Italie après de la
 FAO
 Piazza Margana, 19
 00186 Rome

Drssa Brunella LO TURCO
Segretaria Generale Comitato Nazionale per il
"Codex Alimentarius"
Ministero delle Politiche Agricole, Alimentari e
Forestali
Direzione Generale per la Qualità dei Prodotti
Agrolimentari
Via XX Settembre, 20
00187 Roma
Phone: +39 06 46656041
Email: gpa6@politicheagricole.it

Dr Piergiuseppe FACELLI
Direttore dell' Ufficio III
Rapporti Internazionali del Dipartimento per la
Sanità Pubblica Veterinaria, La Nutrizione e la
Sicurezza degli Alimenti
Ministero della Sanità
Roma
Phone: +39 06 5994 6613
Email: pg.facelli@sanita.it

Dr Ciro IMPAGNATIELLO
Funzionario
Segretariato Generale Comitato Nazionale per il
"Codex Alimentarius"
Ministero delle Politiche Agricole, Alimentari e
Forestali
Direzione Generale per la Qualità dei Prodotti
Agrolimentari
Via XX Settembre, 20
00187 Roma
Phone: +39 06 46656046
Email: c.impagnatiello@politicheagricole.it

Dr Luca RAGAGLINI
Ministero delle Politiche Agricole, Alimentari e
Forestali
Direzione Generale per la Qualità dei Prodotti
Agrolimentari
Via XX Settembre, 20
00187 Roma

Drssa Paola MERCIARO
Ministero delle Politiche Agricole, Alimentari e
Forestali
Direzione Generale per la Qualità dei Prodotti
Agrolimentari
Via XX Settembre, 20
00187 Roma

Mme Sarah GUIZZARDI
Dirigente 1° Livello
Direzione Generale Sicurezza Alimenti
Ministero della Salute
P.zzale Marconi 25
Roma

Drssa Jessyama FORLINI
Ministero delle Politiche Agricole, Alimentari e
Forestali
Via XX Settembre, 20
Roma

JAMAICA - JAMAÏQUE

Dr Camella RHONE
Executive Director
Bureau of Standards Jamaica
6 Winchester Rd
Kingston 10
Jamaica
Phone: +1 876 926 3140
Fax: +1 876 929 4736
Email: crhone@bsj.org.jm

Dr Linnette PETERS
Director Veterinary Public Health
Ministry of Health
2-4 King Street
Kingston
Jamaica
Phone: +876 9671100
Fax: +876 9671280
Email: petersL@moh.gov.jm

Mrs Sheila Yvonne HARVEY
Chief Plant Quarantine Produce Inspector
Ministry of Agriculture and Lands
Plant Quarantine Produce Inspection Division
193 Old Hope Road
Kingston 6
Jamaica
Phone: +876 9770637
Fax: +876 9776992
Email: sharvey@moa.gov.jm
sheharv@yahoo.com

Mrs Fay ANDERSON
Manager of Inspectorate Division
Bureau of Standards Jamaica
6 Winchester Rd
Kingston 10
Jamaica
Phone: +1 876 9263140
Fax: +876 929-4736
Email: fanderson@bsj.org.jm

Dr Michelle HAMILTON
Lecturer
Department of Basic Medical Sciences
Biochemistry Section
University of the West Indies
Mona, Kingston 7
Jamaica
Phone: +1 876 92716
Fax: +1 876 9784737
Email: michelle.hamilton@uwimona.edu.jm

JAPAN - JAPON - JAPÓN

Mr Keiichi NAKABAYASHI
Counsellor
Minister's Secretariat
Ministry of Health, Labour and Welfare
1-2-2 Kasumigaseki
Chiyoda-ku
Tokyo 100 9816
Japan
Phone: +81 3 3595 2326
Fax: +81 3 3503 7965
Email: codexj@mhlw.go.jp

Mr Katsutoshi SARUTA
Coordinator
Food Safety Commission Secretariat
Cabinet Office
Prudential Tower 6F 2-13-10
Nagata-cho
Chiyoda-ku
Tokyo 100-8989
Japan
Phone: +81 3 5251 9138
Fax: +81 3 3591 2236
Email: katsutoshi.saruta@cao.go.jp

Ms Chieko IKEDA
Director
Office of International Food Safety
Policy Planning and Communication Division
Department of Food Safety
Ministry of Health, Labour and Welfare
1-2-2 Kasumigaseki, Chiyoda-ku
Tokyo 100 8916
Phone: +81 3 3595 2326
Fax: +81 3 3503 7965
Email: codexj@mhlw.go.jp

Ms Kazuko FUKUSHIMA
Deputy Director
Policy Planning and Communications Division
Office of International Food Safety
Department of Food Safety
Ministry of Health, Labour and Welfare
1-2-2 Kasumigaseki, Chiyoda-ku
Tokyo 100-8916
Japan
Phone: +81-3-3595-2326
Fax: +81-3-3503-7965
Email: codexj@mhlw.go.jp

Mr Takahiro MAEDA
Section Chief
Policy Planning and Communication Division
Department of Food Safety
Ministry of Health, Labour and Welfare
Kasumigaseki 1-2-2
Chiyoda-ku
Tokyo 100 8916
Japan
Phone: +81 3 3595 2326
Fax: +81 3 3503 7965
Email: codexj@mhlw.go.jp

Mr Hiroshi YOSHIKURA
Adviser
Department of Food Safety
Ministry of Health, Labour and Welfare
1-2-2 Kasumigaseki, Chiyoda-ku
Tokyo, 100-8916
Japan
Phone: +81-3-3595-2326
Fax: +81-3-3503-7965
Email: codexj@mhlw.go.jp

Mr Masahiro MIYAZAKO
Associate Director
International Affairs Division, Food Safety and
Consumer Affairs Bureau
Ministry of Agriculture, Forestry and Fisheries
1-2-1 Kasumigaseki, Chiyoda-ku
Tokyo, 100-8950
Japan
Phone: +81 3 3502 8732
Fax: +81 3 3507 4232
Email: masahiro_miyazako@nm.maff.go.jp

Ms Yoko TAKESHITA
International Affairs Division, Food Safety and
Consumer Affairs Bureau
Ministry of Agriculture, Forestry and Fisheries
1-2-1 Kasumigaseki, Chiyoda-ku
Tokyo, 100-8950
Japan
Phone: +81 3 3502 8732
Fax: +81 3 3507 4232
Email: youko_takeshita@nm.maff.go.jp

Dr Shimmo HAYASHI
Technical Adviser
Japan Food Hygiene Association
1-1-11, Sanwa-cho, Toyonaka
Osaka 561-8588 Japan
Phone: +81 6 6333-0521
Fax: +81 6 6333-3631
Email: shinmo-hayashi@saneigenffi.co.jp

JORDAN - JORDANIE - JORDANIA

Dr Yaseen Muhib KHAYAT
Director-General
Jordan Institution for Standards and Metrology
(JISM)
P.O. Box 941287
Amman 11194
Phone: +00962 6 5680139
Fax: + 00962 6 5681099
Email: ykhayat@jism.gov.jo

Dr Mahmoud ALZU'BI
 Assistant Director-General for Administrative and
 Surveillance Affairs- Director of Standardization
 Department
 Secretary of Jordan National Codex Committee
 Jordan Institution for Standards and Metrology
 (JISM)
 P.O. Box 941287
 Amman 11194
 Phone: +00962 6 5665267
 Fax: +00962 6 5681099
 Email: mzoubi@jism.gov.jo

KENYA

Ms Ann Belinda NYIKULI
 Ambassador
 Permanent Representative to FAO
 Embassy of the Republic of Kenya
 Via Archimede, 164
 00197 Rome

Dr Kioko MANG'ELI
 Managing Director
 Kenya Bureau of Standards
 P.O. Box 54974
 Off Kapiti Road Behind Belleview
 Nairobi 00200
 Kenya
 Phone: +254 020 605490
 Fax: +254 020 609660
 Email: info@kebs.org

Mr Joseph KIMANI MBURU
 Attaché (Agricultural Affairs)
 Alternate Permanent Representative to FAO
 Embassy of the Republic of Kenya
 Via Archimede, 164
 00197 Rome

Mrs Evah ODOUR
 General Manager for Standard Division
 Kenya Bureau of Standards
 P.O. Box 54974 00200
 Kapiti Road behind Belleview
 Nairobi
 Phone: +254 020 605490
 Fax: +254 020 609660/604031
 Email: info@kebs.org
 odourE@kebs.org

Mrs Alice Akoth Okelo ONYANGO
 Acting Head of Codex Office
 Kenya Bureau of Standards
 P.O. Box 54974 off Kapiti Road Behind Belleview
 Nairobi 00200
 Phone: +254 020 605490
 Fax: +254 020 609660
 Email: info@kebs.org
 akothE@kebs.org

Ms Gladys MAINA
 General Manager
 Quality Assurance
 Kenya Plant Health Inspectorate Service (KEPHIS)
 P.O. Box 49542-00100
 Nairobi
 Phone: +254 020 884545
 Fax: +254 020 882265
 Email: director@kephis.org
 gmaina@kephis.org

Mr Samuel OKUCHE
 Senior Public Health Officer
 Ministry of Health
 Box 30016-00100
 Nairobi
 Phone: +254 723559185
 Fax: +0202710055
 Email: okuches@yahoo.com

Mr John Keli WAMBUA
 First Secretary
 Alternate Permanent Representative to FAO
 Embassy of the Republic of Kenya
 Via Archimede, 164
 00197 Rome

KUWAIT - KOWEÏT

Mrs Lamyah Ahmed AL-SAQQAF
 Counsellor
 Permanent Representation of the State of Kuwait to
 FAO
 Viale Aventino 36, int. 8
 00153 Rome

Ms Pamela ALVAREZ
 Permanent Representation of the State of Kuwait to
 FAO
 Viale Aventino 36, int. 8
 00153 Rome

Mr Mohammed AL-WAHAIB
 Permanent Representation of the State of Kuwait to
 FAO
 Viale Aventino 36, int. 8
 00153 Rome

**LAO PEOPLE'S DEMOCRATIC REPUBLIC -
 RÉPUBLIQUE DÉMOCRATIQUE POPULAIRE
 LAO –
 REPÚBLICA DEMOCRÁTICA POPULAR LAO**

Mr Nheune SISAVAD
 Chairman of the Lao Codex Alimentarius Committee
 Department of Intellectual Property Standardization
 and Metrology
 Science Technology and Environment Agency
 Prime Minister's Office
 Nahaiio Rd., P.O. Box2279
 Vientiane
 Laos
 Phone: +856 21 219002
 Fax: +856 21 213472
 Email: nhuene@stea.gov.la
 nheune@hotmail.com

LATVIA - LETTONIE - LETONIA

Mr Maris VALDOVSKIS
Senior Officer
Veterinary and Food Department
Ministry of Agriculture
Republikas laukums 2
Riga, LV-1981 Latvia
Phone: +371 7027196
Fax: +371 7027205
Email: maris.valdovskis@zm.gov.lv

LEBANON - LIBAN - LÍBANO

Mr Habib GHAZIRI
President of the Board
Lebanese Standards Institution- LIBNOR
Sin El-Fil Fouad Chehab av.,
Gedco 3 center Bloc B,
9th Floor
Phone: +9611 485927
Fax: +9611 485929
Email: libnor@libnor.org

LESOTHO

Mrs Mamosala Semakaleng SHALE
First Secretary
Alternate Permanent Representative to FAO
Embassy of the Kingdom of Lesotho
Via Serchio, 8
00198 Rome

LIBERIA - LIBÉRIA

Mrs Musu Jatu RUHLE
Counsellor
Deputy Permanent Representative to FAO
Chargé d'Affaires a.i.
Embassy of the Republic of Liberia
Piazzale delle Medaglie d'Oro, 7
00136 Rome

LITHUANIA - LITUANIE - LITUANIA

Mrs Regina GIRDVAINYTE
Attaché (Agricultural Affairs)
Deputy Permanent Representative to FAO
Embassy of the Republic of Lithuania
Viale di Villa Grazioli, 9
00198 Rome

MALAWI

Mr Charles MALATA-CHIRWA
Director-General
Malawi Bureau of Standards
Moirs Crescent Road
P.O. Box 946, Blantyre
Malawi
Phone: +265 187 0488
Fax: +265 187 0756
Email: charlesmalata@mbsmw.org
mbs@mbsmw.org

MALAYSIA - MALAISIE - MALASIA

Dr Abd Rahim MOHAMAD
Director
Food Safety and Quality Division
Ministry of Health Malaysia
Level 3, Block E7, Parcel E
Federal Government Administration Centre
62590 Putrajaya
Malaysia
Phone: +603 8883 3501
Fax: +603 8889 3815
Email: abd_rahim@moh.gov.my

Mrs Dató Lily ZACHARIAH
Ambassador
Permanent Representative to FAO
Embassy of Malaysia
Via Nomentana 297
00162 Rome
Italy
Phone: +39 06 841 5764
Fax: +39 06 841 5808
Email: mw.rome@embassymalaysia.it

Ms Noraini MOHD. OTHMAN
Deputy Director (Codex)
Food Safety and Quality Division
Department of Public Health
Ministry of Health Malaysia
Level 3, Block E7, Parcel E
Federal Government Administration Centre
62590 Putrajaya
Malaysia
Phone: +603 8883 3500
Fax: +603 8889 3815
Email: noraini_othman@moh.gov.my
noraini_mohdothman@yahoo.co.uk

Ms Shamsinar ABDUL TALIB
Deputy Director (SPS and Regional)
Food Safety and Quality Division
Ministry of Health Malaysia
Level 3, Block E7, Parcel E
Federal Government Administration Centre
62590 Putrajaya
Malaysia
Phone: +603 8883 3508
Fax: +603 8889 3815
Email: shamsinar@moh.gov.my
shamsinar_talib@yahoo.com

Mr Johari BIN RAMLI
Agricultural Attaché
Alternate Permanent Representative to FAO
Embassy of Malaysia
Via Nomentana, 297
00162 Rome
Phone: +39 06 841 9296
Fax: +39 06 856 5110
Email: johari5@hotmail.com

Mr Zamruni MOHD.KHALID
Principal Assistant Secretary
Ministry of Foreign Affairs
Multilateral Economic and Environment Division
Wisma Putra, Precint 2
Pusat Pentadbiran Kerajaan Persekutuan
62602 Putrajaya
Malaysia
Phone: +603 8887 4079
Fax: +603 8889 2843
Email: zamruni@kln.gov.my

Mr Samsuddin NOOR
Principal Assistant Secretary
Ministry of Plantation Industries and Commodities
No.15, level 6-13, Persiaran Perdana, Precint 2
Federal Government Administration Centre
62654 Putrajaya
Malaysia
Phone: +603 8880 3427
Fax: +603 8880 3441
Email: sam@kppk.gov.my

Dr Moktir Singh GARDIR SINGH
Veterinary Officer
Department of Veterinary Services
Ministry of Agriculture and Agro-based Industry
Wisma Tani, Block Podium
Lot 4G1, Precint 4
Federal Government Administration Centre
62630 Putrajaya
Malaysia
Phone: +603 8870 2123
Fax: +603 8888 5755
Email: moktir@jph.gov.my
moktir2001@yahoo.com

Ms Norzidah ABU KHAIR
Assistant Director (Codex)
Food Safety and Quality Division
Ministry of Health Malaysia
Level 3, Block E7, Parcel E
Federal Government Administration Centre
62590 Putrajaya
Malaysia
Phone: +603 8883 3549
Fax: +603 8889 3815
Email: norzidah@moh.gov.my
norzidah@yahoo.com

Dr Salmiah AHMAD
Deputy Director-General Services
Malaysian Palm Oil Board (MPOB)
No. 6, Persiaran Institusi
Bandar Baru Bangi
43000 Kajang, Selangor
Malaysia
Phone: +603 8926 6163
Fax: +603 8926 1329
Email: salmiah@mpob.gov.my

Ms Fauziah ARSHAD
Acting Director
Product Development and Advisory Services
Malaysian Palm Oil Board (MPOB)
No. 6, Persiaran Institusi
Bandar Baru Bangi
43000 Kajang, Selangor
Malaysia
Phone: +603 87694589
Fax: +603 8920 1918
Email: fauziah@mpob.gov.my

Dr Wai Lin SIEW
Senior Research Officer
Malaysian Palm Oil Board (MPOB)
No. 6, Persiaran Institusi
Bandar Baru Bangi
43000 Kajang, Selangor
Malaysia
Phone: +603 8769 4430
Fax: +603 8920 1918
Email: siew@mpob.gov.my

Dr E. Siong TEE
Nutrition Consultant
No. 46, Jalan SS22/32
Damansara Jaya
47400 Petaling Jaya
Selangor
Malaysia
Phone: +603 7728 7287
Fax: +603 7728 7426
Email: president@nutriweb.org.my

Mr Amri ISMAIL
Assistant Agricultural Attaché
Embassy of Malaysia
Via Nomentana, 297
00162 Rome
Phone: +39 06 841 5764
Fax: +39 06 855 5110
Email: agri.aaa@ambasciatamalaysia.191.it

MALI - MALÍ

Prof Boubacar Sidiki CISSE
Conseiller Technique
Ministère de la santé
Bamako
Phone: +223 2225301
Fax: +223 2230203
Email: bscisse@ml.refer.org

M Ousmane TOURE
Directeur Général
Agence Nationale de la Sécurité Sanitaire des
Aliments
Ministère de la Santé
BPE:2362
Quartier du Fleuve
(Ancienne Cour des Grandes Endémies)
Bamako
Mali
Phone: +223 2230183
Fax: +223 2220747
Email: oussou_toure@hotmail.com

M Ibrahim Bocar DAGA
Ambassadeur
Représentant permanent auprès de la FAO
Ambassade de la République du Mali
Via Antonio Bosio, 2
00161 Rome
Phone: +39 06 44254068
Fax: +39 06 44 254029
Email: amb.malirome@tiscalinet.it

M Modibo Mahamane TOURE
Deuxième Conseiller
Représentant permanent adjoint auprès de la FAO
Ambassade de la République du Mali
Via Antonio Bosio, 2
00161 Rome
Phone: +39 06 44254068
Fax: +39 06 44254029
Email: modimah@yahoo.com

MALTA - MALTE

Mr Walter BALZAN
Ambassador
Permanent Representative to FAO
Permanent Representation of the Republic of Malta
to FAO
Via dei Somaschi, 1
00186 Rome

Mr John ATTARD KINGSWELL
Codex Alimentarius National Contact Point
Department of Public Health
Rue d'Argens
Msida, Malta
Phone: +356 21324093/21332225
Fax: +356 21344767
Email: john.attard-kingswell@gov.mt

Ms Ritienne BONAVIA
Alternate Permanent Representative to FAO
Permanent Representation of the Republic of Malta
to FAO
Via dei Somaschi, 1
00186 Rome

MAURITIUS - MAURICE - MAURICIO

Mr Denis CANGY
Alternate Permanent Representative to FAO
Embassy of the Republic of Mauritius
127, rue de Tocqueville, 75017 Paris
c/o Consulate of the Republic of Mauritius
Via G.B. Morgagni 6/A
00161 Rome

MEXICO - MEXIQUE - MÉXICO

Sr Carlos Francisco ARCE MACÍAS
Subsecretario de Normatividad, Inversión Extranjera
y Prácticas Comerciales Internacionales
Secretaría de Economía
Alfonso Reyes 30
C.P. 06140
Col. Hipódromo Condesa
México D.F. Piso 14
Phone: +5255 57299271
Fax: +5255 57299413
Email: carce@economia.gob.mx

Sr Carlos Ramón BERZUNZA SÁNCHEZ
Director de Normalización Internacional
Dirección general de Normas (DGN) de la secretaría
de Economía (SE)
Av. Puente de Tecamachalco 6
Lomas de Temachalco
Sección Fuentes
Naucalpan
Estado de México
Phone: +52 55 57299480
Fax: +52 55 55 209715
Email: cberzunz@economia.gob.mx

Sr Victor D. FLORES
Consejero
Representante Permanente Alterno ante la FAO
Embajada de los Estados Unidos Mexicanos
Via Lazzaro Spallanzani, 16
00161 Roma

Sr Diego Alonso SIMANCAS GUTIÉRREZ
Segundo Secretario
Embajada de los Estados Unidos Mexicanos
Via Lazzaro Spallanzani, 16
00161 Roma

Sra. Rocio del Carmen ALATORRE EDEN-
WYNTER
Comisionada de Evidencia y Manejo de Riesgos
Comisión Federal para la Protección contra Riesgos
Sanitarios (COFEPRIS)
Secretaría de Salud
Av. Monterrey No.33
Piso 9 Colonia Roma
Delegación Cuahtémoc
C.P. 06700
Phone: +52 55 148572
Fax: +52 55 148587
Email: rocioal@salud.gob.mx

Sra. Laura BONILLA REYES
Coordinador Subcomite 8 Etiquetado de los
Alimentos
Canacintra/UNILEVER
Bosques de Toronjos No.150
Col. Bosques de las Lomas C.P. 05120
México, D.F.
Phone: +5255 1105 4487
Fax: +5255 1105 4464
Email: laura.bonilla@unilever.com

Sra. Ingrid MACIEL PEDROTE
Gerente de Normatividad
SIGMA Alimentos
Mariano Escobedo 573 Piso 3
Col. Rincon del Bosque
C.P. 11580
México, D.F.
Phone: +5255 8503 2086
Fax: +5255 8503 2056
Email: imaciel@sigma-alimentos.com

Sr Alfonso MONCADA JIMÉNEZ
Representante
Consejo Agroempresarial de Mesoamerica y el Caribe
Av. División del Norte N° 1419
Col. Santa Cruz Atoyac
México D.F. C.P. 03310
Phone: +52 55 5000 1405
Fax: +52 55 5601 0903
Email: amoji@starmedia.com

Sra. Martha GALICIA ROMERO
Asesor
Consejo Mexicano de la Industria de Productos de Consumo, A.C.
Séneca 65
Colonia Polanco Chapultepec
México, D.F.
C.P. 11560
Phone: +5255 52804335
Fax: +5255 52804335
Email: mgalicia@conmexico.com.mx

Mr Antonio MANTOAN
Assoc. Director Regulatory Affairs Mead Johnson Nutritionals
Av. Revolucion 1267 Col. Tlacopac
México D.F.
C.P. 01049
Phone: +52 55 53372896
Email: antonio.mantoan@bms.com

MOROCCO - MAROC - MARRUECOS

Mme Zakia DRIOUICH
Directrice des Industries de la Pêche Maritime
Département des pêches maritimes
Ministère de l'agriculture, du développement rural et des pêches maritimes
Haut Agdal
Rabat B.P. 476
Phone: +212 37 688295/93
Fax: +212 37 68 82 94
Email: driouich@mpm.gov.ma

M Mohammed MAJDI
Chef de la Division Techniques de la Répression des Fraudes
Ministère de l'agriculture, du développement rural et des pêches maritimes
Rabat B.P. 1308
Phone: +212 37 298150
Fax: +212 37 297544
Email: mohammed_majdi@yahoo.fr

M El-Mâati BENZAOUZ
Laboratoire Officiel d'Analyses et de Recherches Chimiques
Ministère de l'agriculture, du développement rural et des pêches maritimes
25, rue Nichakra Rahal
20000 Casablanca
Phone: +212 22 302196
Fax: +212 22 301972
Email: maatibenazzouz@yahoo.fr

M Mohammed BACHAOUCH
Association Professionnelle des Boissons gazeuses
Rabat

M Mohamed ELJAMALI
Union Nationale des Industries de la Conserve de Poisson (UNICOP)
7, rue El Yarmouk Longchamps
Casablanca
Phone: +212 22 943749
Fax: +212 22 943749
Email: unicop@menara.ma

M Majid JOUNDY
Secrétaire Général
Union Nationale des Industries de la Conserve de Poisson (UNICOP)
7, rue El Yarmouk Longchamps
Casablanca
Phone: +028 84594
Fax: +028 845996/828703
Email: info@belma.ma

M Mohammed HOMMANI
Union Nationale des Industries de la Conserve de Poisson (UNICOP)
7, rue El Yarmouk Longchamps
Casablanca
Phone: +212 22 943749
Email: h.hommani@menara.ma

M Mohamed AIT HMID
Ministre plénipotentiaire
Ambassade du Royaume du Maroc
Via Lazzaro Spallanzani 8-10
Rome

MYANMAR

Mr Hlaing MYINT OO
Alternate Permanent Representative to FAO
Embassy of the Union of Myanmar
Via della Camilluccia, 551
Rome

**NETHERLANDS - PAYS-BAS –
PAÍSES BAJOS**

Mr Ger DE PEUTER
Deputy Director
Department of Food Quality and Animal Health
Ministry of Agriculture, Nature and Food Quality
P.O. Box 20401
2500 EK Den Haag
The Netherlands
Phone: +31 70 3785686
Fax: +31 70 3786177
Email: g.de.peuter@minlnv.nl

Dr Wim VAN ECK
Chief Public Health Officer
Food and Consumer Product Safety Authority
P.O. Box 19506
2500 CM The Hague
The Netherlands
Phone: +31 70 4484814
Fax: +31 70 4484061
Email: wim.van.eck@vwa.nl

Dr Wieke TAS
Senior Policy Officer
Ministry of Health, Welfare and Sport
P.O. Box 20350
2500 EJ The Hague
The Netherlands
Phone: +31 70 340 6365
Fax: +31 70 340 5554
Email: JW.TAS@minvws.nl

**NEW ZEALAND - NOUVELLE-ZÉLANDE -
NUEVA ZELANDIA**

Mr Sundararaman RAJASEKAR
Senior Programme Manager, Codex
Codex Coordinator and Contact Point for New
Zealand
New Zealand Food Safety Authority
P.O. Box 2835
Wellington
New Zealand
Phone: +64 4 894 2576
Fax: +64 4 894 2583
Email: RajasekarS@nzfsa.govt.nz

Dr Steve HATHAWAY
Director, Science
New Zealand Food Safety Authority
P.O. Box 2835
Wellington
Phone: +64 6 867 1144
Fax: +64 6 868 5207
Email: Steve.hathaway@nzfsa.govt.nz

Ms Melissa QUARRIE
Policy Analyst (Codex)
Policy Group
New Zealand Food Safety Authority
Telecom Network House
South Tower, 86 Jervois Quay
Wellington
Phone: +64 4 8942462
Fax: +64 4 894 2583
Email: melissa.quarrie@nzfsa.govt.nz

NIGER - NÍGER

M Illiassou BAKO
Directeur de la normalisation, de la qualité et de la
métrologie
Direction de la Normalisation, de la Qualité et de la
Métrologie
Niamey
Phone: +00227 20736950
Fax: +00227 20732150
Email: bako142002@yahoo.fr

NIGERIA - NIGÉRIA

Mr Edward Dolapo OSUNMAKINDE
Minister
Chargé D'Affaires a.i.
Permanent Representation of the Federal Republic of
Nigeria to FAO
Via Orazio, 14-18
00193 Rome

Mr Yaya Adisa Olaitan OLANIRAN
Minister
Permanent Representative to FAO
Permanent Representation of the
Federal Republic of Nigeria to FAO
Via Orazio, 14-18
00193 Rome

Mrs Margaret E. ESHIETT
Deputy Director
Standards Organization of Nigeria
13/14 Victoria Arobieke Street
Lekki Phase 1
Lagos
Phone: +234 1 2708231-4
Fax: +234 1 2708246
Email: info@sononline-ng.com

Mrs Jane O. NZEOMA
Deputy Director (Regulatory Affairs)
National Agency for Food and Drug Administration
and Control
(NAFDAC)
3/4 Oshodi - Apapa, Expressway
Lagos
Phone: +234 1 4772453(0)/234 1 8033338184
Email: janenzeoma@yahoo.com

Mr Mohammed BELLO-ANDO
Assistant Director
Ministry of Foreign Affairs
Federal Secretariat
Abuja

Mr Salami Henry OLALEKAN
Assistant Controller of Customs
Nigerian Custom Service
Zone 3, Wuse, Abuja
Phone: +080 33334274
Email: sholalekan80@yahoo.com

Mr Charles Emeka NWAGBARA
Assistant Chief Standards Officer
Standards Organization of Nigeria
13/14 Victoria Arobieke Street
Lekki Phase 1
Lagos
Phone: +234 1 2708231-4
Fax: +234 1 2708246
Email: nwagbarace@yahoo.com

Mrs Victoria Fila OJOGBANE
Legal Practitioner
Federal Ministry of Justice
Abuja
Nigeria

NORWAY - NORVÈGE - NORUEGA

Mrs Tone Elisabeth MATHESON
Senior Adviser
Ministry of Agriculture and Food
P.O. Box 8007 Dep
N-0030 Oslo
Phone: +47 22 24 9413
Fax: +47 22 249555
Email: tone.matheson@lmd.dep.no

Mrs Vigdis Veum MOELLERSEN
Advisor
Norwegian Food Safety Authority
Head Office
P.O. Box 383
N-2381 Brummundal
Phone: +47 23 216669
Fax: +47 23 216801

Mr Lennart JOHANSON
Deputy Director-General
Ministry of Fisheries and Coastal Affairs
P.O. Box 8118 Dep
Oslo
Norway
Phone: +47 22242665
Fax: +47 22245678
Email: Lennart.Johanson@fkd.dep.no

Mrs Sissel LYBERG BECKMANN
Deputy Director-General
Ministry of Health and Care Services
P.O. Box 8011 Dep
N-0030 Oslo
Norway
Phone: +47 22 248712
Fax: +47 22 248656
Email: slb@hod.dep.no

Mr Bjørn Røthe KNUDTSEN
Regional Director
Regional Office for Trøndelag, Møre and Romsdal
Norwegian Food Safety Authority
P.O. Box 383
N-2381 Brumunddal
Norway
Phone: +47 74113200
Fax: +47 74113201
Email: bjrkn@mattilsynet.no

OMAN - OMÁN

Mr Rasmi MAHMOUD
Technical Adviser
Embassy of the Sultanate of Oman
Via della Camilluccia, 625
00135 Rome
Phone: +39 06 363 00517
Email: rasmimahmoud@gmail.com

PANAMA - PANAMÁ

Sr Luis BENAVIDES
Jefe del Departamento de Elaboración y Evaluación
de Normas
Autoridad Panameña de Seguridad de Alimentos
Ave. Ricardo J. Alfaro
Sun Towers Mall
2do. piso, local 70
Ciudad de Panamá
Phone: +507 522 0009
Fax: +507 522 0001
Email: aupsa@aupsa.gob.pa

PAPUA NEW GUINEA – PAPOUASIE-NOUVELLE-GUINÉE – PAPUA NUEVA GUINEA

Mr Ian ONAGA
Program Manager/Codex Contact Point
Technical and Field Services Branch
Department of Agriculture and Livestock
P.O Box 2141 Boroko
Papua New Guinea
Phone: +675 3200705/675 3200658
Fax: +675 3211046
Email: onaga_ianonaga@datec.net.pg

PARAGUAY

Dra. Lilian MARTINEZ DE ALONSO
Directora General
Instituto Nacional de Tecnología, Normalización y
Metrología (INTN)
Punto de Contacto del Codex en Paraguay
Avda. Artigas 3973 esquina General Roa
Barrio Santísima Trinidad
C.C. 967 Asunción
Phone: +595 21 297 516
Fax: +595 21 297 516
Email: codex@intn.gov.py

Sra. Liz Haydee CORONEL CORREA
 Consejera
 Representante Permanente Adjunto ante la FAO
 Embajada de la República del Paraguay
 Via Firenze, 43 Scala A, int 17
 00184 Roma
 Phone: +39 06 4741715
 Fax: +39 06 4745473
 Email: embaparoma@virgilio.it

Abog. María Rocio ABED OVIEDO
 Secretaria Ejecutiva de Comité del Codex
 Instituto Nacional de Tecnología, Normalización y
 Metrología (INTN)
 Punto de Contacto del Codex en Paraguay
 Avda. Artigas 3973 esquina General Roa
 Barrio Santísima Trinidad
 C.C. 967 Asunción
 Phone: +595 21 290 873
 Fax: +595 21 290 873
 Email: codex@intn.gov.py

Ing. Luis Daniel FLEITAS BRIZUELA
 Director de Normalización
 Instituto Nacional de Tecnología, Normalización y
 Metrología (INTN)
 Punto de Contacto del Codex en Paraguay
 Avda. Artigas 3973 esquina General Roa
 Barrio Santísima Trinidad
 C.C. 967 Asunción
 Phone: +595 21 290 873
 Fax: +595 21 290 873
 Email: codex@intn.gov.py

Sra. Zully MILLESSI
 Directora de Metrología
 Instituto Nacional de Tecnología, Normalización y
 Metrología (INTN)
 Punto de Contacto del Codex en Paraguay
 Avda. Artigas 3973 esquina General Roa
 Barrio Santísima Trinidad
 C.C. 967 Asunción
 Phone: +595 21 290 873
 Fax: +595 21 290 873
 Email: codex@intn.gov.py

Sr Robert DUARTE
 Jefe del Departamento de Metrología
 Instituto Nacional de Tecnología, Normalización y
 Metrología (INTN)
 Punto de Contacto del Codex en Paraguay
 Avda. Artigas 3973 esquina General Roa
 Barrio Santísima Trinidad
 C.C. 967 Asunción
 Phone: +595 21 290 873
 Fax: +595 21 290 873
 Email: codex@intn.gov.py

Lic. Elsi OVELAR
 Directora Instituto de Alimentación
 Ministerio de Salud Pública y Bienestar Social (MSP
 y BS)
 Phone: +595 21 206 874
 Fax: +595 21 206 874
 Email: directiongeneral_inan@webmail.com.py
 inanpy@hotmail.com

Sra. Sonia Beatriz BIEDERMANN MAYOR
 Tercera Secretaria
 Representante Permanente Alterno ante la FAO
 Embajada de la República del Paraguay
 Via Firenze, 43 Scala A, int 17
 00184 Roma
 Phone: +39 06474 1715
 Fax: +39 06 474 5473
 Email: embaparoma@virgilio.it

Sr Juan Ignacio LIVIERES OCAMPOS
 Ministro de Embajada
 Embajada de la República del Paraguay
 Via Firenze, 43 Scala A, int 17
 00184 Roma
 Phone: +39 06 474 1715
 Fax: +39 06 474 5473
 Email: embajada.paraguay@virgilio.it
 embaparoma@virgilio.it

Ing. Roque Rafael LEGUIZAMON
 Ministerio de Agricultura y Ganadería
 Presidente Franco 475 c/ 14 de Mayo
 Asunción
 Phone: +095 21 582290
 Email: roqueflm@yahoo.com

Lic. Asunción CORONEL
 Directora
 Programas de Nutrición
 Asunción
 Phone: +595 21 294073

Dr Vidal ROMÁN
 Representante
 Ministerio de Relaciones Exteriores
 Pte. Franco y Ayolas P6 OF 610
 Asunción
 Phone: +59521 496786
 Email: vihermosa@mre.gov.py

Sra. Aida ROMERO
 Funcionario de Gobierno
 Asunción

PERU - PÉROU - PERÚ

Ing. Claudia SOLANO ORÉ
 Consultora de Calidad y Normatividad y Miembro
 del Pleno del Comité Nacional del Codex
 Comisión para la Promoción del Perú para la
 Exportación y el Turismo - PROMPERU
 Phone: 511 2221222 anexo 301
 Fax: 511 4213179
 Email: csolano@promperu.gob.pe

PHILIPPINES - FILIPINAS

Mr Noel DE LUNA
 Agricultural Attaché
 Deputy Permanent Representative to FAO
 Embassy of the Republic of the Philippines
 Viale delle Medaglie d'Oro, 112
 00136 Rome

POLAND - POLOGNE - POLONIA

Ms Marta SOBIERAJ
Senior Specialist
Codex Contact Point for Poland
Agricultural and Food Quality Inspection
International Co-operation Department
30 Wspólna Street
00-930 Warsaw
Poland
Phone: +48 22 6232903
Fax: +48 22 6232997
Email: kodeks@ijhars.gov.pl
msobieraj@ijhars.gov.pl

Ms Grazyna MORKIS
Specialist
Institute of Agriculture and Food Economics
National Research Institute
20 Swietokrzyska Street
00-002 Warsaw, Poland
Phone: +48 22 6232903
Fax: +48 22 6232997
Email: kodeks@ijhars.gov.pl

Ms Wioletta KARAS
Specialist
Institute of Agriculture and Food Economics
National Research Institute
20 Swietokrzyska Street
00-002 Warsaw, Poland
Phone: +48 22 6232903
Fax: +48 22 6232997
Email: kodeks@ijhars.gov.pl

PORTUGAL

Dr Luis SALINO
Gabinete de Planeamento e Políticas
Ministério da Agricultura, Desenvolvimento Rural e
Pescas
Rua Padre António Vieira, Nº 1
1099-073 Lisboa
Phone: +00351 213819300
Fax: +00351 213876635
Email: lsalino@gpaa.pt

Mr Antonio DUARTE DE ALMEIDA PINHO
Counsellor
Deputy Permanent Representative to FAO
Embassy of the Portuguese Republic
Viale Liegi, 21
00198 Rome

Mr Kari TÖLLIKKÖ
Principal Administrator
General Secretariat of the Council of the European
Union- The Portuguese Presidency
Rue de la Loi 175
B-1048 Brussels
Belgium
Phone: +32 2 281 7841
Fax: +32 2 281 6198
Email: kari.tollikko@consilium.europa.eu

Ms Rose-Marie ELFQUIST
Secretary
General Secretariat of the Council of the European
Union- The Portuguese Presidency
Rue de la Loi 175
B-1048 Brussels
Belgium
Phone: +32 2 2819508
Fax: +32 2 2817928
Email: rose-marie.elfquist@consilium.europa.eu

QATAR

Soltan Saad S.K. AL-MORAIKHI
Ambassador
Permanent Representative to FAO
Embassy of the State of Qatar
Via Antonio Bosio, 14
Rome

Dr Hassan BAYRAKDAR
Food Expert
Doha Municipality
Ministry of Municipal Affairs and Agriculture
P.O. Box 163
Doha-Qatar
Phone: +971 505537184
Fax: +971 48839880
Email: hassan6264@hotmail.com

Dr Shadi Salah ZEYADAH
Food Consultant
Health Affairs
Doha Municipality
Ministry of Municipal Affairs and Agriculture
P.O. Box 163
Doha-Qatar
Phone: +974 4347540
Fax: +974 4479729
Email: sh.zeyadah@yahoo.com

Hatour AKEEL
Embassy of the State of Qatar
Via Antonio Bosio, 14
Rome

**REPUBLIC OF KOREA –
RÉPUBLIQUE DE CORÉE –
REPÚBLICA DE COREA**

Dr WOO Gun-Jo
Director, Center for Food Safety
Korea Food and Drug Administration
#194 Tongil-ro Eunpyung-gu
Seoul 122-704
Phone: +82 2 380 1685
Fax: +82 2 385 0015
Email: gjwoo@kfda.go.kr

Dr KWAK Hyo Sun
Deputy Director
Korea Food and Drug Administration
#194 Tongil-ro Eunpyung-gu
Seoul 122-704
Phone: +82 2 380 1682
Fax: +82 2 380 1615
Email: kwakhyos@kfda.go.kr

Ms JUNG Seung Yun
Senior Researcher
Korea Food and Drug Administration
#194 Tongil-ro Eunpyung-gu
Seoul 122-704
Phone: +82 2 380 1347
Fax: +82 2 385 2416
Email: jsy0511@kfda.go.kr

Dr LEE Cherl-Ho
Professor
Graduate School of Biotechnology
Korea University
1 Anamdong-5
Sungbukku
Seoul, 136-701
Republic of Korea
Phone: +82 2 3290 3414
Fax: +82 2 927 5201
Email: chlee@korea.ac.kr

Dvm KIM Kyu
Assistant-Director
Bilateral Cooperation Division
International Agriculture Bureau
Ministry of Agriculture and Forestry
Government Gwacheon Complex
Jungang-dong, Gwacheon-City
Gyeonggi-do
Republic of Korea
Phone: +82 2 500 1727
Fax: +82 2 504 6659
Email: kimk@maf.go.kr

Dr HWANG Eun-Sun
Deputy Director
Food Industry Division
Agricultural Marketing Policy Bureau
Ministry of Agriculture and Forestry
Government Gwacheon Complex
Jungang-dong, Gwacheon-City
Gyeonggi-do
Republic of Korea
Phone: +82 2 500 1855
Fax: +82 2 503 7905
Email: ehwang@maf.go.kr

Dr KWAG Sang Ick
Assistant director
Livestock Products Sanitation Division, Livestock
Bureau
Ministry of Agriculture and Forestry
Government Gwacheon Complex
Jungangdong, Gwacheon-City
Gyeonggi Province
Republic of Korea
Phone: +82 2 500 1918
Fax: +82 2 503 0020
Email: kwagsi@maf.go.kr

Mr KIM Sang Kyu
Director
Fisheries Products Quality and Safety Team
Fisheries Policy Bureau
Ministry of Maritime Affairs and Fisheries
140-2 Gye-Dong
Jongno-Gu
Seoul
Republic of Korea
Phone: +82 2 3674 6921
Fax: +82 2 3674 6919
Email: kskyu@momaf.go.kr

Ms PARK Mi Jung
Researcher
National Fisheries Products Quality Inspection
Service
912-7, Joongsan-dong
Ilsan-donggu, Koyang-City
Gyeonggi-do
Republic of Korea
Phone: +82 31 976 3024
Fax: +82 31 976 6391
Email: parkmj@momaf.go.kr

Dr HAN Kyu-Jai
Principal Research Scientist
Korea Food Research Institute
#516 Baekhyeon-dong
Bundang-gu, Seongnam-si
Gyeonggi-do, 436-746
Republic of Korea
Phone: +82 31 780 9120
Fax: +82 31 780 9264
Email: hankj@kfri.re.kr

Dr HA Jaeho
Principal Research Scientist
Korea Food Research Institute
#516, Baekhyeon-dong
Bundang-gu, Seongnam-si
Gyeonggi-do, 436-746
Republic of Korea
Phone: +82 31 780 9127
Fax: +82 31 780 9264
Email: jhkfri@kfri.re.kr

Dr RHO Jeong-Hae
Senior Research Scientist
Korea Food Research Institute
#516, Baekhyeon-dong
Bundang-gu, Seongnam-si
Gyeonggi-do, 436-746
Republic of Korea
Phone: +82 31 780 9060
Fax: +82 31 780 9264
Email: drno@kfri.re.kr

ROMANIA - ROUMANIE - RUMANIA

Mrs Carmen BERGHES
Counsellor of the Minister
Ministry of Agriculture and Rural Development
Carol Street I, no. 2, secto 3
023951 Bucharest

Mrs Monica NEAGU
Director
Division for Standards, Marks and Food Quality
National Sanitary Veterinary and Food Safety
Authority
Negustori Street, no. 1B, Sector 2
023951 Bucharest
Email: neagu@ansv.ro

Mr Alexandru POPESCU
Director
Division for European Affairs and International
Relations
National Sanitary Veterinary and Food Safety
Authority
Negustori Street, no. 1B, Sector 2
023951 Bucharest
Email: popescu@ansv.ro

Ms Alina PUSCARAGIU
Deputy Permanent Representative to FAO
Embassy of Romania
Via Nicolò Tartaglia 36
00197 Rome
Phone: +39 06 8084529
Fax: +39 06 8084995
Email: amdiroma@roembit.org
fao@roembit.org

RWANDA

Dr Kimonyo ANASTASE
Acting Director-General of
Rwanda Bureau of Standards
Kigali
Phone: +00250 08532755
Email: akimonyo@gmx.de

SAUDI ARABIA - ARABIE SAOUDITE – ARABIA SAUDITA

Mr Nabil A. MOLLA
Director-General
Saudi Arabian Standards Organization - SASO
P.O. Box 3437 Riyadh 11471

Mr Yacoub A. AL-TURKI
Director General of Standards General Department
Saudi Arabian Standards Organization - SASO
P.O. Box 3437 Riyadh 11471
Email: alturky@saso.org.sa

Mr Abdulaziz Sh. SHUWAISH
Agrculture and Food Products Department
Saudi Arabian Standards Organization - SASO
P.O. Box 3437 Riyadh 11471
Email: aalshuwaish@saso.org.sa

Dr Ibrahim AL-MOHEZA
Vice Executive President for Food Affairs
Saudi Food and Drug Authority
Riyadh
Email: isarm12@sfd.gov.sa

Dr Salah A. AL-MAIMAN
Consultant on Food Affairs
Saudi Food and Drug Authority
Riyadh
Email: samaiman@sfd.gov.sa

SENEGAL - SÉNÉGAL

M Adama BA
Deuxième Conseiller
Représentant permanent adjoint auprès de la FAO
Ambassade de la République du Sénégal
Via Giulia, 66
00186 Rome
Phone: +39 06 68210342
Fax: +39 06219294
Email: ambasenequiri@tiscali.it

M Alassane WELE
Deuxième Conseiller
Représentant permanent suppléant auprès de la FAO
Ambassade de la République du Sénégal
Via Giulia, 66
00186 Rome
Phone: +39 06 68210342
Fax: +39 06 68219294
Email: ambasenequiri@tiscali.it

SERBIA - SERBIE

Mr Slobodan SIBALIC
Head of Department for Veterinary Public Health
Ministry of Agriculture, Forestry and Water
Management
Veterinary Directorate
1 Omladinskih Brigada St.
11070 New Belgrade
Phone: +381 11 2604629
Fax: +381 11 3117657
Email: s.sibalic@minpolj.sr.gov.yu
ssibalic@sbb.co.yu

SINGAPORE - SINGAPOUR - SINGAPUR

Dr Sin Bin CHUA
 Chief Executive Officer
 Agri-Food and Veterinary Authority, Singapore
 5, Maxwell Road, #04-00, Tower Block
 MND Complex
 Singapore 069110
 Phone: +65 6325 7622
 Fax: +65 6220 6068
 Email: chua_sin_bin@ava.gov.sg

Mr Sin-I CHU
 Head, Food Legislation and Factory Control Branch
 Agri-Food and Veterinary Authority
 5, Maxwell Road, #18-00
 Tower Block, MND Complex
 Singapore 069110
 Phone: +65 6325 8582
 Fax: +65 6324 4563
 Email: chu_sin-i@ava.gov.sg

SLOVAKIA - SLOVAQUIE - ESLOVAQUIA

Dr Zuzana BIROSOVA
 Director of Food Safety Department
 Ministry of Agriculture
 Dobrovcova 12
 81266 Bratislava
 Phone: +004212 59266571
 Email: zuzana.birosova@land.gov.sk

Mr Milan KOVÁČ
 Counsellor
 Permanent Representative to FAO
 Embassy of the Slovak Republic
 Via dei Colli della Farnesina, 144
 00194 Rome
 Phone: +39 06 36715206

SLOVENIA - SLOVÉNIE - ESLOVENIA

Ms Blaza NAHTIGAL
 Ministry of Agriculture, Forestry and Food
 Directorate for Food Safety
 Dunajska, 58
 1000 Ljubljana
 Phone: +00386 14789398
 Fax: +00386 1 4789055
 Email: blaza.nahtigal@gov.si

SOUTH AFRICA - AFRIQUE DU SUD - SUDÁFRICA

Ms Jennifer Morongoe RATHEBE
 Director
 Food Safety and Quality Assurance
 Department of Agriculture
 Private Bag X343
 0001 Pretoria
 South Africa
 Phone: +27 12 319 7000/7304
 Fax: +27 12 319 6764
 Email: DFSQA@nda.agric.za

Mr Andries Wilhelmus Jacobus PRETORIUS
 Director
 Food Control
 Department of Health
 Directorate: Food Control
 Private Bag X828
 0001 Pretoria
 South Africa
 Phone: +27 12 3120185
 Fax: +27 12 3123180
 Email: pretoa@health.gov.za

Mrs Francina MAKHOANE
 Deputy Director
 International Liaison and Regulatory Nutrition
 National Codex Contact Point
 Department of Health
 Private Bag X828
 0001 Pretoria
 South Africa
 Phone: +27 12 3120158
 Fax: +27 12 3123180
 Email: cacpsa@health.gov.za

Mr Michael Jeffrey YOUNG
 Senior Manager
 Food and Associated Industries SABS
 Regulatory Affairs
 South African Bureau of Standards
 14 Railway Road
 Montague Gardens, 7441
 Cape Town
 South Africa
 Phone: +27 21 526 3400
 Fax: +27 21 526 3450
 Email: youngmj@sabs.co.za

Mr Pantelis Nicholas TSELENTIS
 Legal and Regulatory Affairs Manager
 Consumer Goods Council of South Africa
 P.O. Box 41417
 2024 Graighall
 South Africa
 Phone: +27 11 789 5777
 Fax: +27 12 886 4966
 Email: nt@cgcsa.co.za

Dr Nomsa MNISI
 Deputy Director
 Veterinary Public Health
 Private Bag 138
 Pretoria 0001
 Republic of South Africa
 Phone: +2712 2197456/7688
 Fax: +273297218
 Email: nomfundom@nda.agric.za

SPAIN - ESPAGNE - ESPAÑA

Don José Luis DICENTA BALLESTER
Embajador
Representante Permanente ante la FAO
Embajada de España
(Oficina de los Representantes Permanentes Adjunto y Alternos)
Largo dei Lombardi, 21
00186 Roma
Phone: +39 06 6878762
Fax: +39 06 06 6873076
Email: repfao@maparoma.191.it

Sr Juan Carlos CALVO HUERTA
Jefe de Servicio de la CIOA
Subdirección General de Gestión de Riesgos Alimentarios
Alcalá, 56- Despacho 437
28014 Madrid
España
Phone: +91 33 800 40
Fax: +91 33 80883
Email: jcalvo@msc.es

Sra. Elisa REVILLA GARCÍA
Subdirectora General Adjunta
Subdirección General de Planificación y Control Alimentario
Ministerio de Agricultura, Pesca y Alimentación
Pº Infanta Isabel I
28071 Madrid
Phone: +34 91 3474596
Fax: +34 91 3475728
Email: erevilla@mapya.es

Sr Victorio José TERUEL MUÑOZ
Jefe de Area Gestión de Riesgos químicos
Agencia Española de Seguridad Alimentaria y Nutrición
Subdirección General de Gestión de Riesgos Alimentarios
Alcalá, 56- Despacho 498
28014 Madrid
España
Phone: +91 3380122
Email: vteruel@msc.es

Sra. M. Jesús CALCEDO BARBA
Jefe de Sección
Agencia Española de Seguridad Alimentaria y Nutrición
Subdirección General de Gestión de Riesgos Alimentarios
Alcalá, 56
28014 Madrid
Phone: +91 3380819
Email: mcalcedo@msc.es

D. Ernesto RÍOS LÓPEZ
Consejero
Representante Permanente Adjunto ante la FAO
Embajada de España
(Oficina de los Representantes Permanentes Adjunto y Alternos)
Largo dei Lombardi, 21
00186 Roma

D. Ángel OROZCO GÓMEZ
Agregado
Representante Permanente Alternos ante la FAO
Embajada de España
(Oficina de los Representantes Permanentes Adjunto y Alternos)
Largo dei Lombardi, 21
00186 Roma

Dª Icíar RUIZ CASTILLO
Técnica
Representación Permanente de España ante la FAO y el PMA
Largo dei Lombardi, 21
00186 Roma

D. Pedro BONADA
Consejero Comercial
Embajada de España
Largo dei Lombardi, 21
00186 Roma

SRI LANKA

Mrs Saranya Hasanthi Urugodawatte
DISSANAYAKE
Permanent Representative (Designate) and Chargé d'Affaires a.i.
Embassy of the Democratic Socialist Republic of Sri Lanka
Via Adige, 2
00198 Rome
Phone: +39 06 8840801/8554560
Fax: +39 06 84241670
Email: slembassy@tiscali.it

Mr Abeynayakage Wesley Wimal HEMACHANDRA
Minister Counsellor (Commercial)
Embassy of the Democratic Socialist Republic of Sri Lanka
Via Adige, 2
00198 Rome
Phone: +39 06 8554560
Fax: +39 06 84241670
Email: sltrade@virgilio.it/slembassy@tiscali.it

SUDAN - SOUDAN - SUDÁN

Dr Omer ABD ALWAHAB ABDALLA
Under-Secretary
Federal Ministry of Agriculture and Forestry
P.O. Box 285
Al Gamaá Avenue
Khartoum
Phone: +249 912398400
Fax: +249 183 782027
Email: moafus@yahoo.com

Mr Hamdi Abbas IBRAHIM
 Director
 Standards and Quality Control Unit
 Federal Ministry of Agriculture and Forestry
 P.O. Box 285
 Al Gamaá Avenue
 Khartoum
 Phone: +249 918211470
 Fax: +249 183782027
 Email: hamdi20072000@yahoo.com

Dr Bashir TAHA MOHAMED TAHA
 Under Secretary
 Ministry of Animal Resources and Fisheries
 Khartoum
 Phone: +249 183 465920
 Fax: +249 183475996

Mr Zakaria Mohamed SULIMAN SALIH
 Manager
 SSMO Portsudan Branch
 SSMO/Sudan
 Khartoum
 Fax: +249 183 774852
 Email: zak-suliman@yahoo.com

Ms Rahba SAED ABDALLA MOHAMED
 Manager
 International Relations Unit
 Sudanese Standards and Metrology Organization
 SSMO/Sudan
 Khartoum
 Phone: +249 912 207959
 Fax: +249 183 774852
 Email: rahbamohamed563@hotmail.com

Mr Ismail Ahmed ALKAMISH
 Director
 Food Control Administration
 Federal Ministry of Health
 Khartoum
 Phone: +249 12247820
 Email: Kamish2005@hotmail.com

Mr Mohamed ELTAYEB ELFAKI ELNOR
 Counsellor
 Permanent Representative to FAO
 Embassy of the Republic of the Sudan
 Via Prati della Farnesina, 57
 00194 Rome

SWEDEN - SUÈDE - SUECIA

Mrs Kerstin JANSSON
 Deputy Director
 Ministry of Agriculture
 SE-103 33 Stockholm
 Sweden
 Phone: +46 8 4051168
 Fax: +46 8 206496
 Email: kerstin.jansson@agriculture.ministry.se

Mrs Eva Rolfsdotter LÖNBERG
 Codex Coordinator
 National Food Administration
 Box 622
 SE-751 26 Uppsala
 Sweden
 Phone: +46 18175500
 Fax: +46 181 05848
 Email: codex@slv.se

Dr Stuart SLORACH
 Stubbängsvägen 9A
 SE 12553 Älvsjö
 Sweden
 Phone: +468 6469597
 Fax: +468 6469597
 Email: stuart.slorach@gmail.com

SWITZERLAND - SUISSE - SUIZA

Mrs Awilo OCHIENG PERNET
 Codex Alimentarius, International Food Safety and
 Nutrition Issues
 Division of International Affairs
 Swiss Federal Office of Public Health
 CH-3003 Bern
 Switzerland
 Phone: +41 31 3220041
 Fax: +41 31 3229574
 Email: awilo.ochieng@bag.admin.ch

Mr Patrik AEBI
 Head of Quality and Sales Promotion Section
 Principal Division for Production and International
 Affairs
 Federal Office for Agriculture
 Mattenhofstrasse 5
 CH-3003 Bern, Switzerland
 Phone: +41 31 322 2592
 Fax: +41 31 322 2634
 Email: patrik.aebi@blw.admin.ch

Dr Marquard IMFELD
 Bioresco Ltd
 Bundesstrasse 29
 CH-4054 Basel, Switzerland
 Phone: +41 61 2737706
 Fax: +41 61 2737703
 Email: marquard.imfeld@bioresco.ch

Dr Hervé NORDMANN
 Director Scientific and Regulatory Affairs
 Ajinomoto Switzerland AG
 Innere Güterstrasse 2-4
 CH-6304 Zug, Switzerland
 Phone: +4121 8003763
 Fax: +41218004087
 Email: herve.nordmann@asg.ajinomoto.com

Mr Jürg SIMON
 Lenz and Staehelin
 Bleicherweg 58
 CH-8027 Zürich, Switzerland
 Phone: +41 44 2041212
 Fax: +41 44 2041200
 Email: juerg.simon@lenzstaehelin.com

Mrs Ursula TRÜEB
 Representative of the Swiss Consumer Organizations
 Bözli 1
 CH-4312 Magden, Switzerland
 Phone: +41 61 8411256
 Fax: +41 61 8411256
 Email: ursula.trueb@tiscali.ch

THAILAND - THAÏLANDE - TAILANDIA

Mr Somchai CHARNNARONGKUL
 Deputy Secretary General
 National Bureau of Agricultural Commodity and
 Food Standards
 Ministry of Agriculture and Cooperatives
 3 Rajdamnoen Nok Avenue
 Bangkok 10200
 Thailand
 Phone: +66 2 280 3882
 Fax: +66 2 280 3899/66 2 280 3886
 Email: somchaic@acfs.go.th

Dr Nippon POPATTANACHAI
 Deputy Secretary General
 Food and Drug Administration
 Ministry of Public Health
 Tiwanond Rd
 Nonthaburi 10110
 Thailand
 Phone: +66 2 591 8443
 Fax: +66 2 590 7116/66 2 590 7197
 Email: nippon@fda.moph.go.th

Dr Chanin CHAROENPONG
 Scientific Advisor on Standard of Health Products
 Food and Drug Administration
 Ministry of Public Health
 Ministry of Public Health
 Tiwanond Rd
 Nonthaburi 10110
 Thailand
 Phone: + 66 2 591 8447
 Fax: +66 2 591 8460
 Email: chanin@fda.moph.go.th

Mrs Tritaporn KHOMAPAT
 Minister (Agriculture)
 Permanent Representative to FAO
 Office of Agricultural Affairs
 Royal Thai Embassy
 Via Cassia 929 Villino M
 00189 Rome
 Phone: +39 06 3036 3687
 Fax: +39 06 3031 2700
 Email: thagri.rome@flashnet.it

Mrs Oratai SILAPANAPAPORN
 Director
 Office of Commodity and System Standards
 National Bureau of Agricultural Commodity and
 Food Standards
 Ministry of Agriculture and Cooperatives
 3 Rajdamnoen Nok Avenue
 Bangkok 10200, Thailand
 Phone: +66 2 280 3900
 Fax: +66 2 280 3899
 Email: oratai@acfs.go.th

Mr Ronachai JUANGPHANICH
 Director, Livestock Standards Division
 Bureau of Livestock Standards and Certification
 Department of Livestock Development
 Ministry of Agriculture and Cooperatives
 Phaya Thai rd, Rajathevee
 Bangkok 10400 Thailand
 Phone: +66 2 653 4444 ext.3141
 Fax: +66 2 653 444 ext. 3143
 Email: dr_juang@yahoo.com

Mr Pisan PONGSAPITCH
 Standards Officer
 Office of Commodity and System Standards
 National Bureau of Agricultural Commodity and
 Food Standards
 Ministry of Agriculture and Cooperatives
 3 Rajdamnoen Nok Avenue
 Bangkok 10200, Thailand
 Phone: +66 2 283 1681
 Fax: +66 2 280 3899
 Email: pisanp@yahoo.com

Mrs Sairak CHAILANGGAR
 First Secretary (Agriculture)
 Deputy Permanent Representative to FAO
 Office of Agricultural Affairs
 Royal Thai Embassy
 Via Cassia 929 Villino M
 00189 Rome
 Phone: +39 06 3036 3687
 Fax: +39 06 3031 2700
 Email: sairakc@gmail.com

Mr Boonpeng SANTIWATTANATAM
 Vice-Chairman of Food Processing Industry Club
 The Federation of Thai Industries
 Queen Sirikit National Convention Center, Zone C,
 4th Fl.,
 60 New Rachadapisek Rd., Klongtoey
 Bangkok 10110, Thailand
 Phone: +66 2 229 4255 ext. 505
 Fax: +66 2 229 4937
 Email: foodgroup@off.fti.or.th
 boonpeng@cpf.co.th

Mrs Malinee SUBVANICH
 General Secretary of Thai Food Processors'
 Association
 Board of Trade Thailand
 150 Rajbopit Rd., Pranakhon District
 Bangkok 10200
 Thailand
 Phone: +66 2 261 2684-6
 Fax: +66 2 261 2996-7
 Email: thaifood@thaifood.org

TOGO

M Kokou EGUE
 Directeur des Laboratoires
 Division Nutrition et Technologie alimentaire et
 normalisation
 Institut Togolais de Recherche Agronomique
 BP 1163
 Lomé
 Phone: +228 225 2148/225 3096
 Fax: +228 225 1559
 Email: eguekokou@yahoo.fr

TUNISIA - TUNISIE - TÚNEZ

M Mohamed Chokri REJEB
 Directeur Général
 Centre Technique de l'Agro-Alimentaire
 Tunis
 Phone: +216 71940358
 Fax: +2167 1941080
 Email: ctaa@email.ati.tn
 codextunisie@email.ati.tn

M Mohamed LAOUNI
 Directeur
 Direction Générale de la Qualité, du Commerce
 Intérieur et des services et métiers
 Ministère du Commerce et de l'Artisanat
 36 Avenue Khereddine Pacha
 Tunis
 Phone: +21671842029
 Fax: +21671 800382
 Email: codextunisie@email.ati.tn

M Mabrouk NEDHIF
 Directeur
 Direction de l'hygiène du milieu et de la protection
 de l'environnement
 Ministère de la santé publique
 Rue Bab Saadoun
 Tunis
 Phone: +002167 1940081
 Fax: +002167 1941080
 Email: codextunisie@email.ati.tn

M Nabil AMMAR
 Ministre plénipotentiaire
 Représentant permanent suppléant auprès de la FAO
 Ambassade de la République tunisienne
 Via Asmara, 7
 00199 Rome

TURKEY - TURQUIE - TURQUÍA

Mrs Nergiz ÖZBAG
 Food Engineer
 Codex Division
 General Directorate of Protection and Control
 Ministry of Agriculture and Rural Affairs
 Akay Cad. No.3
 Bakanliklar
 Ankara
 Phone: +90 (312)4174176/6202
 Email: nergizo@kkgm.gov.tr

Mrs Ebru EKEMAN
 Second Secretary
 Alternate Permanent Representative to FAO
 Embassy of the Republic of Turkey
 Via Palestro, 28
 00185 Rome

UGANDA - OUGANDA

Dr Terry Epaphura KAHUMA
 Executive Director
 Uganda National Bureau of Standards
 P.O. Box 6329
 Kampala
 Phone: +256 0414 505995
 Fax: +256 0414 286123
 Email: unbs@infocom.co.ug

Mr Ben MANYINDO
 Deputy Executive Director
 Uganda National Bureau of Standards
 P.O. Box 6329 Kampala
 Uganda
 Phone: +256 0414 505995
 Fax: +256 0414 286123
 Email: unbs@infocom.co.ug
 ben.manyindo@unbs.go.ug

Ms Hope KABARISI
 Special Presidential Assistant (Science and
 Technology)
 State House
 Member of the Uganda National Codex Committee
 Uganda National Bureau of Standards
 Plot M217 Nakawa Industrial Area
 P.O. Box 6329
 Kampala
 Uganda
 Phone: +256 772 505 646
 Fax: + 256 414 343277
 Email: kabirisi@yahoo.co.uk

**UNITED ARAB EMIRATES –
ÉMIRATS ARABES UNIS –
EMIRATOS ÁRABES UNIDOS**

Dr Rashid Ahmed BIN-FAHAD
Vice-Chairman
Emirates Authority for Standardization and
Metrology
P.O. Box 2166
Abu Dhabi
Phone: +00971 2 6710999
Fax: +00971 2 6710888
Email: binfahad@gso.org.sa

Mr Khalid Mohammed SHARIF AL AWADI
Head of Food Control Section
Assistant Director of Health Department- Dubai
Municipality
Emirates Authority for Standardization and
Metrology ESMA
P.O. Box 2166
Abu Dhabi
UAE
Phone: +009714 2064200
Fax: +009714 2231905
Email: kmsharif@dm.gov.ae

Ms Amina AHMED MOHAMMED
Head of Food and Environment Lab. Section
Dubai Municipality
Emirates Authority for Standardization and
Metrology ESMA
PO Box 2166 Abu Dhabi
UAE
Phone: +009714 3027373
Fax: +009714 3358448
Email: aamohammed@dm.gov.ae

**UNITED KINGDOM - ROYAUME-UNI - REINO
UNIDO**

Mr Steve WEARNE
Head of Regulation, International and Openness
Division
Food Standards Agency
115B, Aviation House
125 Kingsway
London WC2B 6NH
United Kingdom
Phone: +44 20 7276 8338
Fax: +44 20 7276 8376
Email: steve.wearne@foodstandards.gsi.gov.uk

Mr Bill KNOCK
Head of EU and International Strategy
Food Standards Agency
Aviation House
125 Kingsway
London WC2B 6NH
United Kingdom
Phone: +44 20 7276 8183
Fax: +44 20 7276 8376
Email: bill.knock@foodstandards.gsi.gov.uk

**UNITED REPUBLIC OF TANZANIA -
RÉPUBLIQUE-UNIE DE TANZANIE -
REPÚBLICA UNIDA DE TANZANÍA**

Mr Wilfred NGIRWA
Ambassador
Permanent Representative to FAO
Embassy of the United Republic of Tanzania
Villa Tanzania
Via Cortina D'amezzo, 185
00135 Rome

Mr Linus GEDI
Food Technologist
Small Industries Development Organization
Mfaume/Fire Road
P.O. Box 2476
Dar-es-Salaam
Email: gedilinus@gmail.com

Mrs Perpetua Mary HINGI
Agricultural Attaché
Alternate Permanent Representative to FAO
Embassy of the United Republic of Tanzania
Villa Tanzania
Via Cortina D'amezzo, 185
00135 Rome
Email:
mhingi@yahoo.co.uk/phingi@embassyoftanzania.it

Mr Amon RUVURULA
Minister Counsellor
Embassy of the United Republic of Tanzania
Villa Tanzania
Via Cortina D'amezzo, 185
00135 Rome

**UNITED STATES OF AMERICA –
ÉTATS-UNIS D'AMÉRIQUE –
ESTADOS UNIDOS DE AMÉRICA**

Mr Richard RAYMOND
Under Secretary for Food Safety
Department of Agriculture
1400 Independence Avenue, SW
Room 227E, JLW Bldg
Washington, D.C. 20250
United States of America
Phone: +1-202 720 0351
Fax: +1 202 690 0820
Email: Dick.Raymond@usda.gov

Ambassador Gaddi H. VASQUEZ
Permanent Representative to FAO
U.S. Mission to the U.N. Agencies
United States Embassy
Via Vittorio Veneto, 119/A
Rome, Italy
Phone: +39 06 4674 3529
Fax: +39 06 4674 3535
Email: usunrome@state.gov

Mr Terry BANE
Branch Chief
Fruit and Vegetable Programs
Agricultural Marketing Services
South Building, Room 0709
U.S. Department of Agriculture
1400 Independence Ave, SW
Washington, DC 20250
RM 0709 Stop 0247
Phone: +202 690-0627
Fax: +202 690 1087
Email: Terry.bane@usda.gov

Mr Jack Anthony BOBO
Deputy Chief
Biotechnology Trade Division
United States Department of State
2201 C Street NW
Washington, D.C. 20520
U.S.A.
Phone: +1 202 647 1647
Fax: +1 202 647 1894
Email: boboja@state.gov

Ms Catherine CARNEVALE
Director, International Affairs Staff
Center for Food Safety and Applied Nutrition
U.S. Food and Drug Administration
5100 Paint Branch Parkway
College Park, MD 20740
United States of America
Phone: + 1 301 436 1723
Fax: +1 301 436 2618
Email: Catherine.carnevale@fda.hhs.gov

Ms Amanda EAMICH
Public Affairs Specialist
Food Safety and Inspection Service
South Building, Room 1175
U.S. Department of Agriculture
1400 Independence Ave., SW
Washington, DC 20250
Phone: +202 720 9113
Fax: +202 5704
Email: Amanda.Eamich@fsis.usda.gov

Ms Karen L. HULEBAK
Chief Scientist
Food Safety and Inspection Service
United States Department of Agriculture
1400 Independence Ave., SW
Washington, DC 20250
United States of America
Phone: +202 720 2644
Email: karen.hulebak@fsis.usda.gov

Mr Lee A. BRUDVIG
Deputy Chief of Mission
U.S. Mission to the U.N. Agencies
United States Embassy
Via Vittorio Veneto, 119/A
Rome, Italy
Phone: +39 06 4674 3513
Fax: +39 06 4674 3535
Email: BrudvigLA@state.gov

Mr Robin GRAY
Agriculture Attaché
Office of Agricultural Affairs
United States Embassy
Via Vittorio Veneto, 119/A
Rome, Italy
Phone: +39 06 4674 2362
Fax: +39 4788 7008
Email: Robin.Gray@usda.gov

Ms V. Eileen D. HILL
Lead International Economist
International Trade Administration
U.S. Department of Commerce
14th Street and Constitution Ave., N.W.
Washington, D.C. 20230
U.S.A.
Phone: +1 202 482 5276
Fax: +1 202 482 5939
Email: Eileen_hill@ita.doc.gov

Ms Mary Frances LOWE
Senior Program Advisor
Office of Pesticide Programs
U.S. Environment Protection Agency
1200 Pennsylvania Avenue N.W.
Washington, D.C. 20460
United States of America
Phone: +1 703 305 5689
Fax: +1 703 308 1850
Email: Lowe.MaryFrances@epa.gov

Mr Marcus LOWER
Acting Director, International Regulations and
Standards Division
Office of Scientific and Technical Affairs
Foreign Agricultural Service
U.S. Department of Agriculture
1400 Independence Ave. NW
Washington, DC 20250
United States of America
Phone: +1 202 720 7217
Fax: +1 202 690 0677
Email: Marcus.lower@fas.usda.gov

Ms Barbara MCNIFF
Director, Codex Program Staff
Food Safety and Inspection Service
U.S. Department of Agriculture
1400 Independence Avenue, SW
Washington, DC 20250
United States of America
Phone: +1 202 690 4719
Fax: +1 202 690 3856
Email: Barbara.mcniff@fsis.usda.gov

Mr Richard OWEN
 Director, Agricultural Affairs
 Office of the United States Trade Representative
 600 17th Street, NW
 Washington, D.C.
 Phone: +202 3959615
 Fax: +202 395 4579
 Email: Richard_owen@ustr.eop.gov

Dr F. Edward SCARBROUGH
 Manager, U.S. Codex Office
 Food Safety and Inspection Service
 U.S. Department of Agriculture
 Room 4861 South Building
 1400 Independence Avenue, SW
 Washington, DC 20250
 U.S.A.
 Phone: +1 202 205 7760
 Fax: +1 202 720 3157
 Email: ed.scarbrough@fsis.usda.gov

Mr H. Michael WEHR
 Codex Program Coordinator
 Center for Food Safety and applied Nutrition
 U.S. Food and Drug Administration
 5100 Paint Branch Parkway
 College Park, MD 20740
 United States of America
 Phone: +1 301 436 1724
 Fax: +1 301 436 2618
 Email: michael.wehr@fda.hhs.gov

Mr Kyd BRENNER
 Partner
 DTB Associates, LLP
 901 New York Avenue, N.W.
 Third Floor
 Washington, D.C. 20001
 United States of America
 Phone: +1 202 661 7098
 Fax: +1 202 661 7093
 Email: Kbrenner@dtbassociates.com

Ms Marsha ECHOLS
 Law Office of Marsha Echols
 3286 M Street, N.W.
 Washington, D.C. 20007
 United States of America
 Phone: + 1 202 625 1451
 Fax: +1 202 625 9126
 Email: mechols@earthlink.net

Mr David P. LAMBERT
 Principal
 Lambert Associates
 5105 Yuma Street, N.W.
 Washington, D.C. 20016
 U.S.A.
 Phone: +1 202 966 5056
 Fax: +1 202 966 5094
 Email: lambertdp@yahoo.com

Mr Clayton Lee HOUGH
 Senior Vice President and General Counsel
 International Dairy Foods Association
 1250 H. Street, N.W., Suite 900
 Washington, D.C. 20005
 United States of America
 Phone: +1 202 220 3516
 Fax: +1 202 737 3516
 Email: chough@idfa.org

Mr Philippe CHABOT
 Development Resources Specialist
 1450 Independence Ave
 Washington, D.C.
 20250

Ms Erika A. LUEDIG
 Office of Agricultural Affairs
 U.S. Mission to the U.N. Agencies
 Rome
 Phone: +39 06 4674 3508
 Fax: +39 06 4674 3520
 Email: Erika.Luedig@usda.gov

URUGUAY

Sra. Maria Gabriela CHIFFLET
 Consejera
 Representante Permanente Alterno ante la FAO
 Embajada de la República Oriental del Uruguay
 Via Vittorio Veneto, 183
 00187 Roma

VANUATU

Mr Timothy TUMUKON
 Department of Livestock and Quarantine
 Ministry of Agriculture, Quarantine, Forestry and
 Fisheries
 Private Mail bag 9095
 Port Vila
 Republic of Vanuatu
 Phone: +678 23519
 Fax: +678 23185
 Email: vqisvila@vanuatu.com.vu

VENEZUELA (BOLIVARIAN REPUBLIC OF) VENEZUELA (RÉPUBLIQUE BOLIVARIENNE DU) – VENEZUELA (REPÚBLICA BOLIVARIANA DE)

Ms Achirana ALTUVE
 Segunda Secretaria
 Embajada de la República Bolivariana de Venezuela
 Via Nicolò Tartaglia, 11
 Roma

VIET NAM

Mr THO Pham Van
Deputy Head
Department for Science and Technology
Ministry of Fishery
10 Nguyen Cong Hoan Street
Hanoi
Phone: +84 4 7716270
Fax: +84 4 7716702
Email: phamvantho@mofi.gov.vn

Mr QUYNH Vu Ngoc
Director
Viet Nam Codex Contact Point
Viet Nam National Codex Committee
70 Tran Hung Dao Street
Hanoi
Phone: +84 4 9426605
Fax: +84 4 8222520
Email: vnquynhcodex@tcvn.gov.vn

Mrs HA Nguyen Thi Minh
Officer
Vietnam Codex Contact Point
Vietnam National Codex Committee
70 Tran Hung Dao Street
Hanoi
Vietnam
Phone: +84 4 9 428030
Fax: +84 4 8 222520
Email: codex@tcvn.gov.vn
nguyen_thi_minh_ha@yahoo.com

Mrs TRAN Thi My Hien
Vice Director
Directorate for Standards and Quality
Quality Assurance and Testing centre 3
49 Pasteur, District 1
Hochiminh City
Viet Nam
Phone: +84 8 8294274
Fax: +84 8 9142674
Email: ttm-hien@quatest3.com.vn

Mrs DINH Thi Huong
Engineer
Sepal Food Corporation
364-366 Nguyen Van Luong
Ward 12
District 6
Ho Chi Minh City
Phone: +84 8 8730457
Fax: +84 8 9142674

Mrs CHU Hanh Phuc
Engineer
The Representation of Viet Nam Codex Contact
Point in Hochiminh City
Quality Assurance and Testing Centre 3
49 Pasteur, District 1
Hochiminh City
Viet Nam
Phone: +84 8 829 4274
Fax: +84 8 8294274
Email: ch-phuc@quatest3.com.vn

Mrs SANG Pham Thi
Deputy of Head
Food Office Under Centre for Standardization of Viet
Nam
8 Hoang Quoc Viet Street
Nghia Do- Hanoi
Phone: +84 4 7564605
Fax: +84 4 8361771
Email: tc4tcvn@yahoo.com

Ms DO THI BICH Hanh
Director
SIPHU Company Ltd
36 A 105 Chua-Boc Street
Hanoi
Viet Nam

Mr NGUYEN QUANG Tien
Vice-Director
SIPHU Company Ltd
Hanoi
Viet Nam

Mrs DINH THI HONG Van
Officer, Director Assistant
SIPHU Company Ltd
Hanoi

YEMEN - YÉMEN

Mr Abdulrahman BAMATRAF
Ambassador
Permanent Representative to FAO
Permanent Representation of the Republic of Yemen
to FAO
Via Alessandro Malladra, 10b - Int. 10
00157 Rome

Mr Fuad SAEED MOHESN NASHER
Codex Project Director
Yemen Standardization, Metrology and Quality
Control Organization
Sana'a

ZAMBIA - ZAMBIE

Mr Delphin Mwisha KINKESE
Chief Policy Analyst Food Safety and Cosmetics
Codex Focal Point - Zambia
Ministry of Health
P.O. Box 30205
Lusaka
Zambia
Phone: +260 1254067
Fax: +260 1254067/253344
Email: dminkese@yahoo.co.uk

Mr Edward NYONGOLA
First Secretary
Alternate Permanent Representative to FAO
Embassy of the Republic of Zambia
Via Ennio Quirino Visconti, 8
00193 Rome

ZIMBABWE

Mr Fredy CHINYAVANHU
Deputy Chief Government Analyst- Food Control
Ministry of Health and Child Welfare
Gvt Analyst Laboratory
P.O. Box Cy 231
Causeway
Harare
Phone: +263 4 792026/708526
Fax: +263 4 705261
Email: fchinyavanhu@healthnet.org.zw

Mr Michael NYERE
Alternate Permanent Representative to FAO
Embassy of the Republic of Zimbabwe
Via Virgilio, 8
00193 Rome

**UNITED NATIONS AND SPECIALIZED AGENCIES
NATIONS UNIES ET INSTITUTIONS SPÉCIALISÉES
NACIONES UNIDAS Y ORGANISMOS ESPECIALIZADOS**

**INTERNATIONAL TRADE CENTRE
UNCTAD/WTO
CENTRE DU COMMERCE INTERNATIONAL
CNUCED/OMC
CENTRO DE COMERCIO INTERNACIONAL
UNCTAD/OMC**

Mrs LUDOVICA GHIZZONI
Associate Expert on Standards and Quality
Management
International trade Centre UNCTAD/WTO
54 rue de Montbrillant
1202 Geneva
Switzerland
Phone: +41 22 730 0638
Fax: +41 22 730 0576
Email: ghizzoni@intracen.org

**WORLD TRADE ORGANIZATION
ORGANISATION MONDIALE DU COMMERCE
ORGANIZACIÓN MUNDIAL DEL COMERCIO**

Ms Serra AYRAL
Economic Affairs Officer
World Trade Organization
Rue de Lausanne 154
CH-1211 Geneva
Phone: +0041 227295465
Email: serra.ayral@wto.org

**INTERNATIONAL GOVERNMENTAL ORGANIZATIONS
ORGANISATIONS GOUVERNEMENTALES INTERNATIONALES
ORGANIZACIONES GUBERNAMENTALES INTERNACIONALES**

**AFRICAN UNION
UNION AFRICAINE
UNIÓN AFRICANA**

Mr Mobido TRAORE
Director
AU-IBAR Office
Westland Road, NAIROBI
P.B. 30786
Kenya
Email: mobido.traore@au-ibar.org

**ARAB ORGANIZATION FOR AGRICULTURAL
DEVELOPMENT
ORGANISATION ARABE POUR LE
DÉVELOPPEMENT AGRICOLE
ORGANIZACIÓN ÁRABE PARA EL
DESARROLLO AGRÍCOLA**

Prof. Dr Salah El-Dein ABORAYA
Director of egypt office
Arab Organization for Agriculture Development
Khartoum
Phone: +00202 3365795
Fax: +00202 336799
Email: aborayaoad@yahoo.com

**INTER-AMERICAN INSTITUTE FOR
COOPERATION IN AGRICULTURE
INSTITUT INTERAMÉRICAIN DE
COOPÉRATION POUR L'AGRICULTURE
INSTITUTO INTERAMERICANO DE
COOPERACIÓN PARA LA AGRICULTURA**

Dr Ricardo A. MOLINS
Director, Agricultural Health and Food Safety
Inter-American Institute for Cooperation on
Agriculture
P.O. Box 55 - 2200
San José, Vasquez de Coronado
San isidro 11101
Costa Rica
Phone: +506 216 0184
Fax: +506 216 0173
Email: ricardo.molins@iica.int

**INTERNATIONAL INSTITUTE OF
REFRIGERATION
INSTITUT INTERNATIONAL DU FROID
INSTITUTO INTERNACIONAL DEL FRÍO**

Mr Didier COULOMB
Director
International Institute of Refrigeration
177 boulevard Malesherbes
75017 Paris
Phone: +33 (0) 142273235
Fax: +33 (0) 1 47631798
Email: iif-iir@iifiir.org

**INTERNATIONAL ORGANIZATION OF VINE
AND WINE
ORGANISATION INTERNATIONALE DE LA
VIGNE ET DU VIN
ORGANIZACIÓN INTERNACIONAL DE LA VIÑA
Y DEL VINO**

Mr Federico CASTELLUCCI
Director-General
International Organization of Vine and Wine
18, rue d'Aguesseau
F-75008 Paris
Phone: +33 1 44 948081
Fax: +33 1 42 669063
Email: dgeneral@oiv.int

Dr Jean-Claude RUF
Coordinator Scientific and Technical Department
Head of Unit "Enology- Safety and Health - Methods
of Analysis"
International Organization of Vine and Wine
18, rue d'Aguesseau
F-75008 Paris
Phone: +33 1 44 948094
Fax: +33 1 42 669063
Email: jruf@oiv.int

Dr Ignacio SANCHEZ RECARTE
Head of Unit "Viticulture"
International Organization of Vine and Wine
18, rue d'Aguesseau
F-75008 Paris
Phone: +33 1 44 948082
Fax: +33 1 42 669063
Email: isanchez@oiv.int

**REGIONAL INTERNATIONAL ORGANIZATION
FOR PLANT PROTECTION AND ANIMAL
HEALTH
ORGANISME INTERNATIONAL RÉGIONAL
CONTRE LES MALADIES DES PLANTES ET DES
ANIMAUX
ORGANISMO INTERNACIONAL REGIONAL DE
SANIDAD AGROPECUARIA**

Dr Oscar GARCÍA SUARÉZ
Coordinador Regional de la CRIA-OIRSA
Calle Ramón Belloso
final pasaje Isolde
Colonia Escalón
San Salvador
El Salvador
Email: ogarcia@oirsa.org

**WORLD ORGANISATION FOR ANIMAL
HEALTH
ORGANISATION MONDIALE DE LA SANTÉ
ANIMALE
ORGANIZACIÓN MUNDIAL DE SANIDAD
ANIMAL**

Dr Bernard VALLAT
Director-General
World Organisation for Animal Health (OIE)
12, rue de Prony
75017 Paris
Phone: +33 (0) 1 44151888
Fax: +33 (0) 1 42670987
Email: oie@oie.int

Dr Willem DROPPERS
Chargé de Mission to the Director-General
World Organisation for Animal Health (OIE)
12, rue de Prony
75017 Paris
Phone: +33 (0) 1 44151888
Fax: +33 (0) 1 42670987
Email: w.droppers@oie.int

Dr Francesco BERLINGIERI
Deputy Head
International Trade Department
World Organisation for Animal Health (OIE)
12, rue de Prony
75017 Paris
Phone: +33 (0) 1 44151888
Fax: +33 (0) 1 42670987
Email: f.berlingieri@oie.int

**INTERNATIONAL NON-GOVERNMENTAL ORGANIZATIONS
ORGANISATIONS NON-GOUVERNEMENTALES INTERNATIONALES
ORGANIZACIONES INTERNACIONALES NO GUBERNAMENTALES**

**ASOCIACIÓN LATINOAMERICANA DE
AVICULTURA**

Dr J. Isidro MOLFESE
Secretario Ejecutivo
Observador por ALA al Codex Alimentarius
Coordinador del CISA
Asociación Latinoamericana de Avicultura
Esteban Arce 441 - 3F
C1426BSE Buenos Aires
Phone: +54 11 4774-4770
Fax: +54 9 11 4539-2595
Email: molfese@ciudad.com.ar

**ASSOCIATION OF EUROPEAN COELIAC
SOCIETIES
ASSOCIATION DES SOCIÉTÉS COELIAQUES
EUROPÉENNES
ASOCIACIÓN DE SOCIEDADES CELÍACAS
EUROPEAS**

Mrs Hertha DEUTSCH
Chair of Codex and Labelling Affairs
AOECS Association of European Coeliac Societies
Anton Baumgartner Straße 44/C5/2302
A-1230 Vienna
Austria
Phone: +43 1 66 71 887
Fax: +43 1 66 71 887
Email: hertha.deutsch@utanet.at

**BIOTECHNOLOGY INDUSTRY ORGANIZATION
ORGANISATION DE L'INDUSTRIE
BIOTECHNOLOGIQUE
ORGANIZACIÓN DE LA INDUSTRIA
BIOTECNOLÓGICA**

Dr Michael PHILLIPS
Vice President
Biotechnology Industry Organization
1201 Maryland Avenue SW Suite 900
Washington, DC 20024
Phone: +202 962 9200
Fax: +202 488 6303
Email: mphilips@bio.org

Dr Russell SCHNEIDER
Director, Regulatory Affairs
Monsanto Company
1300 Eye Street NW, Suite 450 East
Washington, DC 20005
Phone: +202 383 2866
Fax: +202 488 6303
Email: russell.p.schneider@monsanto.com

Dr Janet COLLINS
Director, Global Regulatory Affairs
The Solae Company
Solae, LLC
P.O. box 88940
St. Louis, MO 63188
United States of America
Phone: +314 982 3534
Fax: +314 982 3490
Email: jcollins@solae.com

**CONFEDERATION OF THE FOOD AND DRINK
INDUSTRIES IN EU
CONFÉDÉRATION DES INDUSTRIES AGRO-
ALIMENTAIRE DE L'UE
CONFEDERACIÓN DE INDUSTRIAS AGRO-
ALIMENTARIAS**

Ms Clara THOMPSON
Manager
Food Policy, Science and R&D
Confederation of the Food and Drink Industries of
the EU
Avenue des Arts 43
1000 Brussels
Belgium
Phone: +32 2 500 8750
Fax: +322 508 1021
Email: c.thompson@ciao.eu

**CONSUMERS INTERNATIONAL
ORGANISATION INTERNATIONALE DES
UNIONS DE CONSOMMATEURS**

Ms Clara MEYNEN
Referentin Codex Alimentarius
Federation of Germany Consumer Organization
(VZBV)
Markgafenstr 66
10969 Berlin
Phone: +49 30 25800444
Fax: +49 3025800418
Email: meynen@vzbv.de

**COUNCIL FOR RESPONSIBLE NUTRITION
CONSEJO PARA UNA NUTRICIÓN
RESPONSABLE**

Mr John HATHCOCK
Vice-President
Council for Responsible Nutrition
1828 L.St. NW, Suite 900
Washington D.C. 20036
United States of America

Mr Mark MANSOUR
 Partner, Foley & Lardner LLP
 Council for Responsible Nutrition
 1828 L street, NW
 Suite 900
 Washington, D.C. 20036
 Phone: +202 672 5585
 Fax: +202 672 5399
 Email: mmansour@foley.com

CROP LIFE INTERNATIONAL

Ms Lucyna KURTYKA
 Global Head, International Organizations
 Monsanto 6
 1300 I Street, NW, Suite 450E
 Washington, D.C.
 Phone: +1 202 383 2861
 Fax: +1 202 789 1748
 Email: lucyna.k.kurtyka@monsanto.com

EUROPEAN DAIRY ASSOCIATION ASSOCIATION LAITIÈRE EUROPÉENNE

Mr Jean-Claude GILLIS
 European Dairy Association
 Rue Montoyer 14
 1000 Brussels
 Phone: +32 2 549 5043
 Fax: +32 2 549 5049

EUROPEAN FOOD LAW ASSOCIATION ASSOCIATION EUROPÉENNE POUR LE DROIT DE L'ALIMENTATION

Mr Miguel FERNANDES DA SILVA
 Member
 European Food Law Association
 rue de la Loi 235
 1040 Brussels
 Belgium
 Phone: +32 2 230 4845
 Fax: +32 2 2308206
 Email: efla_aeda@hotmail.com

GROUPEMENT INTERNATIONAL DES SOURCES D'EAUX NATURELLES ET D'EAUX CONDITIONÉES

Mrs Patricia FOSSELARD
 EFBW Secretary-General
 Rue de l'Association 32
 1000 Brussels
 Belgium

Ms Annick MOREAU
 Head of Scientific and Regulatory Affairs Directors
 Vice Chairwoman of EFBW's Regulatory and
 Technical Committee
 Danone Beverages Worldwide
 74 503 EVIAN CEDEX

INSTITUTE OF FOOD TECHNOLOGISTS

Dr Rodney J.H. GRAY
 Chair, IFT Global Regulations and Policy Committee
 Vice President, Regulatory Affairs
 Martek Biosciences Corporation
 6480 Dobbin Road
 Columbia, MD 21045
 Phone: + 410 740 0081
 Fax: + 410 740 2985
 Email: rgray@martek.com

Dr Rosetta NEWSOME
 Director, Science and Communications
 Institute of Food Technologists
 World Headquarters
 525 West Van Buren Street
 Chicago, IL 60607-3814
 Phone: 312-782 8424
 Fax: 312-782 8348
 Email: rlnewsome@ift.org

INTERNATIONAL ALLIANCE OF DIETARY/FOOD SUPPLEMENT ASSOCIATIONS

Mr David PINEDA EREÑO
 Manager, Regulatory Affairs
 International Alliance of Dietary/Food Supplement
 Associations
 Rue de l'Association, 50
 1000-Brussels
 Belgium
 Phone: +32 2 2091155
 Fax: +32 2 223 3064
 Email: secretariat@iadsa.be

INTERNATIONAL BABY FOOD ACTION NETWORK RÉSEAU INTERNATIONAL DES GROUPES D'ACTION POUR L'ALIMENTATION INFANTILE RED INTERNACIONAL DE ACCIÓN SOBRE ALIMENTOS DE LACTANTES

Ms Elisabeth STERKEN
 Director
 Infant Feeding Action Coalition (INFACT)
 Canada
 Phone: +416 595 9819
 Fax: +416 591 9355
 Email: esterken@infactcanada.ca

INTERNATIONAL CHEWING GUM ASSOCIATION

M Jean SAVIGNY
 General Counsel
 International Chewing Gum Association (ICGA)
 c/o Keller and Heckman
 523 Av. Louise
 1050 Brussels
 Phone: +32 2 6455071
 Fax: +32 2 6455050
 Email: savigny@khlaw.be

**INTERNATIONAL COOPERATIVE ALLIANCE
ALLIANCE COOPÉRATIVE INTERNATIONALE
ALIANZA COOPERATIVA INTERNACIONAL**

Mr Kazuo ONITAKE
Head of Unit
Safety Policy Service
Japanese Consumers'Co-operative Union
Co-op Plaza, 3-29-8, Shibuya, Shibuyaku
Tokyo, japan 150-8913
Phone: +81 3 5778 8109
Fax: +81 3 5778 8002
Email: kazuo.onitake@jccu.coop

**INTERNATIONAL COUNCIL OF BEVERAGES
ASSOCIATION**

Ms Paivi JULKUNEN
Chair
ICBA Committee for Codex
International Council of Beverages Association
103-0022, 3-3-3- Nihonbashi-Muromachi Chuo-Ku
Tokyo, 103-0022
Phone: +81 3 3270 7300
Fax: +81 3 3270 7306
Email: pjulkunen@na.ko.com

Mr Soichi YAMAMOTO
Technical Adviser
International Council of Beverages Associations
Japan Soft Drink Association
3-3-3 Nihonbashi-Muromachi
Chuo-ku
Tokyo 103-0022
Japan
Phone: +81 2 3270 7300
Fax: +81 3 3270 7306
Email: Soichi_yamamoto@suntory.co.jp

Mr Shuji IWATA
Technical Advisor
Japan Soft Drink Association
3-3-3 Nihonbashi-Muromachi
Chuo-ku
Tokyo 103-0022
Japan
Phone: 81-3-3270-7300
Fax: 81-3-3270-7306
Email: shuji_iwata@suntory.co.jp

**INTERNATIONAL COUNCIL OF GROCERY
MANUFACTURERS ASSOCIATION
CONSEJO INTERNACIONAL DE
ASOCIACIONES DE FABRICANTES DE
COMESTIBLES**

Ms Peggy S. ROCHETTE
Senior Director of International Affairs
Grocery Manufacturers Association
1350 I Street NW
Washington, DC 20005
Phone: +202 639 5921
Fax: +202 639 5991
Email: prochette@fpa-food.org

**INTERNATIONAL DAIRY FEDERATION
FÉDÉRATION INTERNATIONALE DE LAITERIE
FEDERACIÓN INTERNACIONAL DE LECHERÍA**

Mr Christian ROBERT
Director-General
International Dairy Federation
Diamant Building
Boulevard Auguste Reyers, 80
1030 Brussels
Belgium
Phone: +32 2 7339888
Fax: +32 2 7330413
Email: Crobert@fil-idf.org

Mr Michael HICKEY
Irish National Committee of IDF
Derryreigh
Creggane, Charleville
Co. Cork
Ireland
Phone: +353 63 89392
Email: mfhickey@oceanfree.net

Mr Jörg SEIFERT
Technical Director
International Dairy Federation
Diamant Building
Boulevard Auguste Reyers, 80
1030 Brussels
Belgium
Phone: +32 2 7068643
Fax: +32 2 7330413
Email: JSeifert@fil-idf.org

**INTERNATIONAL FEDERATION FOR ANIMAL
HEALTH
FÉDÉRATION INTERNATIONALE POUR LA
SANTÉ ANIMALE**

Dr Peter G.H. JONES
Executive Director
IFAH, International Federation for Animal Health
Rue Defacqz,
1-1000 Brussels (Belgium)
Phone: +32 2 541-0111
Fax: +32 2 541 0119
Email: ifah@ifahsec.org

Mr Robert LIVINGSTON
Director
International Affairs and Regulatory Policy
Animal Health Institute
1325 G Street NW Suite 700
Washington, DC 2005-3104
U.S.A.
Phone: +1 202 637 2440
Fax: +202 393 1667
Email: rlivingston@ahi.org

Dr Olivier ESPEISSE
 Corporate Affairs Manager
 ELANCO Animal Health
 Research and Development
 Stoofstraat 52
 1000 Brussels (Belgium)
 Phone: +32-2 5488606
 Fax: +32 2512510
 Email: espeisse_olivier@lilly.com

Mr Dennis L. ERPELDING
 Manager
 Elanco Government Relations, Public Affairs and
 Communications
 Elanco Animal Health
 2001 West Main Street
 P.O. Box 708
 Greenfield, IN 46140
 United States of America
 Phone: +317 2762721
 Email: erpelding_dennis_1@lilly.com

**INTERNATIONAL FEDERATION OF
 AGRICULTURAL PRODUCERS
 FÉDÉRATION INTERNATIONALE DES
 PRODUCTEURS AGRICOLES
 FEDERACIÓN INTERNACIONAL DE
 PRODUCTORES AGRÍCOLAS**

Mr Nils FARNERT
 Adviser
 International Federation of Agricultural Producers
 60 rue Saint Lazare
 75009 Paris
 Phone: +33 145 260553
 Fax: +33 1 48747212
 Email: ifap@ifap.org

**INTERNATIONAL FEDERATION OF
 ENVIRONMENTAL HEALTH**

Professor Eunice TAYLOR
 Centre Director
 International Federation of Environmental Health
 (IFEH)
 International Centre for HACCP Innovation
 United Kingdom
 Phone: +01611 295 5107
 Email: e.taylor@salford.ac.uk

**INTERNATIONAL FEDERATION OF FRUIT
 JUICE PRODUCERS
 FÉDÉRATION INTERNATIONALE DES
 PRODUCTEURS DE JUS DE FRUITS
 FEDERACIÓN INTERNACIONAL DE LOS
 PRODUCTORES DE JUGOS DE FRUTAS**

Mrs Elisabetta ROMEO-VAREILLE
 Secretary-General
 International Federation of Fruit Juice Producers
 23, Boulevard des Capucines
 75002 PARIS
 France
 Phone: +331 47428280
 Fax: +331 4748281
 Email: ifu@ifu-fruitjuice.com

**INTERNATIONAL FROZEN FOODS
 ASSOCIATION
 ASSOCIATION INTERNATIONALE DES
 DENRÉES CONGELÉES
 ASOCIACIÓN INTERNACIONAL DE
 ALIMENTOS CONGELADOS**

Mr John T. ALLAN
 International Frozen Food Association (IFFA)
 2000 Corporate Ridge
 Suite 1000
 McLean, Virginia 22102
 United States of America
 Phone: +01 703 821 0770
 Fax: +01 703 821 1350
 Email: jallan@affi.com

**INTERNATIONAL GLUTAMATE TECHNICAL
 COMMITTEE
 COMITÉ INTERNATIONALE TECHNIQUE DE
 L'ACIDE GLUTAMIQUE**

Dr Tadashi HIRAKAWA
 IGTC Scientific Advisor
 International Glutamate Technical Committee
 Hatchoubori 3-9-5, Chuo-ku
 Tokyo
 104-0032 Japan
 Phone: +81 3 3667 8311
 Fax: +81 3 3667 2860
 Email: ta-hirakawa@jafa.gr.jp

Mr Miroslav SMRIGA
 Scientific Advisor
 International Glutamate Technical Committee
 153 rue de Courcelles
 75017 Paris
 France

**INTERNATIONAL LACTATION CONSULTANT
 ASSOCIATION
 ASSOCIATION INTERNATIONALE DES
 CONSULTANT(E)S EN LACTATION**

Mrs Maryse ARENDT
 Director
 Initiativ Liewensufank
 20 rue de Contern
 L-5955 Itzig
 Phone: +00352 360598
 Fax: +00352 366134
 Email: secretariat@liewensufank.lu

**INTERNATIONAL ORGANIZATION FOR
STANDARDIZATION
ORGANISATION INTERNATIONALE DE
NORMALISATION
ORGANIZACIÓN INTERNACIONAL DE
NORMALIZACIÓN**

Mrs Marie-Noëlle BOURQUIN
Chief Technical Programme Manager
International Organization for Standardization
1, ch. de la Voie-Creuse
Case postale 56
CH-1211 Geneva 20
Switzerland
Phone: +41 22 749 7293
Fax: +41 22 7497349
Email: bourquin@iso.org

**INTERNATIONAL ORGANIZATION OF THE
FLAVOR INDUSTRY
ORGANISATION INTERNATIONALE DE
L'INDUSTRIE DES PRODUITS AROMATIQUES
ORGANIZACIÓN INTERNACIONAL DE LA
INDUSTRIA AROMÁTICA**

Dr Thierry CACHET
Scientific Director
International Organization of the Flavor Industry
Avenue des arts 6
1210 Brussels
Belgium
Phone: +32 2 2142050
Fax: +32 2 2142069
Email: secretariat@iofiorg.org

**INTERNATIONAL SPECIAL DIETARY FOODS
INDUSTRIES
FÉDÉRATION INTERNATIONALE DES
INDUSTRIES DES ALIMENTS DIÉTÉTIQUES**

Ms Andrée BRONNER
Secretary-General
ISDI
194 rue de Rivoli
F-75001 Paris
Phone: +33 1 53 458787
Fax: +331 53 458780
Email: andree.bronner@isdifederation.org

Ms Amandine DEVERGIES
Scientific and Regulatory Affairs
ISDI
194 rue de Rivoli
F-75001 Paris
Phone: +33 1 53 458787
Fax: +33 1 53 458780
Email: amandine.devergies@isdifederation.oeg

Ms Denise MALONE
Director Regulatory Affairs
International Special Dietary Foods Industries
194 rue de Rivoli
75001 Paris
Phone: +33153 3458787
Fax: +33153 458780
Email: andree.bronner@isdifederation.org

Mr Peter VAN DAEL
ISDI
194 rue de Rivoli
F-75001 Paris
Phone: +33 1 53 458787
Fax: +33 1 53 458780
Email: andree.bronner@isdifederation.org

NATIONAL HEALTH FEDERATION

Mr Scott TIPS
President and General Counsel
National Health Federation
P.O. Box 688
Monrovia, California
91017 United States of America
Phone: +1 626 3572181
Fax: +1 626 303 0642
Email: sct@thenhf.com

Mr Sepp HASSLBERGER
National Health Federation
P.O. Box 688
Monrovia, California
91017 United States of America
Phone: +1 626 357 2181
Fax: +1 626 303 0642

**WORLD SUGAR RESEARCH ORGANIZATION
ORGANISATION MONDIALE DE RECHERCHE
SUR LE SUCRE**

Dr Mary HARRINGTON
Assistant to the Director-General
World Sugar Research Organization
P.O. Box 50134
London SW1 V 3XR
United Kingdom
Phone: +44 (0) 20 7821 6800 Ext 232
Fax: +44 (0) 20 7834 4137
Email: mharrington@wsro.org

**WORLD VETERINARY ASSOCIATION
ASSOCIATION MONDIALE VÉTÉRINAIRE
ASOCIACIÓN MUNDIAL VETERINARIA**

Prof. Dr Leon RUSSELL
World Veterinary Association
Emdrupvej 28A
DK-2100 Copenhagen O
Denmark
Phone: +45 38 710156
Fax: +45 38 710322
Email: LRussell@cvm.tamu.edu
lhrdvm@verizon.net

**SECRETARIAT
SECRETARIAT
SECRETARÍA**

Dr Kazuaki MIYAGISHIMA
Secretary, Codex Alimentarius Commission
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 54390
Fax: +39 06 570 54593
Email: kazuaki.miyagishima@fao.org

Ms Selma H. DOYRAN
Senior Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 55826
Fax: +39 06 570 54593
Email: selma.doyran@fao.org

Mr Tom HEILANDT
Senior Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 54384
Fax: +39 06 570 54593
Email: tom.heilandt@fao.org

Ms Noriko ISEKI
Senior Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 53195
Fax: +39 06 570 54593
Email: noriko.iseki@fao.org

Ms Annamaria BRUNO
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 56254
Fax: +39 06 570 54593
Email: annamaria.bruno@fao.org

Dr Jeronimas MASKELIUNAS
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 53967
Fax: +39 06 570 54593
Email: jeronimas.maskeliunas@fao.org

Ms Gracia BRISCO
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 52700
Fax: +39 06 570 54593
Email: gracia.brisco@fao.org

Mrs Verna CAROLISSEN
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 55629
Fax: +39 06 570 54593
Email: verna.carolissen@fao.org

Mr Masashi KUSUKAWA
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 54796
Fax: +39 06 570 54593
Email: masashi.kusukawa@fao.org

Mr Ym Shik LEE
Food Standards Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 55854
Fax: +39 06 570 54593
Email: ymshik.lee@fao.org

Mr Virgile PACE
Executive Support Officer
Joint FAO/WHO Food Standards Programme
Viale delle Terme di Caracalla
00153 Rome
Italy
Fax: +39.06.570.54593
Email: virgile.pace@fao.org

**LEGAL COUNSEL
CONSEILLER JURIDIQUE
ASESOR JURÍDICO**

Mr Antonio TAVARES
Legal Counsel
Legal Office
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 55132
Fax: +39 06 570 54408
Email: antonio.tavares@fao.org

**FAO PERSONNEL
PERSONNEL DE LA FAO
PERSONAL DE LA FAO**

Mr José M. SUMPSSI
Assistant Director-General
Agriculture and Consumer Protection Department
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 53364
Email: jose.sumpssi@fao.org

Mr Ezzeddine BOUTRIF
Director
Nutrition and Consumer Protection Division
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 56156
Fax: +39 06 570 54593
Email: ezzeddine.boutrif@fao.org

Dr Maria de Lourdes COSTARRICA
Senior Officer
Food Quality Liaison Group
Nutrition and Consumer Protection Division
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 56060
Fax: +39 06 570 5459
Email: lourdes.costarrica@fao.org

Dr Maya PINEIRO
Senior Officer
Food Control and Consumer Protection Group
Nutrition and Consumer Protection Division
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 53308
Fax: +39 06 570 54593
Email: maya.pineiro@fao.org

Mr Julius JACKSON
Project Officer
Food Quality and Standards Service
Nutrition and Consumer Protection Division
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 5705 3653
Fax: +39 06 5705 4593
Email: Julius.Jackson@fao.org

Dr Cheikh NDIAYE
Senior Food and Nutrition Officer
FAO Regional Office for Africa
P.O. Box 1628 Accra
Ghana
Phone: +00233 21 7010930 ext. 3154 or 3406
Fax: +00233 21 7010943
Email: cheikh.ndiaye@fao.org

Mr Georges CODJIA
Food and Nutrition Officer
FAO Sub-Regional Office for Southern and Eastern
Africa
P.O. Box 3730 Harare
Zimbabwe
Email: georges.codjia@fao.org

Mrs Carmen DARDANO
Food and Nutrition Officer
Subregional Office for the Caribbean - SLAC
P.O. Box 631C
Bridgetown
Barbados
Phone: +1246 4292002
Fax: +1246 4276075
Email: carmen.dardano@fao.org

Mr B.K. NANDI
Senior Food and Nutrition Officer
Regional Office for Asia and the Pacific
39 Phra Atit Road
Bangkok
Thailand
Phone: +662 6974143
Fax: +662 6974445
Email: biplab.nandi@fao.org

Ms Annika WENNERBERG
Senior Officer and JECFA Secretary
Food Quality and Standards Service
Nutrition and Consumer Protection Division
FAO
Viale delle Terme di Caracalla
00153 Roma
Italy
Phone: 39.06.57053283
Fax: 39.06.57054593
Email: annika.wennberg@fao.org

Mr Dirk SCHULZ
Food and Nutrition Officer
Subregional Office for the Pacific Islands
Private Mail Bag
Apia
Samoa
Phone: +685 22127
Fax: +685 22126
Email: dirk.schulz@fao.org

Ms Fatima HACHEM
Food and Nutrition Officer
Regional Office for the Near East
P.O. Box 2223
Dokki
Cairo
Phone: +202 3316144
Email: fatima.hachem@fao.org

Ms Sarah CAHILL
Nutrition Officer (Food Microbiology)
Food Quality and Standards Service
Nutrition and Consumer Protection Division
FAO
Viale delle Terme di Caracalla
00153 Roma
Italy
Phone: +39 06 57053614
Fax: +39 06 57054593
Email: sarah.cahill@fao.org

Ms Mary KENNY
Nutrition Officer
Food Quality and Standards Service
Nutrition and Consumer Protection Division
FAO
Viale delle Terme di Caracalla
00153 Roma
Italy
Phone: 39.06.57053653
Fax: 39.06.57054593
Email: mary.kenny@fao.org

Ms Renata CLARKE
Nutrition Officer
Food Quality and Standards Service
Nutrition and Consumer Protection Division
FAO
Viale delle Terme di Caracalla
00153 Roma
Italy
Phone: +39 06 57053653
Fax: +39 06 57054593
Email: mary.kenny@fao.org

Ms Masami TAKEUCHI
Food Quality and Standards Service
Nutrition and Consumer Protection Division
FAO
Viale delle Terme di Caracalla
00153 Roma
Italy
Phone: 39.06.57053076
Fax: 39.06.57054593
Email: masami.takeuchi@fao.org

Dr Grimur T. VALDIMARSSON
Director
Fish Products and Industry Division
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 570 56510
Email: grimur.valdimarsson@fao.org

Ms Daniela BATTAGLIA
Animal Production Officer (Feed Utilization and Safety)
Animal Production Service (AGAP)
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 5705 6773
Email: daniela.battaglia@fao.org

Mr Tony BENNETT
Dairy and Meat Officer (Institutional Development)
Animal production Service (AGAP)
Food and Agriculture Organization (FAO)
Viale delle Terme di Caracalla
00153 Rome
Italy
Phone: +39 06 5705 4701
Email: Anthony.bennett@fao.org

**WHO PERSONNEL
PERSONNEL DE L'OMS
PERSONAL DE LA OMS**

Dr Jørgen SCHLUNDT
Director
Department of Food Safety, Zoonoses and Foodborne
Diseases
World Health Organization (WHO)
20 Avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 3445
Fax: +41 22 791 4807
Email: schlundtj@who.int

Dr Peter BEN EMBAREK
Scientist
Department of Food Safety, Zoonoses and Foodborne
Diseases
World Health Organization (WHO)
20 Avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 4204
Fax: +41 22 791 4807
Email: benembarek@who.int

Mrs Françoise FONTANNAZ
Technical Assistant
Department of Food Safety, Zoonoses and Foodborne
Diseases (FOS)
World Health Organization (WHO)
20 avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 3697
Fax: +41 22 7914807
Email: fontannazf@who.int

Dr Gyanendra GONGAL
Regional Focal Point for Foodborne Disease
Surveillance and Zoonoses
WHO Regional Office for South-East Asia
Indraprastha Estate
New Delhi 110002
India
Email: gongalg@searo.who.int

Dr Hilde KRUSE
Regional Adviser for Food Safety
WHO Regional Office for Europe
c/o European Centre for Environment and Health
Via Francesco Crispi 10
00187 Rome
Italy
Email: foodsafety@ecr.euro.who.int

Dr Gerald MOY
Scientist
Department of Food Safety, Zoonoses and Foodborne
Diseases
World Health Organization (WHO)
20 Avenue Appia
CH-1211 Geneva 27Switzerland
Phone: +41 22 791 3698
Fax: +41 22 791 4807
Email: moyg@who.int

Ms Catherine MULHOLLAND
Codex Trust Fund Administrator
Department of Food Safety, Zoonoses and Foodborne
Diseases (FOS)
World Health Organization (WHO)
20 Avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 3080
Fax: +41 22 791 4807
Email: muholandc@who.int

Ms Jenny MURCOTT
Secretary
Department of Food Safety, Zoonoses and Foodborne
Diseases
World Health Organization (WHO)
20 Avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 3557
Fax: +41 22 791 4807
Email: murcottj@who.int

Dr Chizuru NISHIDA
Scientist
Department of Nutrition for Health and Development
World Health Organization (WHO)
20 avenue Appia
CH-1211 Geneva 27, Switzerland
Phone: +41 22 791 3317
Fax: +41 22 791 4156
Email: nishidac@who.int

Dr Angelika TRITSCHER
Scientist
Chemical Safety
Department of Protection of the Human Environment
World Health Organization (WHO)
20, Av. Appia
1211 Geneva 27, Switzerland
Phone: +41 22 791 3569
Fax: +41 22 791 4848
Email: tritschera@who.int

WHO/DFID CONSULTANT

Mr Robert John CONNOR
DFID Consultant
98 High Road
Lancaster LA 26PU
United Kingdom

APPENDIX II

AMENDMENTS TO THE RULES OF PROCEDURE OF THE CODEX ALIMENTARIUS
COMMISSION**Rule IV Coordinators**

- 1.-2. [no change]
3. The functions of the Coordinators shall be:
- (i) **to appoint the Chairperson of the Coordinating Committee where such committee has been set up under Rule XI.1(b)(ii) for the region or group of countries concerned.**
 - (ii) to assist and coordinate the work of the Codex Committees set up under Rule XI.1(b)(i) in their region or group of countries in the preparation of draft standards, guidelines and other recommendations for submission to the Commission
 - (iii) to assist the Executive Committee and the Commission, as required, by advising them of the views of countries and recognized regional intergovernmental and non-government organizations in their respective regions on matters under discussion or of interest;

Rule IV (paragraph 3 (i) renumbered 3 (ii) as above)

[FRENCH ONLY]

aider aux travaux des comités du Codex créés **pour leur région ou groupe de pays** en vertu de l'Article XI.1b)i) et les coordonner, ~~dans leur région ou groupe de pays~~ en ce qui concerne la préparation de projets de normes, de lignes directrices et autres recommandations à soumettre à la Commission;

Rule V Executive Committee

1. The Executive Committee shall consist of the Chairperson and the Vice-Chairpersons of the Commission, and the Coordinators appointed on the basis of Rule IV together with seven further Members elected by the Commission at regular sessions from among the Members of the Commission, one each coming from the following geographic locations: Africa, Asia, Europe, Latin America and the Caribbean, Near East, North America, South-West Pacific. Not more than one delegate from any one country shall be a member of the Executive Committee. Members elected on a geographic basis shall hold office from the end of the session of the Commission at which they were elected until the end of the second succeeding regular session and shall be eligible for re-election if they have not served for more than two years in their current term, but after having served two consecutive terms shall be ineligible to hold such office for the next succeeding term. **Members elected on a geographic basis are expected to act within the Executive Committee in the interest of the Commission as a whole.**
- 2.-7. [no change]

Rule XI Subsidiary Bodies

- 1.-9. [no change]
10. The Members who shall be responsible for appointing Chairpersons of subsidiary bodies established under Rule XI.1(b)(i) ~~and Rule XI.1(b)(ii)~~ shall be designated at each session by the Commission and shall be eligible for re-designation. All other officers of subsidiary bodies shall be elected by the body concerned and shall be eligible for re-election.
11. [no change]

(Secretariat to take care of possible consequential changes)

APPENDIX III

**AMENDMENTS TO THE GUIDE TO THE PROCEDURE FOR THE REVISION AND
AMENDMENT OF CODEX STANDARDS AND ARRANGEMENTS FOR THE AMENDMENTS
OF CODEX STANDARDS ELABORATED BY CODEX COMMITTEES WHICH HAVE BEEN
ADJOURNED *SINE DIE***

**GUIDE TO THE PROCEDURE FOR THE AMENDMENT REVISION AND REVISION AMENDMENT OF CODEX
STANDARDS AND RELATED TEXTS**

13. The procedure for amending or revising a Codex standard is laid down in paragraph 8 of the Introduction to the Procedure for the Elaboration of Codex Standards and Related Texts. This Guide provides more detailed guidance on the existing procedure for the amendment and revision of Codex standards and related text.

24. When the Commission has decided to amend or revise a standard, the unrevised standard will remain the applicable Codex standard until the amendment to the standard or the revised standard has been adopted by the Commission.

3. For the purpose of this Guide:

Amendment means any addition, change or deletion of text or numerical values in a Codex standard or related text, may be editorial or substantive, and concerns one or a limited number of articles in the Codex text. In particular, amendments of an editorial nature may include but are not limited to:

- correction of an error;
- insertion of an explanatory footnote; and
- updating of references consequential to the adoption, amendment or revision of Codex standards and other texts of general applicability, including the provisions in the Procedural Manual.

Finalization or updating of methods of analysis and sampling as well as alignment of provisions, for consistency, to those in similar standards or related texts adopted by the Commission may be handled by the Commission in the same manner as amendments of an editorial nature, as far as the procedure described in this Guide is concerned.

Revision means any changes to a Codex standard or related text other than those covered under “amendment” as defined above.

The Commission has the final authority to determine whether a proposal made constitutes an amendment or a revision, and whether an amendment proposed is of an editorial or substantive nature.

~~42. Proposals for the amendment or revision of Codex standards and related texts should be submitted to the Commission by the subsidiary body concerned, by the Secretariat, or a member of the Commission where the subsidiary body concerned is not in existence or has been adjourned *sine die*. In the latter case, proposals should be received by the Commission’s Secretariat in good time (not less than three months) before the session of the Commission at which they are to be considered. The proposer of an amendment should indicate the reasons for the proposed amendment and should also state whether the proposed amendment had been previously submitted to and considered by the Codex committee concerned and/or the Commission. If the proposed amendment has already been considered by the Codex committee and/or Commission, the outcome of the consideration of the proposed amendment should be stated. The proposal should be accompanied by a project document (see Part 2 of the Elaboration Procedures) unless the Executive Committee or the Commission decides otherwise. However, if the amendment proposed is of an editorial nature, the preparation of a project document is not required.~~

~~53. Taking into account such information regarding the proposed amendment, as may be supplied in accordance with paragraph 1 above, and the outcome of the on-going critical review conducted by the~~

Executive Committee, the Commission ~~will~~ decides whether the amendment or revision of a standard is necessary. If the Commission decides in the affirmative, one of the following courses of action will be taken:

(i) In the case of an amendment of an editorial nature, it will be open to the Commission to adopt the amendment at Step 8 of the Uniform Procedure (see Part 3 of the Elaboration Procedures).

~~(ii) If the proposer of the amendment is a Codex committee, it would be open to the Commission to decide that the proposed amendment be circulated to governments for comments prior to further consideration by the sponsoring Codex Committee. In the case of an amendment proposed and agreed upon by a subsidiary body Codex Committee, it will also be open to the Commission to adopt the amendment at Step 5 of the Uniform Procedure (see Part 3 of the Elaboration Procedures) or Step 8 as appropriate, where in its opinion the amendment is either of an editorial nature or of a substantive nature but consequential to provisions in similar standards adopted by it at Step 8.~~

(iii) In other cases, the Commission will approve the proposal as new work and the approved new work and the proposer of the amendment is other than a Codex committee, the proposed amendment will be referred for consideration to the appropriate subsidiary body Codex committee, if such body committee is still in existence. If such body committee is not in existence, the Commission will determine how best to deal with the new work proposed amendment.

[Paragraphs 1 and 2 of the "Arrangements" are removed.]

~~65. In the case w~~Where Codex subsidiary bodies have been abolished or dissolved, or Codex committees have been adjourned sine die, the Secretariat keeps under review all Codex standards and related texts elaborated by these bodies originating from Codex Committees adjourned sine die and to determine the need for any amendments, in particular those arising from decisions of the Commission, in particular amendments of the type mentioned in para. 1(a), (b), (c), (d) and those of (e) if of an editorial nature. If a need to amend the standard appears appropriate. If the need for amendments of an editorial nature is identified then the Secretariat should prepare proposed amendments a text for consideration and adoption by in the Commission. If the need for amendments of the type in para (f) and those of (e) of a substantive nature is identified, the Secretariat, in cooperation with the national secretariat of the adjourned Committee if applicable, and, if possible, the Chairperson of that Committee, should agree on the need for such an amendment and prepare a working paper containing the wording of a proposed amendment and the reasons for proposing such amendments and the wording of such amendments as appropriate, and request comments from members of the Commission Member Governments: (a) on the need to proceed with such an amendment and (b) on the proposed amendment itself. If the majority of the replies received from members of the Commission Member Governments is affirmative on both the need to amend the standard and the suitability of the proposed wording for the amendment or an alternative proposed wording, the proposal should be submitted to the Commission with a request to approve the amendment of the standard concerned for consideration and adoption. In cases where replies do not appear to offer an uncontroversial solution then the Commission should be informed accordingly and it would be for the Commission to determine how best to proceed.

AMENDMENTS TO THE GENERAL PRINCIPLES OF THE CODEX ALIMENTARIUS

Purpose of the Codex Alimentarius

1. The Codex Alimentarius is a collection of internationally adopted food standards and related texts¹ presented in a uniform manner. These food standards and related texts aim at protecting consumers' health and ensuring fair practices in the food trade. ~~The Codex Alimentarius also includes provisions of an advisory nature in the form of codes of practice, guidelines and other recommended measures intended to assist in achieving the purposes of the Codex Alimentarius.~~ The publication of the Codex Alimentarius is intended to guide and promote the elaboration and establishment of definitions and requirements for foods to assist in their harmonization and in doing so to facilitate international trade.

Scope of the Codex Alimentarius

2. The Codex Alimentarius includes standards for all the principal foods, whether processed, semi-processed or raw, for distribution to the consumer. Materials for further processing into foods should be included to the extent necessary to achieve the purposes of the Codex Alimentarius as defined. The Codex Alimentarius includes provisions in respect of food hygiene, food additives, ~~pesticide~~ residues of pesticides and veterinary drugs, contaminants, labelling and presentation, methods of analysis and sampling, and import and export inspection and certification. ~~It also includes provisions of an advisory nature in the form of codes of practice, guidelines and other recommended measures.~~

Nature of Codex Standards

3. Codex standards and related texts are not a substitute for, or alternative to national legislation. Every country's laws and administrative procedures contain provisions with which it is essential to comply.

43. Codex standards and related texts contain requirements for food aimed at ensuring for the consumer a safe sound, wholesome food product free from adulteration, correctly labelled and presented. A Codex standard for any food or foods should be drawn up in accordance with the Format for Codex Commodity Standards and contain, as appropriate, the sections ~~criteria~~ listed therein.

Revision of Codex Standards

54. The Codex Alimentarius Commission and its subsidiary bodies are committed to revision as necessary of Codex standards and related texts to ensure that they are consistent with and reflect current scientific knowledge and other relevant information. When required, a standard or related text shall be revised or removed in accordance with the Procedures for the Elaboration of Codex Standards and Related Texts ~~using the same procedures as followed for the elaboration of a new standard~~. Each member of the Codex Alimentarius Commission is responsible for identifying, and presenting to the appropriate committee, any new scientific and other relevant information which may warrant revision of any existing Codex standards or related texts.

¹ These include codes of practice, guidelines and other recommendations.

**AMENDMENTS TO THE PRINCIPLES CONCERNING THE PARTICIPATION OF
INTERNATIONAL NON-GOVERNMENTAL ORGANIZATIONS IN THE WORK OF THE
CODEX ALIMENTARIUS COMMISSION**

1.-5. [no change]

6. Review of "Observer Status"

The Directors-General may terminate observer status if an Organization no longer meets the criteria in sections 3 and 4 above ~~that applied at the time it was granted observer status~~, or for reasons of exceptional nature, in accordance with the procedures set out in this section. [...]

DEFINITIONS FOR THE PURPOSE OF THE CODEX ALIMENTARIUS

[for inclusion in Section I]

Codex maximum level for a contaminant in a food or feed commodity is the maximum concentration of that substance recommended by the Codex Alimentarius Commission to be legally permitted in that commodity.

Good Manufacturing Practice in the Use of Food Additives means that:

- the quantity of the additive added to food does not exceed the amount reasonably required to accomplish its intended physical nutritional or other technical effect in food;
- the quantity of the additive that becomes a component of food as a result of its use in the manufacturing, processing or packaging of a food and which is not intended to accomplish any physical, or other technological effect in the food itself, is reduced to the extent reasonably possible;
- the additive is of appropriate food grade quality and is prepared and handled in the same way as a food ingredient. Food grade quality is achieved by compliance with the specifications as a whole and not merely with individual criteria in terms of safety.

PROCEDURES FOR CONSIDERATION OF THE ENTRY AND REVIEW OF FOOD ADDITIVE PROVISIONS IN THE GENERAL STANDARD FOR FOOD ADDITIVES

[for inclusion in Section II]

SCOPE

The Codex General Standard for Food Additives is intended to include food additive provisions for standardised and non-standardised foods in the Codex Alimentarius.

The following text describes the data and information that should be submitted to the Codex Committee on Food Additives when requesting the Committee to initiate work to add or revise food additive provisions in the Codex General Standard for Food Additives. The decisions required to establish acceptance or rejection of new proposals are also elaborated.

Provisions for the use of processing aids (e.g., most enzyme preparations, clarifying and filtering aids, extraction solvents) are not included in the General Standard for Food Additives.

INITIATION OF WORK

Revision

The food additive provisions of the General Standard for Food Additives may be revised by the Committee on Food Additives after requests submitted by Codex Committees, Codex members, or the Codex Alimentarius Commission. Information to support amendment of the General Standard for Food Additives shall be provided by the proposing body. Supporting information provided to the Committee on Food Additives should include, as appropriate:

- Specifications for the food additive;
- A summary of the Joint FAO/WHO Expert Committee on Food Additives (JECFA) safety evaluation of the food additive;
- The food categories or sub-categories in which the additive is intended to be used;
- An indication of the technological need / justification for the additive, referencing one or more of the General Principles for the Use of Food Additives of the GSFA (Section 3);
- Maximum use levels for the food additive in the specified food categories:
 - For additives with a numerical Acceptable Daily Intake (ADI), a numerical maximum use level for each specified use although for certain cases, a level of GMP may be appropriate;
 - For additives with an ADI Not Specified or Not Limited, a recommendation to list the additive in Table 3 accompanied by additional proposals for inclusion in Tables 1 and 2 for use in the food categories listed in the Annex to Table 3, as appropriate;
 - For additives with an “acceptable” ADI, either a numerical maximum use level for the acceptable level of treatment of a food or a level of GMP, consistent with the JECFA evaluation.
- A justification of the maximum use levels from a technological point-of-view; and an indication, by means of the procedure indicated in Annex A of the General Standard for Food Additives or an exposure assessment, that this level meets the safety requirements enumerated in Section 3.1 of the General Standard for Food Additives.
- A reasoned statement that consumers will not be misled by the use of the additive.

The Committee on Food Additives shall consider all amendments to the General Standard for Food Additives proposed by Codex Committees, Codex members, or the Codex Alimentarius Commission.

Review

The food additive provisions for the General Standard for Food Additives shall be reviewed by the Committee on Food Additives on a regular basis and revised as necessary in light of revisions of the risk assessment by JECFA or of changing technological need and justification for use.

- If JECFA changes an ADI to a Temporary ADI, the food additive provisions of the General Standard for Food Additives may remain unchanged until the ADI has been withdrawn or the full status has been restored by JECFA.
- If JECFA withdraws an ADI the food additive provisions of the General Standard for Food Additives shall be amended by removing all provision for the use of the additive.

The following additional guidance is provided regarding the information to be submitted:

- Identity of the food additive
 - Food additives shall have been evaluated by JECFA and either assigned a full numerical or non-numerical (“not specified” or “not limited”) ADI, or deemed to be acceptable for a particular use.
 - Food additives shall have been assigned an International Numbering System number.
- Functional effect of the food additive
 - The functional class list used in *Class Names and the International Numbering System* (CAC/GL 36-1989) should be used.
- Proposed use of the food additive
 - The appropriate food categories from the food category system (Annex B of the General Standard for Food Additives) and maximum use levels should be specified.
 - With regard to the acceptable maximum use level:
 - A numerical use level should be provided for a food additive assigned a numerical ADI. However, in some cases, reporting the use level as good manufacturing practice (“GMP”) may be appropriate.
 - For a food additive assigned a non-numerical (“not specified” or “not limited”) ADI that is listed in Table 3 of the General Standard for Food Additives, a numerical or good manufacturing practice (“GMP”) use level should be provided for any request to list the additive in a food category in the Annex to Table 3.
 - For some food additives, the ADI has been reported on a specific basis (e.g., “as phosphorus” for phosphates; “as benzoic acid” for benzoates). For consistency, the maximum use level for these additives should be reported on the same basis as the ADI.
- Justification for the use and technological need of the food additive
 - Supporting information based on the criteria in Section 3.2 of the Preamble of the General Standard for Food Additives should be included.
- Safe use of the food additive
 - An intake assessment of the proposed use of the food additive, in accordance with Section 3.1 of the Preamble of the General Standard for Food Additives, should be included as appropriate.
- Justification that the use does not mislead the consumer
 - A reasoned statement that consumers will not be misled by the use of the additive should be provided.

DOES THE FOOD ADDITIVE USE MEET THE CRITERIA OF SECTION 3.2 OF THE PREAMBLE OF THE GENERAL STANDARD FOR FOOD ADDITIVES?

Section 3.2 of the Preamble of the General Standard for Food Additives establishes the criteria for justifying the use of a food additive. Adherence to these criteria is necessary for the inclusion of the food additive in the General Standard for Food Additives. If the use of the additive does not meet these criteria, it is not considered further and the work is discontinued. If the information provided to justify the use of the additive is inadequate for the Codex Committee on Food Additives to reach a decision, further information on the use and technological justification and need for the food additive will be requested for consideration at the Committee's next session. If this information is not provided by the next session, work on the provision is discontinued.

IS THE FOOD ADDITIVE USED IN STANDARDIZED FOOD?

The Codex Committee on Food Additives, asks the relevant Codex commodity committee to consider the functional classes of additives, additives, and their technological justification for the commodity and to refer back this information by the next available session.. In light of this information, the Codex Committee on Food Additives recommends appropriate conditions of use based on proposals of the commodity committee.

In certain cases, however, it may be appropriate for the Codex commodity committee to develop a list of food additives with associated functional classes and acceptable maximum use levels that would be forwarded to the Codex Committee on Food Additives for endorsement and, ultimately, incorporation into the General Standard for Food Additives. The development of such food additive lists should be consistent with the principles used in the development of the General Standard for Food Additives. However, the development of food additive lists in commodity standards should be restricted as much as possible. For example, an additive may be listed in a commodity standard if it is needed to achieve a technical effect that is not achievable by the use of other additives of the same functional class. Additives may also be listed in a commodity standard if there is a need, based on a safety assessment, to limit the use of the additive. Justification for such exceptions should be provided by the Codex commodity committees to the Codex Committee on Food Additives for consideration.

If the Codex commodity committee has been adjourned, the Codex Committee on Food Additives may revise the food additive provisions in commodity standards under the purview of the adjourned committee, as necessary.

The Codex Committee on Food Additives would consider any proposed revision in light of the principles of technological justification for the use of additives as indicated in Section 3.2 of the Preamble of the General Standard for Food Additives. These revisions, once adopted by the Commission, would be incorporated into the General Standard for Food Additives.

HAS A NON-NUMERICAL (“NOT SPECIFIED” OR “NOT LIMITED”) ADI BEEN ASSIGNED?

Yes - Non-Numerical (“Not Specified” or “Not Limited”) ADI:

Food additives assigned a non-numerical ADI are proposed for inclusion in Table 3 of the General Standard for Food Additives. Requests for the use of these additives in the food categories listed in the Annex to Table 3 are made by proposing provisions for inclusion in Tables 1 and 2 of the General Standard for Food Additives. These proposals are considered by the Codex Committee on Food Additives according to the criteria described under “**Consideration of Conditions of Use in the Specific Food Categories**”, below.

No - Numerical ADI or Acceptable for Limited Use:

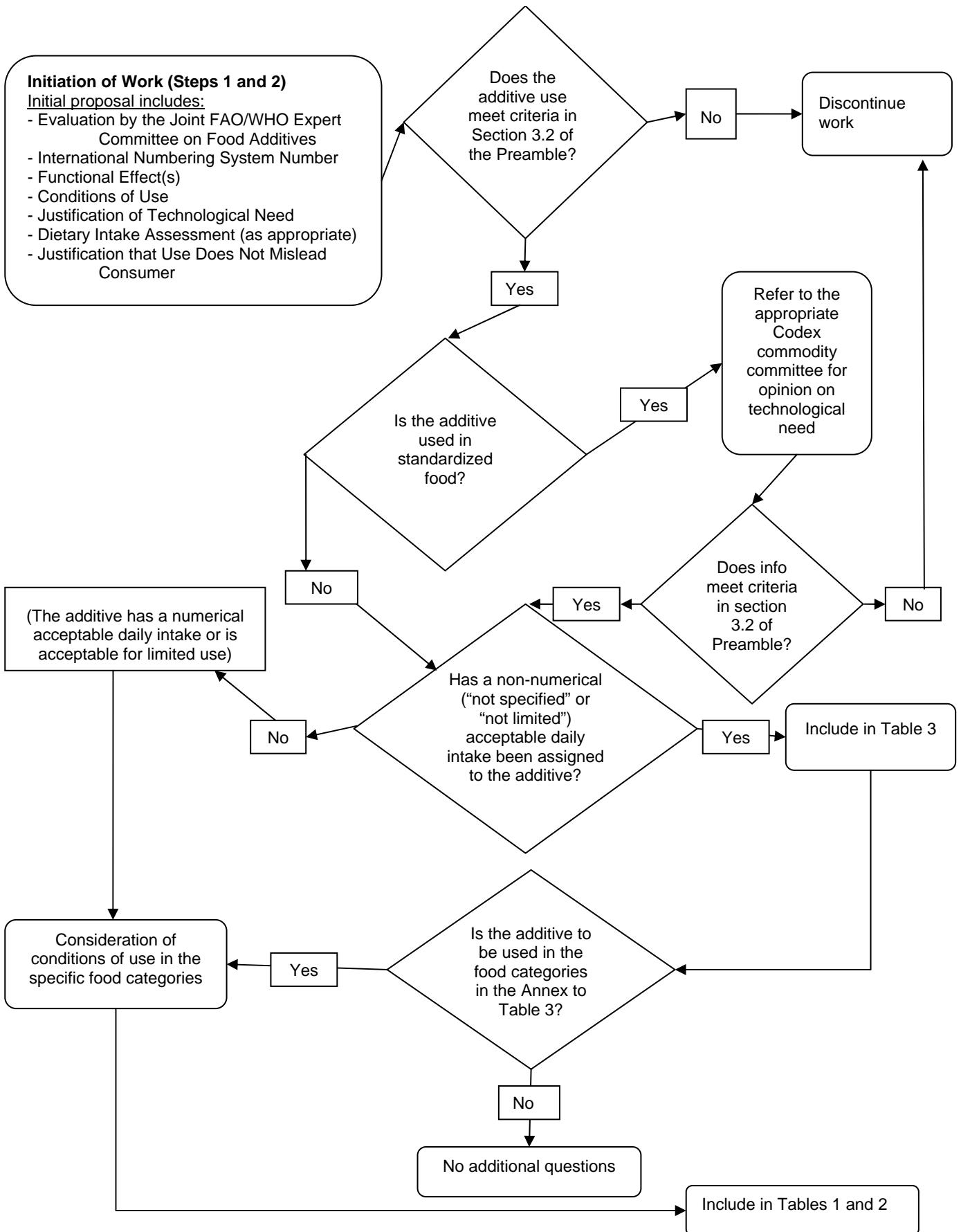
Food additives assigned a numerical ADI or evaluated to be acceptable for one or more particular uses are proposed for inclusion in Tables 1 and 2 of the General Standard for Food Additives. These proposals are considered by the Codex Committee on Food Additives according to the criteria described under “**Consideration of Conditions of Use in the Specific Food Categories**”, below.

CONSIDERATION OF CONDITIONS OF USE IN THE SPECIFIC FOOD CATEGORIES

The Codex Committee on Food Additives identifies and recommends appropriate food categories and use levels for inclusion in Tables 1 and 2 of the General Standard for Food Additives. For this purpose, the Committee will consider the following general principles for the inclusion of a food additive provision in Tables 1 and 2 of the General Standard for Food Additives:

1. Food additives that share a numerical group ADI will be considered as a group without further restrictions on the use of individual additives in that group. However, in some cases, restrictions on the use of individual additives in that group could be appropriate (e.g., because of public health concerns).
2. Food additives that have multiple functional classes will be considered without further restrictions to their functional class.
3. In general, a numerical use level for a proposed use of a food additive in a food category is given preference over a use level reported as good manufacturing practice (“GMP”). However, exceptions, as noted under “**Initiation of Work**”, shall also be taken into account by the Codex Committee on Food Additives on a case-by-case basis.
4. When establishing the acceptable maximum level of use for an additive in a specified food category, the Codex Committee on Food Additives considers the technological justification for the proposed level and the exposure assessment in accordance with Sections 3.1 and 3.2 of the Preamble of the General Standard for Food Additives. If more than one maximum use level is proposed, and the Committee cannot reach consensus on the appropriate maximum use level, the delegations supporting and the delegations opposing the proposed maximum use level should provide additional justification for their proposed levels to address any specific concerns raised by the Committee, by the next available session, to the Codex Committee on Food Additives, for consideration in its next session. Proposals lacking justification will no longer be considered, and the proposed level for which justification has been provided will be forwarded for adoption.
5. To resolve questions related to dietary exposure of food additives, the Codex Committee on Food Additives may request JECFA to perform exposure assessments for the additives based on the acceptable maximum use levels under consideration by the Codex Committee on Food Additives.
6. Acceptable maximum use levels are established as described in the previous sections and the food additive provisions are entered in the General Standard for Food Additives. Each use level represents the highest acceptable maximum use level in the broadest food category for which the use is technologically justified. To the extent possible, the hierarchical structure of the food category system will be used to simplify the listing of the food additive provisions in Tables 1 and 2 of the General Standard of Food Additives. In this regard:
 - If the new use of a food additive is for a broader food category and at a maximum use level that is higher than or equal to those in the sub-categories of the broad food category that are already listed in the General Standard for Food Additives, then the new use in the broader food category supersedes the already-listed provisions. These provisions are discontinued (if proposed draft or draft provisions), or revoked upon adoption of the proposed use at Step 8 (if adopted provision at Step 8).
 - If the new use of a food additive is for a broader food category and at a lower maximum use level than for the sub-categories of the broad food category that already exist in the General Standard for Food Additives, then the provisions listed in the General Standard for Food Additives are determined according to the hierarchy of the food category system. The highest maximum use level in each food sub-category, whether from an existing provision or from the new use in the broader food category, is entered into the General Standard for Food Additives. Any existing provisions that are superseded by the new use are discontinued (if proposed draft or draft provisions), or revoked upon adoption of the proposed use at Step 8 (if adopted provision at Step 8).
 - If the new use of a food additive, together with the already-listed provisions in the General Standard for Food Additives, represents use in all of the sub-categories of a broader food category at the same maximum use level, then the use in the broader food category will be listed in the General Standard for Food Additives. The already-listed provisions in the sub-categories are discontinued (if proposed draft or draft provisions), or revoked upon adoption of the provision in the broader food category at Step 8 (if adopted provision at Step 8).

Diagram of procedure for consideration of the entry and review of food additives in the Codex General Standard for Food Additives



AMENDMENTS TO THE PRINCIPLES FOR THE ESTABLISHMENT OR SELECTION OF CODEX SAMPLING PROCEDURES

PURPOSE OF CODEX METHODS OF SAMPLING

Codex Methods of Sampling are designed to ensure that fair and valid sampling procedures are used when food is being tested for compliance with a particular Codex commodity standard. The sampling methods are intended for use as international methods designed to avoid or remove difficulties which may be created by diverging legal, administrative and technical approaches to sampling and by diverging interpretation of results of analysis in relation to lots or consignments of foods, in the light of the relevant provision(s) of the applicable Codex standard.

METHODS OF SAMPLING

Types of Sampling Plans and Procedures

(a) Sampling Plans for Commodity Defects:

~~Such plans~~ These are normally applied to visual defects (e.g. loss of colour, ~~mis-graded for misgrading of size, etc.~~) and extraneous matter. They ~~are~~ will normally be attributes plans, and plans such as those included in Section 3.1 and 4.2 of the ~~FAO/WHO Codex Alimentarius Sampling Plans for Prepackaged Foods (AQL 6.5) General Guidelines on Sampling (CAC/GL 50-2004)~~ (hereinafter referred to as "General Guidelines") may be applied.

(b) Sampling Plans for Net Contents:

~~These~~ Such plans are ~~sampling plans~~ those which apply to pre-packaged foods generally and are intended to serve to check compliance of lots or consignments with provisions for net contents. Plans such as those included in Section 3.3 and 4.4 of the General Guidelines may be applied.

(c) Sampling Plans for Compositional Criteria:

Such plans are normally applied to analytically determined compositional criteria (e.g., loss on drying in white sugar, etc.). They are predominantly based on variable procedures with unknown standard deviation. Plans such as those included in Section 4.3 of the General Guidelines may be applied.

(d) Specific Sampling Plans for Health-related Properties:

Such plans are ~~generally~~ normally applied to heterogeneous conditions, e.g., in the assessment of microbiological spoilage, microbial by-products or sporadically occurring chemical contaminants.

General Instructions for the Selection of Methods of Sampling

~~(a) Official methods of sampling as elaborated by international organizations occupying themselves with a food or a group of foods are preferred. Such methods, when attracted to Codex standards, may be revised using Codex recommended sampling terms (to be elaborated).~~

(a) Sampling methods described in the General Guidelines or official methods of sampling elaborated by international organizations occupying themselves with a food or a group of foods are preferred. Such official methods may be written using the General Guidelines when attracted to Codex standards.

(b) When selecting appropriate sampling plans, Table 1 in the General Guidelines may be utilized.

~~(b)~~ The appropriate Codex Commodity Committee should indicate, before it elaborates any sampling plan, or before any plan is endorsed by the Codex Committee on Methods of Analysis and Sampling, the following:

- (i) the basis on which the criteria in the Codex Commodity standards have been drawn up (e.g. whether on the basis that every item in a lot, or a specified high proportion, shall comply with the provision in the standard or whether the average of a set of samples extracted from a lot must comply and, if so, whether a minimum or maximum tolerance, as appropriate, is to be given);

- (ii) whether there is to be any differentiation in the relative importance of the criteria in the standards and, if so, what is the appropriate statistical parameter each criterion should attract, and hence, the basis for judgement when a lot is in conformity with a standard.

(ed) Instructions on the procedure for the taking of samples should indicate the following:

- (i) the measures necessary in order to ensure that the sample taken is representative of the consignment or of the lot;
- (ii) the size and the number of individual items forming the sample taken from the lot or consignment;
- (iii) the administrative measures for taking and handling the sample.

(de) The sampling protocol may include the following information:

- (i) the statistical criteria to be used for acceptance or rejection of the lot on the basis of the sample;
- (ii) the procedures to be adopted in cases of dispute.

GENERAL CONSIDERATIONS

(a) The Codex Committee on Methods of Analysis and Sampling should maintain closest possible relations with all interested organizations working on methods of analysis and sampling.

(b) The Codex Committee on Methods of Analysis and Sampling should organize its work in such a manner as to keep under constant review all methods of analysis and sampling published in the Codex Alimentarius.

(c) In the Codex methods of analysis, provision should be made for variations in reagent concentrations and specifications from country to country.

(d) Codex methods of analysis which have been derived from scientific journals, theses, or publications, either not readily available or available in languages other than the official languages of FAO and WHO, or which for other reasons should be printed in the Codex Alimentarius *in extenso*, should follow the standard layout for methods of analysis as adopted by the Codex Committee on Methods of Analysis and Sampling.

(e) Methods of analysis which have already been printed as official methods of analysis in other available publications and which are adopted as Codex methods need only be quoted by reference in the Codex Alimentarius.

AMENDMENTS TO THE FORMAT FOR CODEX COMMODITY STANDARDS

FOOD ADDITIVES

This section should contain ~~the names of the additives permitted and, where appropriate, the maximum amount permitted in the food. It should be prepared in accordance with guidance given in the section on Food Additives and Contaminants in the *Relations between Commodity Committees and General Committees*, a general reference to the corresponding sections of the General Standard for Food Additives which and may should~~ take the following form:

~~“The following provisions in respect of food additives and their specifications as contained in section of the Codex Alimentarius are subject to endorsement [have been endorsed] by the Codex Committee on Food Additives.”~~

“[Food Additive functional class] used in accordance with Tables 1 and 2 of the Codex General Standard of Food Additives in food category x.x.x.x [food category name] or listed in Table 3 of the General Standard for Food Additives are acceptable for use in foods conforming to this standard.”

Exceptions from, or addition to, the General Standard for Food Additives that are necessary for its interpretation with respect to the product concerned should be justified fully, and should be restricted where possible. In cases where it is necessary to explicitly list food additives in a commodity standard, the names of the additives/functional classes permitted and, where appropriate, the maximum amount permitted in the food should be prepared in accordance with guidance given in the section on Food Additives in the *Relations between Commodity Committees and General Committees*, and may take the following form:

“The following provisions in respect of food additives and their specifications as contained in section of the Codex Alimentarius are subject to endorsement [have been endorsed] by the Codex Committee on Food Additives.”

Then should follow a tabulation, viz.:

“INS number, name of additive, maximum level (in percentage or mg/kg), grouped by functional classes.”

In this section, provisions for flavourings and processing aids should also be included.

CONTAMINANTS

Pesticide Residues:

~~This section should include, by reference, any levels for pesticide residues that have been established by the Codex Alimentarius Commission for the product concerned.~~

Other Contaminants:

~~In addition, this section should contain the names of other contaminants and where appropriate the maximum level permitted in the food, and the text to appear in the standard may take the following form:~~

~~“The following provisions in respect of contaminants, other than pesticide residues, are subject to endorsement [have been endorsed] by the Codex Committee on Contaminants in Foods.”~~

~~Then should follow a tabulation, viz.:~~

~~“Name of contaminant, maximum level (in percentage or mg/kg).”~~

This section should include the following statement:

“The products covered by this Standard shall comply with the Maximum Levels of the Codex General Standard for Contaminants and Toxins in Foods (CODEX/STAN 193-1995) and the maximum residue limits for pesticides and veterinary drugs established by the CAC.”

AMENDMENTS TO THE RELATIONS BETWEEN COMMODITY COMMITTEES AND GENERAL COMMITTEES

Codex Committees may ask the advice and guidance of committees having responsibility for matters applicable to all foods on any points coming within their province.

The Codex Committees on Food Labelling; Food Additives; Contaminants in Foods; Methods of Analysis and Sampling; Food Hygiene; Nutrition and Foods for Special Dietary Uses; and Food Import and Export Inspection and Certification Systems may establish general provisions on matters within their terms of reference. These provisions should only be incorporated into Codex Commodity Standards by reference unless there is a need for doing otherwise.

Codex Commodity standards shall contain sections on hygiene, labelling and methods of analysis and sampling and these sections should contain all of the relevant provisions of the standard. Provisions of Codex General Standards, Codes or Guidelines shall only be incorporated into Codex Commodity Standards by reference unless there is a need for doing otherwise. Where Codex Committees are of the opinion that the general provisions are not applicable to one or more commodity standards, they may request the responsible Committees to endorse deviations from the general provisions of the Codex Alimentarius. Such requests should be fully justified and supported by available scientific evidence and other relevant information. Sections on hygiene, labelling, food additives and methods of analysis and sampling which contain specific provisions or provisions supplementing the Codex General Standards, Codes or Guidelines shall be referred to the responsible Codex Committees at the most suitable time during Steps 3, 4 and 5 of the Procedure for the Elaboration of Codex Standards and Related Texts, though such reference should not be allowed to delay the progress of the standard to the subsequent steps of the Procedure.

Subject and commodity Committees should refer to the principles and guidelines developed by the Codex Committee on Food Import and Export Inspection and Certification Systems when developing provisions and/or recommendations on inspection and certification and make any appropriate amendments to the standards, guidelines and codes within the responsibility of the individual committees at the earliest convenient time.

FOOD LABELLING [no change]

FOOD ADDITIVES AND CONTAMINANTS

~~Codex commodity committees should prepare a section on food additives in each draft commodity standard and this section should contain all the provisions in the standard relating to food additives. The section should include the names of those additives which are considered to be technologically necessary or which are widely permitted for use in the food within maximum levels where appropriate.~~ shall examine the General Standard for Food Additives with a view toward incorporating a reference to the General Standard. All proposals for additions or revisions to the General Standard in order to establish a reference to the General Standard shall be referred to the Codex Committee on Food Additives. The Codex Committee on Food Additives shall consider such proposals for endorsement. Revisions of a substantive nature that are endorsed by the Food Additives Committee will be referred back to the commodity committee in order to achieve consensus between both committees at an early stage of the step procedure.

Should the Codex commodity committee consider that a general reference to the General Standard for Food Additives does not serve its purpose, a proposal should be prepared and forwarded to the Codex Committee on Food Additives for consideration and endorsement. The commodity committee shall provide a justification for why a general reference to the General Standard would not be appropriate in light of the criteria for the use of food additives established in the Preamble of the General Standard, in particular Section 3.

All provisions in respect of food additives (including processing aids) ~~and contaminants~~ contained in Codex commodity standards should be referred to the Codex Committee on Food Additives ~~or on Contaminants in Foods~~ preferably ~~after~~ before the Standards have been advanced to Step 5 of the Procedure for the Elaboration of Codex Standards or before they are considered by the Commodity Committee concerned at Step 7, though such ~~reference~~ referral should not be allowed to delay the progress of the Standard to the subsequent Steps of the Procedure.

All provisions in respect of food additives contained in commodity standards will require ~~to be endorsed~~ endorsement by the Codex Committee on Food Additives, on the basis of technological justification

submitted by the commodity committees and ~~of~~ on the recommendations of the Joint FAO/WHO Expert Committee on Food Additives concerning the safety-in-use (acceptable daily intake (ADI) and other restrictions) and an estimate of the potential and, where possible, the actual intake of the food additives, ensuring conformity with the ~~General Principles for the Use of Food Additives~~ Preamble of General Principles for the Use of Food Additives.

~~In preparing working papers for the~~ When forwarding a food additive section of a commodity standard for endorsement by Codex Committee on Food Additives, the Secretariat should prepare ~~make~~ a report to the Committee concerning the endorsement of provisions for food additives (including processing aids), ~~on the basis of the General Principles for the Use of Food Additives. Provisions for food additives should~~ that includes the functional classes and technological justification. With regard to exceptional cases where specific food additives and their maximum levels are given, the report should also indicate the International Numbering System (INS) number, the Acceptable Daily Intake (ADI) assigned by the Joint FAO/WHO Expert Committee on Food Additives, technological justification, proposed level, and whether the additive was previously endorsed (or temporarily endorsed) by the Codex Committee on Food Additives.

When commodity standards are sent to governments for comment at Step 3, they should contain a statement that the provisions “in respect of food additives ~~and contaminants~~ are subject to endorsement by the Codex Committees on Food Additives ~~or on Contaminants in Foods~~ and to incorporation into the General Standard for Food Additives ~~or the General Standard for Contaminants and Toxins in Foods~~”.-

~~When establishing provisions for food additives, Codex committees should follow the General Principles for the Use of Food Additives and the Preamble of the General Standard for Food Additives. Full explanation should be provided for any departure from the above recommendations.~~

When an active commodity committee exists, proposals for the use of additives in any commodity standard under consideration should be prepared by the committee concerned, and forwarded to the Codex Committee on Food Additives for endorsement and inclusion in the General Standard for Food Additives. When the Codex Committee on Food Additives decides not to endorse specific additives provisions (~~use of the additive, or level in the end product~~), the reason should be clearly stated. The section under consideration should be referred back to the commodity ~~C~~committee concerned if further information is needed, or for information if the Codex Committee on Food Additives decides to amend the provision.

When no active commodity committee exists, proposals for new additive provisions or amendment of existing provisions for inclusion in the General Standard for Food Additives should be forwarded directly by Codex member ~~countries~~ to the Codex Committee on Food Additives.

Good Manufacturing Practice means that:

- ~~the quantity of the additive added to food does not exceed the amount reasonably required to accomplish its intended physical nutritional or other technical effect in food;~~
- ~~the quantity of the additive that becomes a component of food as a result of its use in the manufacturing, processing or packaging of a food and which is not intended to accomplish any physical, or other technological effect in the food itself, is reduced to the extent reasonably possible;~~

~~the additive is of appropriate food grade quality and is prepared and handled in the same way as a food ingredient. Food grade quality is achieved by compliance with the specifications as a whole and not merely with individual criteria in terms of safety.~~

Move the above definition of Good Manufacturing Practice in the Use of Food Additives to section “Definitions for the Purposes of the Codex Alimentarius”

FOOD HYGIENE [no change]

METHODS OF ANALYSIS AND SAMPLING [no change]

RISK ANALYSIS PRINCIPLES APPLIED BY THE CODEX COMMITTEE ON PESTICIDE RESIDUES

[for inclusion in Section III]

SCOPE

1. This document addresses the respective applications of risk analysis principles by the Codex Committee on Pesticide Residues (CCPR) as the risk management body and the Joint FAO/WHO Meeting on Pesticide Residues (JMPR) as the risk assessment body and facilitates the uniform application of the Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius. This document should be read in conjunction with the Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius.

ROLES OF CCPR AND JMPR IN RISK ANALYSIS

INTERACTION BETWEEN CCPR AND JMPR

2. In addressing pesticide residue issues in Codex, providing advice on risk management is the responsibility of the Codex Alimentarius Commission (CAC) and CCPR while conducting risk assessment is the responsibility of JMPR.

3. CCPR and JMPR recognize that an adequate communication between risk assessors and risk managers is an essential requirement for successfully performing their risk analysis activities.

4. CCPR and JMPR should continue to develop procedures to enhance communication between the two bodies.

5. CCPR and JMPR should ensure that their respective contributions to the risk analysis process result in outputs that are scientifically based, fully transparent, thoroughly documented and available in a timely manner to members².

6. JMPR, in consultation with CCPR, should continue to explore developing minimum data requirements necessary for JMPR to perform risk assessments.

7. These requirements should be used by CCPR as a fundamental criterion as described in the Annex in preparing its Priority List for JMPR. The JMPR Secretariat should consider whether these minimum data requirements have been met when preparing the provisional agenda for meetings of JMPR.

ROLE OF CCPR

8. CCPR is primarily responsible for recommending risk management proposals for adoption by the CAC.

9. CCPR shall base its risk management recommendations, such as MRLs, to the CAC following JMPR's risk assessments of the respective pesticides, and considering, where appropriate, other legitimate factors such as relevant to the health protection of consumers and for the promotion of fair practices in food trade.

10. In cases where JMPR has performed a risk assessment and CCPR or the CAC determines that additional scientific guidance is necessary, CCPR or CAC may make a specific request to JMPR to provide further scientific guidance necessary for a risk management decision.

11. CCPR's risk management recommendations to the CAC shall take into account the relevant uncertainties as described by JMPR.

² Submission and evaluation of pesticide residues data for the estimation of maximum residue levels in food and feed; FAO Plant Production and Protection Paper, 170, 2002, ISBN 92-5-104759-6

12. CCPR shall consider maximum residue limits (MRLs) only for those pesticides for which JMPR has completed a full safety evaluation.
13. CCPR shall base its recommendations on the GEMS/Food diets used to identify consumption patterns on a global scale when recommending MRLs in food. The GEMS/Food diets are used to assess the risk of chronic exposure. The acute exposure calculations are not based on those diets, but available consumption data provided by members.
14. When establishing its standards, CCPR shall clearly state when it applies any considerations based on other legitimate factors in addition to JMPR's risk assessment and recommended maximum residue levels and specify its reasons for doing so.
15. CCPR shall consider the following when preparing its priority list of compounds for JMPR evaluation:
 - CCPR's Terms of Reference;
 - JMPR's Terms of Reference;
 - The Codex Alimentarius Commission's Strategic Plan;
 - The Criteria for the Establishment of Work Priorities;
 - The Criteria for Inclusion of Compounds on the Priority List;
 - The Criteria for Selecting Food Commodities for which Codex MRLs or Extraneous Maximum Residue Limits (EMRLs) should be Established;
 - The Criteria for Evaluation of New Chemicals;
 - The Criteria for Prioritization Process of Compounds for Evaluation by JMPR
 - A commitment to provide the necessary data for the evaluation in time.
16. When referring substances to JMPR, the CCPR shall provide background information and clearly specify the reasons for the request when chemicals are nominated for evaluation.
17. When referring substances to JMPR, the CCPR may also refer a range of risk management options, with a view toward obtaining JMPR's guidance on the attendant risks and the likely risk reductions associated with each option.
18. CCPR shall request JMPR to review any methods and guidelines being considered by CCPR for assessing maximum limits for pesticides.

ROLE OF JMPR

19. The Joint FAO/WHO Meeting on Pesticide Residues (JMPR) consists of the FAO Panel of Experts on Pesticide Residues in Food and the Environment and the WHO Core Assessment Group. It is an independent scientific expert body convened by both Directors General of FAO and WHO according to the rules of both organizations, charged with the task to provide scientific advice on pesticide residues.
20. This guidance document applies to the work of JMPR in the context of Codex and in particular as it relates to advice requests from CCPR.
21. JMPR is primarily responsible for performing the risk assessments upon which CCPR and ultimately the CAC base their risk management decisions. JMPR also proposes MRLs based on Good Agricultural Practices (GAPs)/ registered uses or in specific cases, such as EMRLs, based on monitoring data.
22. JMPR provides CCPR with science-based risk assessments that include the four components of risk assessment as defined by CAC and safety assessments that can serve as the basis for CCPR's risk-management discussions. JMPR should continue to use its risk assessment process for establishing Acceptable Daily Intakes (ADIs) and Acute Reference Doses (ARfDs) where appropriate.

23. JMPR should identify and communicate to CCPR in its assessments any information on the applicability and any constraints of the risk assessment to the general population and to particular sub-populations and will as far as possible identify potential risks to populations of potentially enhanced vulnerability (e.g. children).
24. JMPR is responsible for evaluating exposure to pesticides. JMPR should strive to base its exposure assessment and hence the dietary risk assessments on global data, including that from developing countries. In addition to GEMS/Food data, monitoring data and exposure studies may be used. The GEMS/Food diets are used to assess the risk of chronic exposure. The acute exposure calculations are not based on those diets, but on the available high percentile consumption data as provided by members.
25. JMPR should communicate to CCPR the magnitude and source of uncertainties in its risk assessments. When communicating this information, JMPR should provide CCPR a description of the methodology and procedures by which JMPR estimated any uncertainty in its risk assessment.
26. JMPR should communicate to CCPR the basis for all assumptions used in its risk assessments.

ANNEX: LIST OF RISK MANAGEMENT POLICIES USED BY CCPR

1. This part of the document addresses the risk management policy that is used by the Codex Committee on Pesticides Residues (CCPR) when discussing the risk assessments, the exposure to pesticides and the proposals for MRLs which are the outcomes of the Joint FAO/WHO Meeting on Pesticides Residues (JMPR).

ESTABLISHMENT OF MRLs/EMRLs

Procedure for Proposing Pesticides for Codex Priority Lists

2. CCPR has developed a policy document in relation to establishing a priority list of pesticides for evaluation or re-evaluation by JMPR³.
3. Before a pesticide can be considered for the Priority List, it must:
- be available for use as a commercial product; and
 - not have been already accepted for consideration.
4. To meet the criteria for inclusion in the priority list, the use of the pesticide must: give rise to residues in or on a food or feed commodity moving in international trade, the presence of which is (or may be) a matter of public health concern and thus create (or have the potential to create) problems in international trade.
5. When prioritising new chemicals for evaluation by the JMPR, the Committee will consider the following criteria:
- (i) If the chemical has a reduced acute and/or chronic toxicity risk to humans compared with other chemicals in its classification (insecticide, fungicide, herbicide);
 - (ii) The date when the chemical was nominated for evaluation;
 - (iii) Commitment by the sponsor of the compound to provide supporting data for review with a firm date for data submission;
 - (iv) The availability of regional/national reviews and risk assessments, and coordination with other regional/national lists; and
 - (v) Allocating priorities to new chemicals, so that at least 50% of evaluations are for new chemicals, if possible.

³ Criteria for Prioritization Process of Compounds for Evaluation by JMPR, Procedural Manual

6. When prioritising chemicals for periodic re-evaluation by the JMPR, the Committee will consider the following criteria:

- (i) If the intake and/or toxicity profile indicate some level of public health concern;
- (ii) Chemicals that have not been reviewed toxicologically for more than 15 years and/or not having a significant review of maximum residue limits for 15 years;
- (iii) The year the chemical is listed in the list for Candidate Chemicals for Periodic Re-evaluation –Not Yet Scheduled;
- (iv) The date that data will be submitted;
- (v) Whether the CCPR has been advised by a national government that the chemical has been responsible for trade disruption;
- (vi) If there is a closely related chemical that is a candidate for periodic re-evaluation that can be evaluated concurrently; and
- (vii) The availability of current labels arising from recent national re-evaluations.

7. Once the JMPR has reviewed a chemical, three scenarios may occur:

- the data confirm the existing Codex MRL, it remains in place, or
- a new MRL is recommended or an amendment of an existing MRL. The new or amended proposal enters at Step 3 of the Codex procedure. The existing MRL remains in place for no more than four years, or
- insufficient data have been submitted to confirm or amend an existing Codex MRL. The Codex MRL is recommended for withdrawal. However, the manufacturer or countries may provide a commitment to the JMPR and CCPR to provide the necessary data for review within four years. The existing Codex MRL is maintained for a period of no more than four years pending the review of the additional data. A second period of four years is not granted.

MRLs for Commodities of Animal Origin

8. Farm animal metabolism studies are required whenever a pesticide is applied directly to livestock, to animal premises or housing, or when significant residues remain in crops or commodities used in animal feed, in forage crops, or in plant parts that could be used in animal feeds. The results of farm animal feeding studies and residues in animal feed serve also as a primary source of information for estimating maximum residue levels in animal products.

9. If no adequate studies are available, no MRLs will be established for commodities of animal origin. MRLs for feeds (and the primary crops) should not be established in the absence of animal transfer data. Where the exposure of livestock to pesticides through feeds leads to residues at the limit of quantitation, MRLs at the LOQ must be established for animal commodities. MRLs should be established for all mammalian species where pesticides on feeds are concerned and for specific species (e.g. cattle, sheep) where direct treatments of pesticides are concerned.

10. Where the recommended maximum residue limits for animal commodities resulting from direct treatment of the animal, regardless of whether they are recommended by JMPR or JECFA, and from residues in animal feed do not agree, the higher recommendation will prevail.

MRLs for Processed or Ready-to-eat Foods or Feeds

11. CCPR agreed not to establish MRLs for processed foods and feeds unless separate higher MRLs are necessary for specific processed commodities.

MRLs for spices

12. CCPR agreed that MRLs for spices can be established on the basis of monitoring data in accordance with the guidelines established by JMPR.

MRLs for fat-soluble pesticides

13. If a pesticide is determined as “fat soluble” after consideration of the following factors, it is indicated with the text “The residues are fat soluble” in the residue definition:

- When available, it is the partitioning of the residue (as defined) in muscle versus fat in the metabolism studies and livestock feeding studies that determines the designation of a residue as being “fat soluble”.
- In the absence of useful information on the distribution of residues in muscle and fat, residues with $\log Pow > 3$ are likely to be “fat soluble”

14. For fat soluble pesticides, two MRLs are recommended if data permit: one for whole milk and one for milk fat. For enforcement purposes, a comparison can be made either of the residue in milk fat with the MRL for milk fat or of the residue in whole milk with the MRL for milk.

Establishment of MRLs

15. The CCPR is entrusted with the elaboration of Maximum Residue Limits (MRLs) of pesticide residues in food and feed. The JMPR is using the WHO Guidelines for predicting dietary intake of pesticides residues (revised)(1997)⁴. The JMPR is recommending MRLs establishing Supervised Trial Median Residues (STMRs) for new and periodic review compounds for dietary intake purposes. In cases the intake exceeds the Acceptable Daily Intake (ADI) in one or more of the regional diets, the JMPR, when recommending MRLs, flags this situation indicating the type of data which may be useful to further refine the dietary intake estimate.

16. When the ADI is exceeded in one or more regional diets, then the MRLs will not advance to Step 8 pending further refinement of the intake at the international level. If further refinement is not possible then MRLs are withdrawn until the remaining MRLs give no longer rise to intake concerns. This procedure should be reviewed at regular interval.

17. The JMPR is currently routinely establishing acute reference doses (ARfDs), where appropriate, and indicates cases where an ARfD is not necessary. The 1999 JMPR for the first time calculated the short-term dietary intake estimates following an approach using the International and National Estimates of Short-term Intake (IESTI, NESTI). The procedure allows for estimating the short-term risk for relevant subgroups of the population, like children. The JMPR flags cases when the IESTI for a given commodity exceeds the ARfD.

18. When the ARfD is exceeded for a given commodity, then the MRLs will not advance to Step 8 pending further refinement of the intake at the international level.

19. When a Draft MRL has been returned to Step 6 three times, the CCPR should ask JMPR to examine residue data from other appropriate GAPs and to recommend MRLs which cause no dietary intake concerns if possible.

20. If further refinement is not possible then MRLs are withdrawn. More sophisticated methodologies such as probabilistic approaches are under investigation at the moment.

21. The estimate of the short-term dietary intake requires substantial food consumption data that currently are only sparsely available. Governments are urged to generate relevant consumption data and to submit these data to the WHO.

Utilization of Steps 5/8 for elaboration of MRLs

22. Preconditions for utilization of Step 5/8 Procedure

- New MRL circulated at Step 3
- JMPR report available electronically by early February
- No intake concerns identified by JMPR

23. Steps 5/8 Procedure (Recommendation to omit Steps 6 and 7 and adopt the MRL at Step 8)

⁴ Programme of Food Safety and Food Aid; WHO/FSF/FOS/97.7

- If the preconditions listed above are met.
- If a delegation has a concern with advancing a given MRL, a concern form should be completed detailing the concern along with a description of the data that will be submitted to substantiate the concern preferably as comments at Step 3, or at the latest, one month after the CCPR session.
- If the JMPR Secretariat or the CCPR can address that concern at the upcoming CCPR session, and the JMPR position remains unchanged, the CCPR will decide if the MRL will be advanced to Step 5/8.
- If the concern cannot be addressed at the meeting, the MRL will be advanced to Step 5 at the CCPR session and the concern will be addressed by the JMPR as soon as possible but the rest of the MRLs should be advanced to Step 5/8.
- The result of the consideration of the concern by the JMPR will be considered at the next CCPR session. If the JMPR position remains unchanged, the CCPR will decide if the MRL will be advanced to Step 8.

Establishment of EMRLs

24. The Extraneous Maximum Residue Limit (EMRL) refers to a pesticide residue or a contaminant arising from environmental sources (including former agricultural uses) other than the use of the pesticide or contaminant substance directly or indirectly on the commodity. It is the maximum concentration of a pesticide residue that is recommended by the Codex Alimentarius Commission to be legally permitted or recognized as acceptable in or on a food, agricultural commodity or animal feed.

25. Chemicals for which EMRLs are most likely to be needed are persistent in the environment for a relatively long period after uses have been discontinued and are expected to occur in foods or feeds at levels of sufficient concern to warrant monitoring.

26. All relevant and geographically representative monitoring data (including nil-residue results) are required to make reasonable estimates to cover international trade. JMPR has developed a standard format for reporting pesticide residues monitoring data⁵.

27. The JMPR compares data distribution in terms of the likely percentages of violations that might occur if a given EMRL is proposed to the CCPR.

28. Because residues gradually decrease, CCPR evaluates every 5 years, if possible, the existing EMRLs, based on the reassessments of the JMPR.

29. The CCPR generally agreed at the 30th Session on the potential elements for inclusion in a set of criteria for estimation of EMRLs while it also agreed not to initiate a full exercise of criteria elaboration.

Periodic Review Procedure

30. The Committee agreed on the Periodic Review Procedure, which was endorsed by the CAC and attached to the list of MRLs prepared for each session of the CCPR. Those Codex MRLs confirmed by JMPR under the Periodic Review shall be distributed to members and interested organizations for comments.

Deleting Codex MRLs

31. Every year new compounds are introduced. These compounds are often new pesticides which are safer than existing ones. Old compounds are then no longer supported/produced by industry and existing Codex MRLs can be deleted.

⁵ Submission and evaluation of pesticide residues data for the estimation of maximum residue levels in food and feed; FAO Plant Production and Protection Paper, 170, 2002, ISBN 92-5-104759-6

32. If information is delivered between two sessions of CCPR, that a certain compound is no longer supported, this information will be shared during the first coming session (t=0). The proposal will be to delete the existing MRLs at the following session (t=0+1 year).

33. It may happen that compounds are no longer supported in Codex, but are supported in some selected countries. If there is no international trade in commodities where the active compounds may have been used, CCPR will not establish MRLs.

MRLs AND METHODS OF ANALYSIS

34. JMPR needs data and information for their evaluations. Among these are methods of analysis. Methods should include specialized methods used in supervised trials and enforcement methods.

35. If no methods of analysis are available for enforcing MRLs for a specific compound, no MRLs will be established by CCPR.

RISK ANALYSIS PRINCIPLES APPLIED BY THE CODEX COMMITTEE ON RESIDUES OF VETERINARY DRUGS IN FOODS

[for inclusion in Section III]

SECTION 1. PURPOSE – SCOPE

1. The purpose of this document is to specify Risk Analysis Principles applied by the Codex Committee on Residues of Veterinary Drugs in Foods.

SECTION 2. PARTIES INVOLVED

2. The *Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius* has defined the responsibilities of the various parties involved. The responsibility for providing advice on risk management concerning residues of veterinary drugs lies with the Codex Alimentarius Commission and its subsidiary body, the Codex Committee on Residues of Veterinary Drugs in Foods (CCRVDF), while the responsibility for risk assessment lies primarily with the Joint FAO/WHO Expert Committee on Food Additives (JECFA).

3. According to its mandate, the responsibilities of the CCRVDF regarding veterinary drug residues in food are:

- (a) to determine priorities for the consideration of residues of veterinary drugs in foods;
- (b) to recommend maximum residue limits (MRLs) for such veterinary drugs;
- (c) to develop codes of practice as may be required;
- (d) to consider whether available methods of sampling and analysis for the determination of veterinary drug residues in foods.

4. The CCRVDF shall base its risk management recommendations to the Codex Alimentarius Commission on JECFA's risk assessments of veterinary drugs in relation to proposed MRLs.

5. The CCRVDF is primarily responsible for recommending risk management proposals for adoption by the Codex Alimentarius Commission.

6. JECFA is primarily responsible for providing independent scientific advice, the risk assessment, upon which the CCRVDF base their risk management decisions. It assists the CCRVDF by evaluating the available scientific data on the veterinary drug prioritised by the CCRVDF. JECFA also provides advice directly to FAO and WHO and to Member governments.

7. Scientific experts from JECFA are selected in a transparent manner by FAO and WHO under their rules for expert committees on the basis of the competence, expertise, experience in the evaluation of compounds used as veterinary drugs and their independence with regard to the interests involved, taking into account geographical representation where possible.

SECTION 3. RISK MANAGEMENT IN CCRVDF

8. Risk management should follow a structured approach including:

- preliminary risk management activities;
- evaluation of risk management options; and
- monitoring and review of decisions taken.

9. The decisions should be based on risk assessment, and take into account, where appropriate, other legitimate factors relevant for the health protection of consumers and for fair practices in food trade, in accordance with the *Criteria for the Consideration of the Other Factors Referred to in the Second Statement of Principles*⁶.

⁶ *Statements of Principle Concerning the Role of Science in the Codex Decision-making Process and the Extent to Which Other Factors are Taken into Account*, Codex Procedural Manual - Appendix

3.1 Preliminary risk management activities

10. This first phase of risk management covers:
- Establishment of risk assessment policy for the conduct of the risk assessments;
 - Identification of a food safety problem;
 - Establishment of a preliminary risk profile;
 - Ranking of the hazard for risk assessment and risk management priority;
 - Commissioning of the risk assessment; and
 - Consideration of the result of the risk assessment.

3.1.1 Risk Assessment Policy for the Conduct of the Risk Assessment

11. The responsibilities of the CCRVDF and JECFA and their interactions along with core principles and expectations of JECFA evaluations are provided in *Risk Assessment Policy for the Setting of MRLs in Food*, established by the Codex Alimentarius Commission.

3.1.2 Establishment of Priority List

12. The CCRVDF identifies, with the assistance of Members, the veterinary drugs that may pose a consumer safety problem and/or have a potential adverse impact on international trade. The CCRVDF establishes a priority list for assessment by JECFA.

13. In order to appear on the priority list of veterinary drugs for the establishment of a MRL, the proposed veterinary drug shall meet some or all of the following criteria:

- A Member has proposed the compound for evaluation;
- A Member has established good veterinary practices with regard to the compound;
- The compound has the potential to cause public health and/or international trade problems;
- It is available as a commercial product; and
- There is a commitment that a dossier will be made available.

14. The CCRVDF takes into account the protection of confidential information in accordance with WTO Agreement on Trade-Related Aspects of Intellectual Property Rights (TRIPS) - Section 7: Protection of Undisclosed Information - Article 39, and makes every effort to encourage the willingness of sponsors to provide data for JECFA assessment.

3.1.3 Establishment of a Preliminary Risk Profile

15. Member(s) request(s) the inclusion of a veterinary drug on the priority list. The available information for evaluating the request shall be provided either directly by the Member(s) or by the sponsor. A preliminary risk profile shall be developed by the Member(s) making the request, using the template presented in the Annex.

16. The CCRVDF considers the preliminary risk profile and makes a decision on whether or not to include the veterinary drug in the priority list.

3.1.4 Ranking of the Hazard for Risk Assessment and Risk Management Priority

17. The CCRVDF establishes an ad-hoc Working Group open to all its Members and observers, to make recommendations on the veterinary drugs to include into (or to remove from) the priority list of veterinary drugs for the JECFA assessment. The CCRVDF considers these recommendations before agreeing on the priority list, taking into account pending issues such as temporary Acceptable Daily Intakes (ADIs) and/or MRLs. In its report, the CCRVDF shall specify the reasons for its choice and the criteria used to establish the order of priority.

18. Prior to development of MRLs for new veterinary drugs not previously evaluated by JECFA, a proposal for this work shall be sent to the Codex Alimentarius Commission with a request for approval as new work in accordance with the *Procedures for the Elaboration of Codex Standards and Related Texts*.

3.1.5 Commissioning of the Risk Assessment

19. After approval by the Codex Alimentarius Commission of the priority list of veterinary drugs as new work, the CCRVDF forwards it to JECFA with the qualitative preliminary risk profile as well as specific guidance on the CCRVDF risk assessment request. JECFA, WHO and FAO experts then proceed with the assessment of risks related to these veterinary drugs, based on the dossier provided and/or all other available scientific information.

3.1.6 Consideration of the Result of the Risk Assessment

20. When the JECFA risk assessment is completed, a detailed report is prepared for the subsequent session of the CCRVDF for consideration. This report shall clearly indicate the choices made during the risk assessment with respect to scientific uncertainties and the level of confidence in the studies provided.

21. When the data are insufficient, JECFA may recommend temporary MRL on the basis of a temporary ADI using additional safety considerations⁷. If JECFA cannot propose an ADI and/or MRLs due to lack of data, its report should clearly indicate the gaps and a timeframe in which data should be submitted, in order to allow Members to make an appropriate risk management decision.

22. The JECFA assessment reports related to the concerned veterinary drugs should be made available in sufficient time prior to a CCRVDF meeting to allow for careful consideration by Members. If this is, in exceptional cases, not possible, a provisional report should be made available.

23. JECFA should, if necessary, propose different risk management options. In consequence, JECFA should present, in its report, different risk management options for the CCRVDF to consider. The reporting format should clearly distinguish between the risk assessment and the evaluation of the risk management options.

24. The CCRVDF may ask JECFA any additional explanation.

25. Reasons, discussions and conclusions (or the absence thereof) on risk assessment should be clearly documented, in JECFA reports, for each option reviewed. The risk management decision taken by the CCRVDF (or the absence thereof) should also be fully documented.

3.2 Evaluation of Risk Management Options

26. The CCRVDF shall proceed with a critical evaluation of the JECFA proposals on MRLs and may consider other legitimate factors relevant for health protection and fair trade practices in the framework of the risk analysis. According to the 2nd statement of principle, the criteria for the consideration of other factors should be taken into account. These other legitimate factors are those agreed during the 12th session of the CCRVDF⁸ and subsequent amendments made by this Committee.

27. The CCRVDF either recommends the MRLs as proposed by JECFA, modifies them in consideration of other legitimate factors, considers other measures or asks JECFA for reconsideration of the residue evaluation for the veterinary drug in question.

28. Particular attention should be given to availability of analytical methods used for residue detection.

3.3 Monitoring and Review of the Decisions Taken

29. Members may ask for the review of decisions taken by the Codex Alimentarius Commission. To this end, veterinary drugs should be proposed for inclusion in the priority list. In particular, review of decisions may be necessary if they pose difficulties in the application of the *Guidelines for the Establishment of a Regulatory Programme for the Control of Veterinary Drug Residues in Foods* (CAC/GL 16-1993).

30. The CCRVDF may request JECFA to review any new scientific knowledge and other information relevant to risk assessment and concerning decisions already taken, including the established MRLs.

⁷ Definition of "Codex maximum limit for residues of veterinary drugs", Codex Procedural Manual.

⁸ ALINORM 01/31 paragraph 11.

31. The risk assessment policy for MRL shall be reconsidered based on new issues and experience with the risk analysis of veterinary drugs. To this end, interaction with JECFA is essential. A review may be undertaken of the veterinary drugs appearing on prior JECFA agendas for which no ADI or MRL has been recommended.

SECTION 4. RISK COMMUNICATION IN THE CONTEXT OF RISK MANAGEMENT

32. In accordance with the *Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius*, the CCRVDF, in cooperation with JECFA, shall ensure that the risk analysis process is fully transparent and thoroughly documented and that results are made available in a timely manner to Members. The CCRVDF recognises that communication between risk assessors and risk managers is critical to the success of risk analysis activities.

33. In order to ensure the transparency of the assessment process in JECFA, the CCRVDF provides comments on the guidelines related to assessment procedures being drafted or published by JECFA.

ANNEX: TEMPLATE FOR INFORMATION NECESSARY FOR PRIORITIZATION BY CODEX COMMITTEE ON RESIDUES OF VETERINARY DRUGS IN FOODS

Administrative information

1. Member(s) submitting the request for inclusion
2. Veterinary drug names
3. Trade names
4. Chemical names
5. Names and addresses of basic producers

Purpose, scope and rationale

6. Identification of the food safety issue (residue hazard)
7. Assessment against the criteria for the inclusion on the priority list

Risk profile elements

8. Justification for use
9. Veterinary use pattern
10. Commodities for which Codex MRLs are required

Risk assessment needs and questions for the risk assessors

11. Identify the feasibility that such an evaluation can be carried out in a reasonable framework
12. Specific request to risk assessors

Available information⁹

13. Countries where the veterinary drugs is registered
14. National/Regional MRLs or any other applicable tolerances
15. List of data (pharmacology, toxicology, metabolism, residue depletion, analytical methods) available

Timetable

16. Date when data could be submitted to JECFA

⁹ When preparing a preliminary risk profile, Member(s) should take into account the updated data requirement, to enable evaluation of a veterinary drug for the establishment of an ADI and MRLs, published by JECFA.

RISK ASSESSMENT POLICY FOR THE SETTING OF MAXIMUM LIMITS FOR RESIDUES OF VETERINARY DRUGS IN FOODS

[for inclusion in Section III]

Role of JECFA

1. The Joint FAO/WHO Expert Committee on Food Additives (JECFA) is an independent scientific expert body convened by both Directors-General of FAO and WHO according to the rules of both organizations, charged with the task to provide scientific advice on veterinary drug residues in food.
2. This annex applies to the work of JECFA in the context of Codex and in particular as it relates to advice requests from the Codex Committee on Residues of Veterinary Drugs in Foods (CCRVDF).
 - (a) JECFA provides CCRVDF with science-based risk assessments conducted in accordance with the *Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius* and incorporating the four steps of risk assessment. JECFA should continue to use its risk assessment process for establishing Acceptable Daily Intakes (ADIs) and proposing Maximum Residues Limits (MRLs).
 - (b) JECFA should take into account all available scientific data to establish its risk assessment. It should use available quantitative information to the greatest extent possible and also qualitative information.
 - (c) Constraints, uncertainties and assumptions that have an impact on the risk assessment need be clearly communicated by JECFA.
 - (d) JECFA should provide CCRVDF with information on the applicability, public health consequences and any constraints of the risk assessment to the general population and to particular sub-populations and, as far as possible, should identify potential risks to specific group of populations of potentially enhanced vulnerability (e.g. children).
 - (e) Risk assessment should be based on realistic exposure scenarios.
 - (f) When the veterinary drug is used both in veterinary medicine and as a pesticide, a harmonised approach between JECFA and the Joint FAO/WHO Meeting on Pesticide Residues (JMPR) should be followed.
 - (g) MRLs, that are compatible with the ADI, should be set for all species based on appropriate consumption figures. When requested by CCRVDF, extension of MRLs between species will be considered if appropriate data are available.

Data Protection

3. Considering the importance of intellectual property in the context of data submission for scientific evaluation, JECFA has established procedures to cover the confidentiality of certain data submitted. These procedures enable the sponsor to declare which data is to be considered as confidential. The procedure includes a formal consultation with the sponsor.

Expression of Risk Assessment Results in terms of MRLs

4. MRLs have to be established for target animal tissues (e.g. muscle, fat, or fat and skin, kidney, liver), and specific food commodities (e.g. eggs, milk, honey) originating from the target animals species to which a veterinary drug can be administered according to good veterinary practice.
5. However, if residue levels in various target tissues are very different, JECFA is requested to consider MRLs for a minimum of two. In this case, the establishment of MRLs for muscle or fat is preferred to enable the control of the safety of carcasses moving in international trade.
6. When the calculation of MRLs to be compatible with the ADI may be associated with a lengthy withdrawal period, JECFA should clearly describe the situation in its report.

AMENDMENTS TO THE RISK ANALYSIS PRINCIPLES APPLIED BY THE CODEX COMMITTEE ON FOOD ADDITIVES AND CONTAMINANTS

RISK ANALYSIS PRINCIPLES APPLIED BY THE CODEX COMMITTEE ON FOOD ADDITIVES AND THE CODEX COMMITTEE ON CONTAMINANTS IN FOODS ~~CODEX COMMITTEE ON FOOD ADDITIVES AND CONTAMINANTS~~

SECTION 1. SCOPE

- 1) This document addresses the respective applications of risk analysis principles by the ~~Codex Committee on Food Additives and Contaminants (CCFAC)~~ Codex Committee on Food Additives (CCFA) and the Codex Committee on Contaminants in Foods (CCCF) and the Joint FAO/WHO Expert Committee on Food Additives (JECFA). For matters which cannot be addressed by JECFA, this document does not preclude the possible consideration of recommendations arising from other internationally recognized expert bodies, as approved by the Commission.
- 2) This document should be read in conjunction with the *Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius*.

SECTION 2. ~~CCFAC~~ CCFA/CCCF and JECFA

- 3) ~~CCFAC~~ CCFA/CCCF and JECFA recognize that communication between risk assessors and risk managers is critical to the success of their risk analysis activities.
- 4) ~~CCFAC~~ CCFA/CCCF and JECFA should continue to develop procedures to enhance communication between the two committees.
- 5) ~~CCFAC~~ CCFA/CCCF and JECFA should ensure that their contributions to the risk analysis process involve all interested parties and are fully transparent and thoroughly documented. While respecting legitimate concerns to preserve confidentiality, documentation should be made available, upon request, in a timely manner to all interested parties.
- 6) JECFA, in consultation with ~~CCFAC~~ CCFA/CCCF, should continue to explore developing minimum quality criteria for data requirements necessary for JECFA to perform risk assessments. These criteria are used by ~~CCFAC~~ CCFA/CCCF in preparing ~~its~~ their Priority List for JECFA. The JECFA Secretariat should consider whether these minimum quality criteria for data have been met when preparing the provisional agenda for meetings of JECFA.

SECTION 3. ~~CCFAC~~ CCFA/CCCF

- 7) ~~CCFAC~~ CCFA/CCCF ~~is~~ are primarily responsible for recommending risk management proposals for adoption by the CAC.
- 8) ~~CCFAC~~ CCFA/CCCF shall base ~~its~~ their risk management recommendations to the CAC on JECFA's risk assessments, including safety assessments¹⁰, of food additives, naturally occurring toxicants, and contaminants in food.
- 9) In cases where JECFA has performed a safety assessment and ~~CCFAC~~ CCFA/CCCF or the CAC determines that additional scientific guidance is necessary, ~~CCFAC~~ CCFA/CCCF or CAC may make a more specific request to JECFA to obtain the scientific guidance necessary for a risk management decision.
- 10) ~~CCFAC~~ CCFA's risk management recommendations to the CAC with respect to food additives shall be guided by the principles described in the Preamble and relevant annexes of the Codex General Standard for Food Additives.

¹⁰ A Safety Assessment is defined as a scientifically-based process consisting of: 1) the determination of a NOEL (No Observed Effect Level) for a chemical, biological, or physical agent from animal feeding studies and other scientific considerations; 2) the subsequent application of safety factors to establish an ADI or tolerable intake; and 3) comparison of the ADI or tolerable intake with probable exposure to the agent (Temporary definition to be modified when JECFA definition is available).

- 11) ~~CCFAC~~ CCCF's risk management recommendations to the CAC with respect to contaminants and naturally occurring toxicants shall be guided by the principles described in the Preamble and relevant annexes of the Codex General Standard for Contaminants and Naturally Occurring Toxins in Food.
- 12) ~~CCFAC~~ CCFA/CCCF's risk management recommendations to the CAC that involve health and safety aspects of food standards shall be based on JECFA's risk assessments and other legitimate factors relevant to the health protection of consumers and to ensuring fair practices in food trade in accordance with the *Criteria for the Consideration of the Other Factors Referred to in the Second Statement of Principles*.
- 13) ~~CCFAC~~ CCFA/CCCF's risk management recommendations to the CAC shall take into account the relevant uncertainties and safety factors described by JECFA.
- 14) ~~CCFAC~~ CCFA shall endorse maximum use levels only for those additives for which 1) JECFA has established specifications of identity and purity and 2) JECFA has completed a safety assessment or has performed a quantitative risk assessment.
- 15) ~~CCFAC~~ CCCF shall endorse maximum levels only for those contaminants for which 1) JECFA has completed a safety assessment or has performed a quantitative risk assessment and 2) the level of the contaminant in food can be determined through appropriate sampling plans and analysis methods, as adopted by Codex. ~~CCFAC~~ CCCF should take into consideration the analytical capabilities of developing countries unless public health considerations require otherwise.
- 16) ~~CCFAC~~ CCFA/CCCF shall take into account differences in regional and national food consumption patterns and dietary exposure as assessed by JECFA when recommending maximum use levels for additives or maximum levels for contaminants and naturally occurring toxicants in food.
- 17) Before finalising proposals for maximum levels for contaminants and naturally occurring toxicants, ~~CCFAC~~ CCCF shall seek the scientific advice of JECFA about the validity of the analysis and sampling aspects, about the distribution of concentrations of contaminants and naturally occurring toxicants in foods and about other relevant technical and scientific aspects, including dietary exposure, as necessary to provide for a suitable scientific basis for its advice to ~~CCFAC~~ CCCF.
- 18) When establishing its standards, codes of practice, and guidelines, ~~CCFAC~~ CCFA/CCCF shall clearly state when it applies any other legitimate factors relevant to the health protection of consumers and to ensuring fair practices in food trade in accordance with the *Criteria for the Consideration of the Other Factors Referred to in the Second Statement of Principles*, in addition to JECFA's risk assessment, and specify its reasons for doing so.
- 19) ~~CCFAC~~ CCFA/CCCF's risk communication with JECFA includes prioritising substances for JECFA review with the view towards obtaining the best available risk assessment for purposes of elaborating safe conditions of use for food additives and elaborating safe maximum levels or codes of practice for contaminants and naturally occurring toxicants in food.
- 20) ~~CCFAC~~ CCFA/CCCF shall consider the following when preparing their priority list of substances for JECFA review:
 - Consumer protection from the point of view of health and prevention of unfair trade practices;
 - ~~CCFAC~~ CCFA/CCCF's Terms of Reference;
 - JECFA's Terms of Reference;
 - The Codex Alimentarius Commission's Strategic Plan , its relevant plans of work and *Criteria for the Establishment of Work Priorities*;
 - The quality, quantity, adequacy, and availability of data pertinent to performing a risk assessment, including data from developing countries;
 - The prospect of completing the work in a reasonable period of time;
 - The diversity of national legislation and any apparent impediments to international trade;
 - The impact on international trade (i.e., magnitude of the problem in international trade);
 - The needs and concerns of developing countries; and,

– Work already undertaken by other international organizations;

- 21) When referring substances to JECFA, ~~CCFAC~~ CCFA/CCCF shall provide background information and clearly explain the reasons for the request when chemicals are nominated for evaluation;
- 22) ~~CCFAC~~ CCFA/CCCF may also refer a range of risk management options, with a view toward obtaining JECFA's guidance on the attendant risks and the likely risk reductions associated with each option.
- 23) ~~CCFAC~~ CCFA/CCCF requests JECFA to review any methods and guidelines being considered by ~~CCFAC~~ CCFA/CCCF for assessing maximum use levels for additives or maximum levels for contaminants and naturally occurring toxicants. ~~CCFAC~~ CCFA/CCCF makes any such request with a view toward obtaining JECFA's guidance on the limitations, applicability, and appropriate means for implementation of a method or guideline for ~~CCFAC~~ CCFA/CCCF's work.

SECTION 4. JECFA

- 24) JECFA is primarily responsible for performing the risk assessments upon which ~~CCFAC~~ CCFA/CCCF and ultimately the CAC base their risk management decisions.
- 25) JECFA's scientific experts should be selected on the basis of their competence and independence, taking into account geographical representation to ensure that all regions are represented.
- 26) JECFA should strive to provide ~~CCFAC~~ CCFA/CCCF with science-based risk assessments that include the four components of risk assessment as defined by CAC and safety assessments that can serve as the basis for ~~CCFAC~~ CCFA/CCCF's risk-management discussions. For contaminants and naturally occurring toxicants, JECFA should determine to the extent possible the risks associated with various levels of intake. Because of the lack of appropriate information, including data in humans, however, this may be possible in only a few cases for the foreseeable future. For additives, JECFA should continue to use its safety assessment process for establishing ADIs.
- 27) JECFA should strive to provide ~~CCFAC~~ CCFA/CCCF with science-based quantitative risk assessments and safety assessments for food additives, contaminants, and naturally occurring toxicants in a transparent manner.
- 28) JECFA should provide ~~CCFAC~~ CCFA/CCCF with information on the applicability and any constraints of the risk assessment to the general population to particular sub-populations and should as far as possible identify potential risks to populations of potentially enhanced vulnerability (e.g., children, women of child-bearing age, the elderly).
- 29) JECFA should also strive to provide ~~CCFAC~~ CCFA with specifications of identity and purity essential to assessing risk associated with the use of additives.
- 30) JECFA should strive to base its risk assessments on global data, including data from developing countries. These data should include epidemiological surveillance data and exposure studies.
- 31) JECFA is responsible for evaluating exposure to additives, contaminants, and naturally occurring toxicants.
- 32) When evaluating intake of additives or contaminants and naturally occurring toxicants during its risk assessment, JECFA should take into account regional differences in food consumption patterns.
- 33) JECFA should provide to ~~CCFAC~~ CCCF its scientific views on the validity and the distribution aspects of the available data regarding contaminants and naturally occurring toxicants in foods which have been used for exposure assessments, and should give details on the magnitude of the contribution to the exposure from specific foods as may be relevant for risk management actions or options of ~~CCFAC~~ CCCF.
- 34) JECFA should communicate to ~~CCFAC~~ CCFA/CCCF the magnitude and source of uncertainties in its risk assessments. When communicating this information, JECFA should provide ~~CCFAC~~ CCFA/CCCF with a description of the methodology and procedures by which JECFA estimated any uncertainty in its risk assessment.
- 35) JECFA should communicate to ~~CCFAC~~ CCFA/CCCF the basis for all assumptions used in its risk assessments including default assumptions used to account for uncertainties.

- 36) JECFA's risk assessment output to ~~CCFAC~~ CCFA/CCCF is limited to presenting its deliberations and the conclusions of its risk assessments and safety assessments in a complete and transparent manner. JECFA's communication of its risk assessments should not include the consequences of its analyses on trade or other non-public health consequence. Should JECFA include risk assessments of alternative risk management options, JECFA should ensure that these are consistent with the Working Principles for Risk Analysis for the Application in the Framework of the Codex Alimentarius and Risk Analysis Principles applied by the ~~Codex Committee on Food Additives and Contaminants~~ Codex Committee on Food Additives and the Codex Committee on Contaminants in Foods.
- 37) When establishing the agenda for a JECFA meeting, the JECFA Secretariat work closely with ~~CCFAC~~ CCFA/CCCF to ensure that ~~CCFAC~~ CCFA/CCCF's risk management priorities are addressed in a timely manner. With respect to food additives, the JECFA Secretariat should normally give first priority to compounds that have been assigned a temporary ADI, or equivalent. Second priority should normally be given to food additives or groups of additives that have previously been evaluated and for which an ADI, or equivalent, has been estimated, and for which new information is available. Third priority should normally be given to food additives that have not been previously evaluated. With respect to contaminants and naturally occurring toxicants, the JECFA Secretariat should give priority to substances that present both a significant risk to public health and are a known or expected problem in international trade.
- 38) When establishing the agenda for a JECFA meeting, the JECFA Secretariat should give priority to substances that are known or expected problems in international trade or that present an emergency or imminent public health risk.

AMENDMENTS TO THE CCFAC POLICY FOR EXPOSURE ASSESSMENT OF CONTAMINANTS AND TOXINS IN FOODS OR FOOD GROUPS

CCFAC POLICY OF THE CODEX COMMITTEE ON CONTAMINANTS IN FOODS FOR EXPOSURE ASSESSMENT OF CONTAMINANTS AND TOXINS IN FOODS OR FOOD GROUPS

SECTION 1. INTRODUCTION

1. Maximum ~~Levels~~ ~~Limits~~ (MLs) do not need to be set for all foods that contain a contaminant or a toxin. The Preamble of the Codex General Standard for Contaminants and Toxins in Foods (GSCTF) states in Section 1.3.2 that “maximum levels (MLs) shall only be set for those foods in which the contaminant may be found in amounts that are significant for the total exposure of the consumer. They should be set in such a way that the consumer is adequately protected”. Setting standards for foods that contribute little to dietary exposure would mandate enforcement activities that do not contribute significantly to health outcomes.
2. Exposure assessment is one of the four components of risk assessment within the risk analysis framework adopted by Codex as the basis for all standard-setting processes. The estimated contribution of specific foods or food groups to the total dietary exposure to a contaminant as it relates to a quantitative health hazard endpoint (e.g. PMTDI, PTWI) provides further information needed for the setting of priorities for the risk management of specific foods/food groups. Exposure assessments must be guided by clearly articulated policies elaborated by Codex with the aim of increasing the transparency of risk management decisions.
3. The purpose of this Annex is to outline steps in contaminant data selection and analysis undertaken by JECFA when requested by ~~CCFAC~~ the Codex Committee on Contaminants in Foods (CCCF) to conduct a dietary exposure assessment.
4. The following components highlight aspects of JECFA’s exposure assessment of contaminants and toxins that contribute to ensuring transparency and consistency of science-based risk assessments. Exposure assessments of contaminants and toxins in foods are performed by JECFA at the request of ~~CCFAC~~CCCF. ~~CCFAC~~CCCF will take this information into account when considering risk management options and making recommendations regarding contaminants and toxins in foods.

SECTION 2. ESTIMATION OF TOTAL DIETARY EXPOSURE TO A CONTAMINANT OR TOXIN FROM FOODS/FOOD GROUPS

5. JECFA uses available data from member countries and from GEMS/Food Operating Program for analytical laboratories system on contaminant levels in foods and the amount of foods consumed to estimate total dietary exposure to a contaminant or toxin. This is expressed as a percentage of the tolerable intake (e.g. PTDI, PTWI, or other appropriate toxicological reference point). For a carcinogen with no clear threshold, JECFA uses available data on intake combined with data on carcinogenic potency to estimate potential population risks.
6. Median/mean contaminant levels in foods are determined from available analytical data submitted by countries and from other sources. These data are combined with information available for the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets to generate dietary exposure estimates for regions in the world. JECFA provides an estimate as to which of the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets are likely to approach or exceed the tolerable intake.
7. In some cases, available national contaminant and/or individual food consumption data may be used by JECFA to provide more accurate estimates of total dietary exposure, particularly for vulnerable groups such as children.
8. JECFA performs exposure assessments if requested by ~~CCFAC~~CCCF using the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets and, if needed, available national consumption data to estimate the impact on dietary exposure of proposed alternative maximum levels to inform ~~CCFAC~~CCCF about these risk management options.

SECTION 3. IDENTIFICATION OF FOODS/FOOD GROUPS THAT CONTRIBUTE SIGNIFICANTLY TO TOTAL DIETARY EXPOSURE OF THE CONTAMINANT OR TOXIN

9. From dietary exposure estimates JECFA identifies foods/food groups that contribute significantly to the exposure according to ~~CCFAC~~CCCCF's criteria for selecting food groups that contribute to exposure.
10. The ~~CCFAC~~CCCCF determines criteria for selecting foods/food groups that contribute significantly to total dietary exposure of a contaminant or toxin. These criteria are based upon the percentage of the tolerable intake (or similar health hazard endpoint) that is contributed by a given food/food group and the number of geographic regions (as defined by the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets) for which dietary exposures exceed that percentage.
11. The criteria are as follows:
 - a) Foods or food groups for which exposure to the contaminant or toxin contributes approximately 10%¹¹ or more of the tolerable intake (or similar health hazard endpoint) in one of the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets;
 - or,
 - b) Foods or food groups for which exposure to the contaminant or toxin contributes approximately 5%¹ or more of the tolerable intake (or similar health hazard endpoint) in two or more of the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets;
 - or,
 - c) Foods or food groups that may have a significant impact on exposure for specific groups of consumers, although exposure may not exceed 5% of the tolerable intake (or similar health hazard endpoint) in any of the ~~GEMS/Food Regional diets~~GEMS/Food Consumption Cluster Diets. These would be considered on a case-by-case basis.

SECTION 4. GENERATION OF DISTRIBUTION CURVES FOR CONCENTRATIONS OF THE CONTAMINANT IN SPECIFIC FOODS/FOOD GROUPS (CONCURRENT WITH SECTION 2, OR SUBSEQUENT STEP)

12. If requested by ~~CCFAC~~CCCCF, JECFA uses available analytical data on contaminant or toxin levels in foods/food groups identified as significant contributors to dietary exposure to generate distribution curves of contaminant concentrations in individual foods. ~~CCFAC~~CCCCF will take this information into account when considering risk management options and, if appropriate, for proposing the lowest achievable levels for contaminants/toxins in food on a global basis.
13. Ideally, individual data from composite samples or aggregated analytical data would be used by JECFA to construct the distribution curves. When such data are not available, aggregated data would be used (for example mean and geometric standard deviation). However, methods to construct distribution curves using aggregated data would need to be validated by JECFA.
14. In presenting the distribution curves to ~~CCFAC~~CCCCF, JECFA should, to the extent possible, provide a comprehensive overview of the ranges of contamination of foods (i.e., both the maximum and outlier values) and of the proportion of foods/food groups that contain contaminants/toxins at those levels.

SECTION 5. ASSESSMENT OF THE IMPACT OF AGRICULTURAL AND PRODUCTION PRACTICES ON CONTAMINANT LEVELS IN FOODS/FOOD GROUPS (CONCURRENT WITH SECTION 2, OR SUBSEQUENT STEP)

15. If requested by ~~CCFAC~~CCCCF, JECFA assesses the potential impact of different agricultural and production practices on contaminant levels in foods to the extent that scientific data are available to support such assessments. ~~CCFAC~~CCCCF takes this information into account when considering risk management options and for proposing Codes of Practice.

¹¹ Rounded to the nearest 1/10th of a percent.

16. Taking this information into account, ~~CCFACCCCF~~ proposes risk management decisions. To refine them, ~~CCFACCCCF~~ may request JECFA to undertake a second assessment to consider specific exposure scenarios based on proposed risk management options. The methodology for assessing potential contaminant exposure in relation to proposed risk management options needs to be further developed by JECFA.

AMENDMENTS TO SECTION IV OF THE PROCEDURAL MANUAL**FAO/WHO COORDINATING COMMITTEE FOR EUROPE (CX-706)*****Membership:***

Membership of the This Committee is open to all Member Nations Governments and Associate Members of FAO and/or WHO which are members of the Codex Alimentarius Commission, within the geographic location area of Europe, including Israel, Turkey and the Russian Federation and its Chairperson is, ex officio, the Coordinator for Europe.

Terms of reference: [no change]

CODEX COMMITTEE ON FOOD ADDITIVES (CX-711)***Terms of reference:***

- (a) to establish or endorse acceptable ~~permitted~~ maximum levels for individual food additives;
- (b)-(f): [no change]

CODEX COMMITTEE ON CONTAMINANTS IN FOODS (CX-735)***Terms of reference:***

- (a) to establish or endorse permitted maximum levels, ~~or~~ and where necessary revise existing guideline levels, for contaminants and naturally occurring toxicants in food and feed;
- (b) to prepare priority lists of contaminants and naturally occurring toxicants for risk assessment by the Joint FAO/WHO Expert Committee on Food Additives;
- (c) to consider and elaborate methods of analysis and sampling for the determination of contaminants and naturally occurring toxicants in food and feed;
- (d) to consider and elaborate standards or codes of practice for related subjects; and
- (e) to consider other matters assigned to it by the Commission in relation to contaminants and naturally occurring toxicants in food and feed.

APPENDIX IV

**LIST OF STANDARDS AND RELATED TEXTS ADOPTED BY THE THIRTIETH SESSION OF
THE CODEX ALIMENTARIUS COMMISSION**

Part 1 – Standards and Related Texts Adopted at Step 8

Standard and Related Text	Reference	Status
Maximum Levels for Tin in Canned Foods (other than beverages) and in Canned Beverages	ALINORM 07/30/41 Appendix IX	Adopted
Food Additive Provisions of the General Standard for Food Additives (GSFA)	ALINORM 07/30/12 Rev. Appendix VII	Adopted
Standard for Table Grapes	ALINORM 07/30/35 Appendix IV and V	Adopted
Code of Hygienic Practice for Eggs and Egg Products	ALINORM 07/30/13 Appendix II	Adopted
Guidelines on the Application of General Principles of Food Hygiene to the Control of <i>Listeria monocytogenes</i> in Ready-to-Eat Foods	ALINORM 07/30/13 Appendix III	Adopted with amendments (see para. 51)
Principles and Guidelines for the Conduct of Microbiological Risk Management	ALINORM 07/30/13 Appendix IV	Adopted
Amendment to the Guidelines for the Production, Processing, Labelling and Marketing of Organically Produced Foods: Annex 2 – Permitted Substances: Table 3	ALINORM 07/30/22 Appendix II	Adopted
Standard for Fat Spreads and Blended Spreads	ALINORM 07/30/17 Appendix II	Adopted with amendments (see para. 55)
Regional Standard for Canned Humus with Tehena	ALINORM 07/30/40 Appendix II	Adopted with amendments (see para. 61)
Regional Standard for Canned Foul Medames	ALINORM 07/30/40 Appendix III	Adopted with amendments (see para. 61)
Regional Standard for Tehena	ALINORM 07/30/40 Appendix IV	Adopted with amendments (see para. 61)
Revised Standard for Infant Formula and Formula for Special Medical Purposes Intended for Infants	ALINORM 07/30/26 Appendix II	Adopted
Standard for Pickled Fruits and Vegetables	ALINORM 07/30/27 Appendix II	Adopted
Standard for Processed Tomato Concentrates	ALINORM 07/30/27 Appendix III	Adopted
Standard for Preserved Tomatoes	ALINORM 07/30/27 Appendix IV	Adopted
Standard for Certain Canned Citrus Fruits	ALINORM 07/30/27 Appendix V	Adopted
Maximum Residue Limits for Pesticides	ALINORM 07/30/24 Appendix II	Adopted with amendments (see para. 69)
Revised Standard for Cheddar (C-1)	ALINORM 06/29/11 Appendix VI	Adopted with amendments from endorsement process

Standard and Related Text	Reference	Status
Revised Standard for Danbo (C-3)	ALINORM 06/29/11 Appendix VII	of food additive provisions (see paras. 74 and 77-79)
Revised Standard for Edam (C-4)	ALINORM 06/29/11 Appendix IX	
Revised Standard for Gouda (C-5)	ALINORM 06/29/11 Appendix X	
Revised Standard for Havarti (C-6)	ALINORM 06/29/11 Appendix XI	
Revised Standard for Samsø (C-7)	ALINORM 06/29/11 Appendix XII	
Revised Standard for Emmental (C-9)	ALINORM 06/29/11 Appendix XIII	
Revised Standard for Tilsiter (C-11)	ALINORM 06/29/11 Appendix XIV	
Revised Standard for Saint-Paulin (C-13)	ALINORM 06/29/11 Appendix XV	
Revised Standard for Provolone (C-15)	ALINORM 06/29/11 Appendix XVI	
Revised Standard for Cottage Cheese (C-16)	ALINORM 06/29/11 Appendix XVII	
Revised Standard for Coulommiers (C-18)	ALINORM 06/29/11 Appendix XVIII	
Revised Standard for Cream Cheese (C-31)	ALINORM 06/29/11 Appendix XIX	
Revised Standard for Camembert (C-33)	ALINORM 06/29/11 Appendix XX	
Revised Standard for Brie (C-34)	ALINORM 06/29/11 Appendix XXI	
Standard for Mozzarella	ALINORM 06/29/11 Appendix XXII	

Part 2 – Standards and Related Texts Adopted at Step 5/8 (with omission of Step 6 and 7)

Standards and Related Text	Reference	Status
Code of Practice for the Prevention and Reduction of Ochratoxin A Contamination in Wine (N05-2006)	ALINORM 07/30/12 Appendix VIII	Adopted
Food Additive provisions of the General Standard for Food Additives (GSFA)	ALINORM 07/30/12 Rev. Appendix VII	Adopted
Amendments to the International Numbering System for Food Additives	ALINORM 07/30/12 Rev. Appendix XIII	Adopted
Specifications for the Identity and Purity of Food Additives arising from the 65 th JECFA meeting	ALINORM 07/30/12 Rev. Appendix XIV Part 1	Adopted

Standards and Related Text	Reference	Status
Code of Practice for Fish and Fishery Products (Quick Frozen Coated Products, Salted Fish and relevant Definitions)	ALINORM 07/30/18 Appendix II	Adopted
Sections 2.1.2 - Maturity Requirements and 3.1 - Minimum Bunch Weight (Standard for Table Grapes)	ALINORM 07/30/35 Appendix V	Adopted
Guidelines for Design, Production, Issuance and Use of Generic Official Certificates (Revision of the <i>Guidelines for Generic Official Certificate Formats and the Design, Production, Issuance and Use of Certificates</i> (N05-2005))	ALINORM 07/30/30 Appendix II	Adopted with amendments (see paras 52-54)
Working Principles for Risk Analysis for Food Safety for Application by Governments	ALINORM 07/30/33 Appendix VIII	Adopted
Maximum Residue Limits for Pesticides	ALINORM 07/30/24 Appendix III	Adopted with amendments (see para. 69)

Guidelines for Design, Production, Issuance and Use of Generic Official Certificates

Part 3 – Standards and Related Texts Adopted at Step 5 of the Accelerated Procedure

Standard and Related Text	Reference	Status
Amendment to the Standard for Canned Sardines and Sardine-Type Products	ALINORM 07/30/18 Appendix III	Adopted

Part 4 – Other Standards and Related Texts Adopted

Standard and Related Text	Reference	Status
Methods of Analysis in Codex Standards at different steps	ALINORM 07/30/23 Appendix III	Adopted
Amendments to Schedule I of the General Standard for Contaminants and Toxins in Foods	ALINORM 07/30/41, para. 46	Adopted (see para. 191)
Amendment to the Annex to Table 3 of the General Standard for Food Additives	ALINORM 07/30/12 Rev., para. 71	Adopted (see para. 203)
Amendment to the General Standard for Food Additives - Entries in Food Category 02.1.1	ALINORM 07/30/12 Rev., para. 167	Adopted (see para. 204)

APPENDIX V

**LIST OF DRAFT STANDARDS AND RELATED TEXTS ADOPTED AT STEP 5 BY THE
THIRTIETH SESSION OF THE CODEX ALIMENTARIUS COMMISSION**

Standards and Related Texts	Reference	Status
Draft Guidelines for the Use of Flavourings (N03-2006) (with the exception of Section 4 and Annexes A and B)	ALINORM 07/30/12 Rev. para. 123 and Appendix XI	Adopted
Draft Regional Standard for Gochujang (N03-2004)	ALINORM 07/30/15, para. 42 and Appendix II	Adopted
Draft Regional Standard for Ginseng Product (N01-2004)	ALINORM 07/30/15, para. 68 and Appendix III	Adopted
Draft Code of Practice for Fish and Fishery Products (Live and Raw Bivalve Molluscs, Lobsters and Crabs and relevant Definitions)	ALINORM 07/30/18, para. 92 and Appendix IV	Adopted
Draft Standard for Live and Raw Bivalve Molluscs	ALINORM 07/30/18, para. 111, Appendix V	Adopted
Draft Amendment to the Guidelines for the Production, Processing, Labelling and Marketing of Organically Produced Foods: Annex 1 (inclusion of Ethylene) (N10-2006)	ALINORM 07/30/22, para. 96 and Appendix IV	Adopted
Draft Amendment to the General Standard for the Labelling of Prepackaged Foods: Quantitative Declaration of Ingredients	ALINORM 07/30/22, para. 133 and Appendix V	Adopted
Draft Definition of Advertising in Relation to Nutrition and Health Claims (N11-2006)	ALINORM 07/30/22, para. 140 and Appendix VI	Adopted
Draft Maximum Residue Limits for Pesticides	ALINORM 07/30/24, paras 44-136 and Appendix IV	Adopted
Advisory Lists of Nutrient Compounds for Use in Foods for Special Dietary Uses Intended for Infants and Young Children	ALINORM 07/30/26, para. 130 and Appendix V	Adopted
Draft Standard for Jams, Jellies and Marmalades	ALINORM 07/30/27, para. 146 and Appendix VI	Adopted
Draft Standard for Certain Canned Vegetables (general provisions)	ALINORM 07/30/27, para. 114 and Appendix VII	Adopted
Draft Standard for Bitter Cassava (N03-2005)	ALINORM 07/30/35, para. 82 and Appendix VI	Adopted
Draft Guidelines for the Inspection and Certification of Fresh Fruits and Vegetables for Conformity to Quality Standards	ALINORM 07/30/35, para. 92 and Appendix VII	Adopted
Draft Maximum Levels for 3-MCPD in Liquid Condiments containing Acid-Hydrolyzed Vegetable Proteins (excluding Naturally Fermented Soy Sauce) (N08-2004)	ALINORM 07/30/41, para. 88 and Appendix X	Adopted
Draft Code of Practice for the Reduction of 3-Monochloropropane-1,2-diol (3-MCPD) during the Production of Acid-Hydrolyzed Vegetable Protein (Acid-HVPs) and Products that Contain Acid-HVPs (N09-2005)	ALINORM 07/30/41, para. 93 and Appendix XI	Adopted

APPENDIX VI

**LIST OF STANDARDS AND RELATED TEXTS REVOKED BY THE THIRTIETH SESSION OF
THE CODEX ALIMENTARIUS COMMISSION**

Standard and Related Texts	Reference
Codex Specifications for Identity and Purity of Food Additives	ALINORM 07/30/12, para. 157 and Appendix XIV Part 2
Certain existing Codex MRLs for pesticides	ALINORM 07/30/24, paras 44-136 and Appendix V
Maximum levels for cadmium (CODEX STAN 248-2005)	ALINORM 07/30/41, para. 46
Standard for Canned Grapefruit (CODEX STAN 15-1981)	ALINORM 07/30/27, para. 89
Standard for Canned Mandarin Oranges (CODEX STAN 68-1981)	
Standard for Margarine (CODEX STAN 32-1981)	ALINORM 07/30/17, para. 63
Standard for Minarine (CODEX STAN 135-1981)	
Code of Practice for Frozen Battered and/or Breaded Fishery Products (CAC/RCP 35-1985)	ALINORM 07/30/18, para. 91
Code of Practice for Salted Fish (CAC/RCP 26-1979)	

APPENDIX VII

**LIST OF DRAFT STANDARDS AND RELATED TEXTS APPROVED AS NEW WORK BY THE
THIRTIETH SESSION OF THE CODEX ALIMENTARIUS COMMISSION**

Responsible Committee	Standard and Related Texts	Job Code
CCFFP	Revision of the Procedure for the Inclusion of Additional Species in Standards for Fish and Fishery Products	Procedure
CCFFP	Amendment to the Standard for Quick Frozen Fish Sticks, Fish Portions and Fish Fillets – Breaded or in Batter (Nitrogen Factors)	N01-2007
CCFFP	Standard for Fish Sauce	N02-2007
CCFFP	Standard for Fresh/Live and Frozen Abalone (<i>Haliotis spp</i>)	N03-2007
CCPFV	Sampling Plan including Methodological Provisions for Controlling Minimum Drained Weight of Canned Fruits and Vegetables in Packing Media	N04-2007
CCNFSDU	Establishment and Application of Risk Analysis Principles by the Committee on Nutrition and Foods for Special Dietary Uses	Procedure
CCASIA	Regional Standard for Chili Sauce	N05-2007
CCASIA	Regional Standard for Edible Sago Flour	N06-2007
TFFBT	Annex to the Guideline for the Conduct of Food Safety Assessment of Foods Derived from Recombinant-DNA Plants on Low-level Presence of Recombinant-DNA Plant Material	N07-2007
CCFH	Guidelines for Control of <i>Campylobacter</i> and <i>Salmonella</i> spp. in Chicken Meat	N08-2007
CCFO	Amendment to the Standard for Named Vegetable Oils: Palm Kernel Olein and Palm Kernel Stearin	N09-2007
CCCF	Code of Practice for the Prevention and Reduction of Aflatoxin Contamination in Dried Figs	N10-2007
CCFA	Revision of the Food Category System (FCS) of the Codex General Standard for Food Additives	N11-2007
CCPR	Priority List of Pesticides for Evaluation by JMPR (New Pesticides and Pesticides under Periodic Review)	Ongoing
CCNMW	Amendments to Section 3.2 "Health-Related Limits for Certain Substances" of the Standard for Natural Mineral Waters (CODEX STAN 108-1981)	N12-2007

APPENDIX VIII

LIST OF WORK DISCONTINUED BY THE THIRTIETH SESSION OF THE CODEX
ALIMENTARIUS COMMISSION

Responsible Committee	Standard and Related Texts	Reference
CCFA	Draft and Proposed Draft Food Additive Provisions of the GSFA	ALINORM 07/30/12, para. 107 and Appendix VIII
CCFL	Proposed Draft Amendment to the Guidelines for the Production, Processing, Labelling and Marketing of Organically Produced Foods: Annex 2 – Permitted Substances : Table 1 (Natural Sodium Nitrate)	ALINORM 07/30/22, para. 92
CCPR	Draft and Proposed Draft MRLs for Pesticides	ALINORM 07/30/24, paras 44-136 and Appendix IX

APPENDIX IX

**CODEX ALIMENTARIUS COMMISSION
STRATEGIC PLAN 2008-2013****PART 1****STRATEGIC VISION STATEMENT**

The Codex Alimentarius Commission envisages a world afforded the highest attainable levels of consumer protection including food safety and quality. To this end, the Commission will develop internationally agreed standards and related texts for use in domestic regulation and international trade in food that are based on scientific principles and fulfil the objectives of consumer health protection and fair practices in food trade.

INTRODUCTION

1. This document sets out the strategic plan for the Codex Alimentarius Commission (CAC), stating strategic goals of the Commission (Part 1) and incorporating a list of programme areas and planned activities with a clearly defined timetable (Part 2). The strategic vision and goals for the CAC underpin the high priority attached to food safety by its parent organizations, the Food and Agriculture Organization of the United Nations (FAO) and the World Health Organization (WHO). *The Strategic Framework for FAO: 2000-2015* accords high priority to promoting policy and regulatory frameworks for food at the international and national levels. Several resolutions adopted by the World Health Assembly recognized the need to highlight health considerations in international food trade and acknowledged the importance of the CAC for ensuring the highest levels of consumer health protection. These resolutions and related documents¹ urged WHO to work towards integrating food safety as one of its essential public health functions with the goal of developing sustainable, integrated food safety systems for the reduction of health risks along the entire food chain. It is understood that Codex, when elaborating standards, guidelines and recommendations, gives full consideration to those resolutions and decisions from WHO and FAO that are relevant within the framework of the Codex mandate. The fundamental mandate of the CAC is to develop international standards, guidelines and other recommendations for protecting the health of consumers and ensuring fair practices in the food trade.
2. The CAC has always operated in an environment of change and technological advancement. The growth in world food trade, advances in modern communication and increasing mobility of populations are all contributing to elevating the profile and significance of food safety and regulation. There is growing international concern related to a perceived emergence of or increase in food-borne diseases. Consumers around the world are seeking ever-greater assurances about the safety and quality of foods they eat. In its endeavour to promote food safety and quality, the CAC needs to ensure more effective participation and involvement of all members in setting globally relevant standards and to consider opportunities for strengthening partnerships with all stakeholders, in particular consumers and their representative organizations, at the global and national levels. It is also likely that developing countries will account for an increasing proportion of global food and agricultural trade. CAC, FAO and WHO are striving to respond to the new challenges and keep abreast of most recent developments.²
3. The recognition and status that Codex standards, guidelines and other recommendations acquired under the World Trade Organization (WTO) Agreement on the Application of Sanitary and Phytosanitary Measures has presented challenges and brought responsibilities, including the need to ensure that Codex standards and related texts are based on scientific principles and meet the needs and mandate of the organization. The WTO Agreement on Technical Barriers to Trade is also of great relevance, given the significance of the provisions pertaining to product description, labelling,

¹ WHO Global Strategy for Food Safety (WHO, 2002).

² Joint FAO/WHO Evaluation of the Codex Alimentarius and Other FAO and WHO Food Standards Work.

packaging and quality descriptors for consumer information and fair practices in trade. The CAC has an important role in providing for essential composition and quality requirements that are not more trade-restrictive than necessary. The CAC needs to maintain its pre-eminent status as the internationally recognized body for food standard-setting and to call for the use of its standards to the widest extent possible by all members as a basis for domestic regulation and international trade. This will help members to be more aware of the importance of the international harmonization of food safety and quality standards, as well as the enhancement of food control systems for ensuring food safety and quality.

DECISION-MAKING BASED ON SCIENTIFIC EVIDENCE

4. The CAC, as a risk management body, does not undertake scientific evaluations *per se* but relies on the opinions of scientific expert bodies convened by FAO and WHO on specific issues. These expert bodies such as the Joint FAO/WHO Expert Committee on Food Additives (JECFA), the Joint FAO/WHO Meetings on Pesticide Residues (JMPR) and the Joint FAO/WHO Expert Meetings on Microbiological Risk Assessment (JEMRA) and other *ad hoc* expert consultations are functionally separate from the CAC and its subsidiary bodies and do not directly fall within the scope of the present Strategic Plan. The mandates, functions, composition and agendas of these bodies are established by FAO and WHO. The independence of the expert bodies is critical to the objectivity of their opinions, and meetings of these bodies should interact with the CAC in accordance with the Working Principle for Risk Analysis for Application in the Framework of the Codex Alimentarius. There is considerable synergy between the scientific bodies of FAO and WHO and the intergovernmental bodies of the CAC in order to take decisions based on scientific evidence.

STRATEGIC GOALS AND SHARED RESPONSIBILITIES

5. To enable the overall achievement of the strategic vision, the CAC must take action jointly with its parent organizations and its members. The Commission urges FAO and WHO to mobilise sufficient resources to allow the CAC to fulfil its mandate. Their other key roles are to provide scientific advice requested by the CAC and to offer technical assistance to developing members so that they can effectively participate in the standard-setting process and build capacity for the development of sound food control systems. The Commission fully recognises the efforts of the members of the CAC, especially those which provide significant financial and other support to the work of the CAC as host governments of subsidiary bodies or as contributors to extra-budgetary programmes of FAO and WHO. In close cooperation with the partners above, the Commission will focus on the following goals to achieve its strategic vision.

Goal 1: Promoting Sound Regulatory Frameworks

6. An effective food control system is critical in enabling all countries to ensure the safety of their foods entering international trade and to ensure that imported foods conform to national requirements. International harmonization based on Codex standards, guidelines and recommendations is essential to promoting a global approach to consumer health protection, including systems for the reduction of food-borne risks, and minimizing the negative effects of technical regulations on international trade. For this purpose, the CAC will provide essential guidance for its members through the continued development of international standards and guidelines relating to food safety and hygiene, nutrition, labelling, and import/export inspection and certification and quality of food stuff. This will require sustained commitment and effort in the following key directions:
 - The CAC will develop international standards, guidelines, and recommendations based on scientific principles for the reduction of health risks along the entire food chain, including feed when appropriate. In strengthening the strategic focus of the CAC in the development of standards and related texts based on risk and performance for broad application across a range of commodities, the CAC must give priority to establishing a coherent and integrated set of food standards covering the entire food chain. Such an approach can serve as a model for the members of the CAC to pursue food regulatory systems that provide consumers with safe food and ensure fair practices in the food trade;
 - Codex standards and related texts for food safety and quality, including labelling aspects, should be carefully prepared to reflect global variations. Codex standards for food quality

should focus on essential characteristics of products to ensure that they are not overly prescriptive and that the standards are not more trade restrictive than necessary; and

- The CAC, when elaborating and deciding upon Codex standards and related texts should take into consideration the technical and economic implications for all members as well as the special needs of developing countries including infrastructure, resources and technical and legal capabilities, Codex standards and related texts should not have the effect of creating unnecessary, unjustified or discriminatory obstacles to the exports of developing countries.

7. In many countries, effective food control is undermined by the existence of fragmented legislation, multiple jurisdictions and weaknesses in surveillance, monitoring and enforcement. Sound national food control and regulatory systems are essential to ensuring the health and safety of the domestic population as well as ensuring the safety and quality of foods entering international trade. The FAO and WHO have made significant advances in promoting sound regulatory frameworks at the national level. The Commission, while encouraging members to use relevant Codex standards, strongly encourages FAO and WHO to continue to promote national regulatory systems that are based on international principles and guidelines and address all components of the food chain. The development of sound food control and regulatory infrastructure including human resources is particularly important for developing countries as they seek to achieve higher levels of food safety and nutrition and will require high-level political and policy commitment.³ Successful negotiation of bilateral mutual recognition and equivalence of food control systems also depends on the abilities of countries to assure each other of the integrity and international conformity of their regulatory systems.

Goal 2: Promoting Widest and Consistent Application of Scientific Principles and Risk Analysis

8. The scientific basis of decision-making by the CAC is spelled out in the Statements of Principle on the Role of Science in the Codex Decision-Making process and the Extent to Which Other Factors are Taken into Account and in the Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius⁴. The CAC will ensure their consistent application by relevant Codex subsidiary bodies, in order to maintain its focus on this Goal. Risk analysis as it applies to food safety across the food chain is an internationally accepted discipline and will require ongoing and sustained inputs from the CAC, its parent organisations and national governments to promote its understanding and application at the international and national levels.
9. In recent years the scope of scientific advice sought by the CAC from the parent organisations increased considerably and went beyond chemical and microbiological hazards. FAO and WHO responded to these requests through several FAO/WHO *ad hoc* consultations on topics such as foods derived from genetically modified organisms and antimicrobial resistance. The Commission requests FAO and WHO to continue to promote the understanding of risk analysis and to continue to explore new areas of work, such as nutritional risk assessment, so as to provide the scientific advice relevant to CAC activities for standard setting.
10. The timely availability of scientific advice is a prerequisite for the CAC to fulfil its mandate. The Commission will continue to encourage FAO and WHO to make sufficient resources available to ensure that the scientific advice to the CAC can be provided in a timely and sustainable way. To make more efficient and effective use of the FAO/WHO expert bodies and *ad hoc* consultations, particularly given the rapidly expanding scope of scientific advice requested from FAO and WHO, the CAC will continue to strengthen the interaction between the risk managers (relevant Codex subsidiary bodies) and the risk assessors (FAO/WHO expert bodies and *ad hoc* expert consultations). The Commission has agreed to recommend to FAO and WHO a set of criteria for the prioritization of requests from the CAC for scientific advice and will review the usefulness of this approach. The CAC, in close cooperation with its parent organizations, will enhance its capacity to respond efficiently to emerging food borne risks by strengthening its work management capabilities (see Goal 3).

³ Report of the Conference on International Food Trade beyond 2000: Science based Decisions, Harmonization, Equivalence and Mutual Recognition, Melbourne, Australia, 11-15 October, 1999, Appendix 1.

⁴ Codex Alimentarius Procedural Manual.

11. The CAC has the goal of elaborating standards that cover the needs of its entire membership to ensure these standards are applicable globally. A constraint to this goal is the persistent lack of relevant data from all major parts of the world. The CAC will continue to encourage countries from both the developed and developing worlds to submit relevant data to the CAC and the parent organizations. The Commission recommends that FAO and WHO build on the achievements already accomplished⁵ and take meaningful steps to ensure that scientific advice is provided more quickly, with even higher quality, that more requests are addressed, and that the process is conducted with enhanced transparency. The Commission particularly encourages FAO and WHO to explore new approaches to enhance participation of experts and the use of data from developing countries in the elaboration of scientific advice. Where relevant data are not available from developing countries, the CAC encourages FAO and WHO to assist the developing countries in generating such data.

Goal 3: Strengthening Codex Work-Management Capabilities

12. Attention to food safety and global food trade has continued to increase among countries and among the international governmental and non-governmental organizations concerned with these matters. More expeditious and efficient work by the CAC is necessary to provide members and international organizations with the standards, guidelines, and recommendations that they need.
13. The CAC has already made several important advances towards achieving more efficient work-management procedures, such as strengthening the role of the Executive Committee as a strategic and standards-management body, holding annual Commission sessions, and instituting more effective use of information technology. But the CAC must take additional steps to keep pace with international developments by better managing its work so that it addresses high priority issues in a timely manner and that standards development work is completed within specified time frames.
14. The implementation of new Codex work-management procedures⁶ must make the CAC more effective and efficient, while maintaining the valuable reputation that the CAC has earned as an open, fair, transparent, and rules-based body. Key features of continuing enhancements include⁷:
 - Enhancing the capabilities of the Executive Committee with respect to strategic oversight, direction, and cross coordination of the work programmes of all subsidiary bodies through recommendations to the Commission;
 - Ensuring that the Commission and its subsidiary bodies make decisions about work prioritization using criteria that enable effective decision making, taking into consideration the need to initiate new work and to revise existing standards;
 - Ensuring that new work and standard-revision work is completed within defined time frames. Work progress is monitored by the Executive Committee, and in the event that work exceeds specified time frames, the Executive Committee recommends to the Commission that corrective actions be taken as necessary;
 - Exploring mechanisms for progressing the work of subsidiary bodies in between sessions, while maintaining transparency and inclusiveness;
 - Promoting consensus-based decision-making; and
 - Strengthening the Secretariat of the CAC to ensure effective operation and work management of the Commission and its subsidiary bodies and to maintain effective communication with the Codex Contact Points.

⁵ FAO/WHO Consultative Process for the Provision of Scientific Advice to Codex and Member Countries.

⁶ As suggested by both the Report of the Evaluation of the Codex Alimentarius and Other FAO and WHO Food Standards Work and the Report on the Review of Codex Committee Structure and Mandates of Codex Committees and Task Forces and as endorsed by the Commission.

⁷ Key features are not listed in priority order.

Goal 4: Promoting Cooperation between Codex and Relevant International Organizations

15. The CAC must work closely on matters of common interest with other relevant international organizations, including those whose work has indirect but significant implications for food-standard issues. Monitoring by the CAC of activities of other organizations that are relevant to food standards, and coordination with them, where appropriate and consistent with Codex procedures, is necessary to achieve complementarity, avoid duplication and prevent development of contradictory standards or guidelines. Such collaboration is also critical to the development of health-protection and food-trade measures that address the food chain from farm to table in a coherent and seamless manner.
16. The WTO recognizes the CAC as the pre-eminent international body for establishing food safety standards. The Commission must, therefore, play a leadership role in establishing international food standards for protecting the health of consumers and ensuring fair practices in food trade, while taking due account of international regulatory initiatives of international governmental and non-governmental organizations. The CAC also has a responsibility to provide its technical input and expertise towards the building of international consensus on food standards and regulatory policy matters. Establishment or promotion of cooperation, between the CAC and other relevant international intergovernmental organizations, in particular, OIE and IPPC, should be considered, where appropriate, to ensure effective collaboration and coordination, and that such cooperation should be in line with the Guidelines on Cooperation between Codex Alimentarius Commission and International Intergovernmental Organizations in the Elaboration of Standards and Related Texts.

Goal 5: Promoting Maximum and Effective Participation of Members

17. Full participation by all Codex Members and other interested parties in the work of the CAC is now more important than ever. The participation of all members and relevant intergovernmental and international non-governmental organizations is critical to sound decision-making and ensuring that Codex standards and related texts take account of the full range of interests and viewpoints. Since the early 1990s there has been a significant increase in the membership of the CAC with developing countries now constituting a significant proportion of total membership. The Commission welcomes some initiatives undertaken so far to mitigate the financial and human resource constraints hitherto hampering the effective participation of developing countries and countries with economies in transition in the activities of the CAC. Such initiatives include the establishment of the Joint FAO/WHO Project and Trust Fund for Enhanced Participation in Codex, and the development of training manuals and other Codex related capacity building tools. Capacity building programmes under FAO and WHO also have a bearing on strengthening these countries' participation in Codex activities. The Trust Fund and other FAO and WHO programmes are efforts aimed at enabling the members to further gain experience in the Codex process. The Commission strongly urges beneficiary members to take these opportunities offered and create sustainability towards more effective participation, by making firm commitments to adequate allocation of national resources towards Codex work.
18. There is a continuing need for FAO and WHO to implement capacity building programmes in a coherent manner, especially in developing countries and countries with economies in transition, aimed at strengthening national administrative and consultative structures on Codex (e.g. Codex Contact Point, National Codex Committee) and enhancing technical expertise required for effective participation in international standards development. The CAC will play an advisory role in facilitating the efforts made by FAO and WHO so that those efforts address the needs of the CAC and its members.
19. In addition to actions to promote participation of member countries, the CAC will continue to enhance inclusiveness and transparency of the Codex process by furthering its efforts to encourage the participation of consumers and public interest groups in its processes at the international level and encourage governments to take action at the national level. The CAC will take advantage of any information technological developments for advancement of inclusiveness and transparency in the Codex process.

PART 2

PROGRAMME AREAS AND PLANNED ACTIVITIES 2008-2013

Goal 1: Promoting Sound Regulatory Frameworks**1.1 : Review and develop Codex standards and related texts for food safety**

Description: Review and develop Codex standards and related texts for food safety, taking into account scientific and technological developments, to ensure that they: emphasize a horizontal approach; employ an approach to food safety that is based on risk and that addresses the entire food chain; and reflect global variations so as to avoid being more trade restrictive than necessary, while respecting the basic objectives of the CAC, taking into consideration the technical and economic implications for all members as well as the special needs of developing countries including infrastructure, resources and technical and legal capabilities.

Timeline : Continuing

Responsible parties: CCFH, CCFA, CCCF, CCPR, CCRVDF, CCNFSDU, relevant Task Forces and Commodity Committees

1.2 : Review and develop Codex standards and related texts for food quality

Description: Review and develop Codex standards and related texts for food quality, taking into account scientific and technological developments, to ensure that they are generic in nature and whilst maintaining inclusiveness, reflect global variations and focus on essential characteristics so as to avoid being overly prescriptive and not more trade restrictive than necessary, while respecting the basic objectives of the CAC, taking into consideration the technical and economic implications for all members as well as the special needs of developing countries including infrastructure, resources and technical and legal capabilities.

Timeline : Continuing

Responsible parties: Relevant Task Forces, Commodity Committees and FAO/WHO Coordinating Committees

1.3 : Review and develop Codex standards and related texts for food labelling and nutrition

Description: Review and develop Codex standards and related texts for food labelling and nutrition, taking into account scientific and technological developments and the WHO Global Strategy on Diet, Physical Activity and Health, to ensure that they: emphasize a horizontal approach and the need to maintain inclusiveness, and address food labelling and nutrition so as to avoid being overly prescriptive and not more trade restrictive than necessary, while respecting the basic objectives of the CAC, taking into consideration the technical and economic implications for all members as well as the special needs of developing countries including infrastructure, resources and technical and legal capabilities.

Timeline: Continuing

Responsible parties: CCFL, CCNFSDU

1.4 : Review and develop Codex standards and related texts for food inspection and certification, and methods of sampling and analysis
Description: Review and develop Codex standards and related texts for food inspection and certification as well as methods of sampling, including guidance on equivalence, mutual recognition and traceability / product tracing, taking into account scientific and technological developments, to ensure that they: emphasize a horizontal approach and the need to maintain inclusiveness, and reflect global variations so as to avoid being overly prescriptive and not more trade restrictive than necessary, while respecting the basic objectives of the CAC, taking into consideration the technical and economic implications for all members as well as the special needs of developing countries including infrastructure, resources and technical and legal capabilities.
Timeline: Continuing
Responsible parties: CCMAS, CCFICS
1.5 : Develop guidance for safe and prudent non-human antimicrobial usage for containment of resistance
Description: Develop guidance within the remit of the Codex mandate for safe and prudent antimicrobial usage for containment of resistance in food production which focuses on public health, is based on sound science and follows risk analysis principles, and takes into account the work of other international organisations.
Timeline: Completion by 2011
Responsible parties: Existing relevant Codex Committees, <i>ad hoc</i> Intergovernmental Task Force on Antimicrobial Resistance
1.6 : Explore innovative risk management frameworks
Description: Explore innovative risk management frameworks in establishing MRLs of veterinary drugs and pesticides and share the results of new approaches among Codex Committees.
Timeline: Completion by 2009
Responsible parties: CCRVDF, CCPR
1.7 : Encourage FAO/WHO to expand capacity building programmes
Description: Encourage FAO/WHO to strengthen their programmes to enhance food control infrastructures and to provide technical assistance including assistance on generating data to countries in need to promote application or use of Codex standards and related texts at the national and regional level. Request FAO/WHO to report to the Commission on the implementation status of their capacity building activities.
Timeline: Continuing
Responsible parties: CAC, CCEXEC and FAO/WHO Coordinating Committees
1.8 : Publish and disseminate the Codex Alimentarius
Description: Ensure timely publication and availability of Codex standards, guidelines and recommendations to all interested parties through the Internet and other appropriate means.
Timeline: Continuing
Responsible parties: Codex Secretariat, Codex Contact Points

Goal 2: Promoting Widest and Consistent Application of Scientific Principles and Risk Analysis

2.1 Review the consistency of risk analysis principles elaborated by the relevant Codex Committees

Description: Review the risk analysis principles elaborated by the relevant Codex Committees for consistency with the Working Principles for Risk Analysis for Application in the Framework of the Codex Alimentarius. The review might result in the CAC advising Codex Committees to amend their risk analysis principles document relevant to their area of work.

Timeline: Completion by 2011.

Responsible parties: CCGP

2.2 Review risk analysis principles developed by relevant Codex Committees

Description: Review risk analysis principles developed by relevant Codex Committees in the light of the experience gained when all relevant Codex Committees have elaborated risk assessment policies pertaining to their area of work, these policies having been adopted by the CAC.

As these risk assessment policies play a pivotal role in the interaction between risk managers and risk assessors, communication between these two parties should be further improved, where appropriate. The result of such a review may be revised documents on risk analysis principles for adoption by the Commission. The review should also take into account the outcome of the activities described under 2.1 and 2.3.

Timeline: Completion by 2013

Responsible parties: CAC, CCEXEC, CCFA, CCCF, CCPR, CCRVDF, CCFH, CCNFSDU

2.3 Enhance communication among relevant Codex subsidiary bodies and the FAO/WHO scientific expert bodies

Description: Enhance communication between the risk managers and risk assessors in accordance with paragraph 38 of the Working Principle for Risk Analysis for Application in the Framework of the Codex Alimentarius.

Timeline: Ongoing

Responsible parties: CCFA, CCCF, CCPR, CCRVDF, CCFH, CCNFSDU

2.4 Review the set of criteria recommended to FAO and WHO for prioritization of requests from Codex for scientific advice.

Description: Review the usefulness of the criteria agreed upon by the 28th Session of the CAC for use by FAO/WHO to prioritize requests from Codex for scientific advice.

Timeline: Completion by 2009.

Responsible parties: CCEXEC

2.5 Encourage countries to channel their requests for scientific advice to FAO / WHO through the CAC

Description: Encourage countries to channel their requests through the CAC in order to make the best use of the limited resources available at FAO and WHO for the provision of scientific advice. Encourage FAO and WHO to inform the CCEXEC and the CAC of all requests on provision of scientific advice on food safety received from member countries.

Timeline: Continuing

Responsible parties: CAC, CCEXEC, all subsidiary bodies

2.6 Encourage FAO/WHO to provide training and capacity building on risk analysis for food safety

Description: Assistance with capacity building activities aimed at effective implementation of the principles

Timeline: Continuing

Responsible parties: CAC, CCEXEC, FAO/WHO Coordinating Committees and Codex Members

Goal 3: Strengthening Codex Work-Management Capabilities

3.1 Review the Criteria for the Establishment of Work Priorities and procedures of the critical review carried out by the CCEXEC

Description: Review and revise, if necessary, the Criteria for the Establishment of Work Priorities and the effectiveness of the critical review process.

Timeline: Completion of analysis by CCEXEC by 2009; if there is need for revision, completion of revision by CCGP by 2011

Responsible parties: CCEXEC, CCGP

3.2 Ensure effective standards management

Description: Annually review progress of all subsidiary-body activities (i.e., standards, codes of practice, codes of hygienic practice, guidelines) against specified timeframes for completion of each activity, and recommend corrective actions to the Commission for activities that appear likely to exceed or have exceeded specified timeframes.

Timeline: Continuing

Responsible parties: CCEXEC

3.3 Develop committee-specific decision making and priority setting criteria

Description: Develop committee-specific decision-making and priority-setting criteria and use these criteria for management of work. Implement and review criteria where necessary.

Timeline: Completion of decision making and priority setting criteria by 2008. Review of criteria, beginning in 2008.

Responsible parties: All General Subject Committees and some other subsidiary bodies as appropriate

3.4 Analyse work-management approaches that facilitate advancement of texts in the Codex step process.

Description: Analyse work-management approaches used by committees that facilitate advancement of texts in the Codex step procedure in the light of the criteria referred to in 3.3 and their use by these committees.

Timeline: Completion of analysis by 2009

Responsible parties: Analysis work to be done possibly either by Codex Secretariat or by consultant for next step (3.5)

3.5 Adopt approaches proven to facilitate advancement of texts in the Codex step procedure by subsidiary bodies not currently using such approaches.

Description: Recommend adoption of approaches proven to facilitate advancement of texts in the Codex step procedure by subsidiary bodies not currently using such approaches, in the light of the analysis undertaken as per 3.4.

Timeline: Completion by 2011

Responsible parties: CCEXEC and CAC

3.6 Implement priority-ranked comprehensive accounting of all requests for scientific advice

Description: Request FAO and WHO annually to produce a priority-ranked comprehensive accounting (including budget information as it has impacts on Codex work) of all requests for scientific advice (i.e. continuing, ad hoc, requested by subsidiary bodies, or requested by members). The criteria that shall be used for priority ranking are those agreed to at the 55th Session of the Executive Committee (ALINORM 05/28/3). FAO and WHO are also requested to include budget information relevant to provision of scientific advice.

Timeline: Continuing

Responsible parties: CCEXEC, CAC, FAO and WHO

3.7 Evaluate the capacity of the Codex Secretariat to perform its function effectively

Description: Evaluate the effectiveness and resource needs of the Codex Secretariat in the operation and work management of the Commission and its subsidiary bodies, and in communicating with and serving the needs of Codex Contact Points.

Timeline: Completion by 2009

Responsible parties: Codex Secretariat, CCEXEC, CAC

3.8 Streamline Codex Commodity work

Description: Implement the decisions of the Commission on how to streamline Codex work on commodities through an improved structure of Codex subsidiary bodies.

Timeline: Completion by 2010

Responsible parties: CCEXEC, CAC

Goal 4: Promoting Cooperation between Codex and Other Relevant International Organizations

4.1 Track the activities of other international standard-setting bodies
Description: Track the activities of other international standard-setting bodies to identify areas of potential complementarities, gaps, duplication, or conflict. A summary of such activities relevant to Codex shall be reported to the Executive Committee and to the Commission annually.
Timeline: Continuing
Responsible parties: CAC, CCEXEC, Codex Secretariat, subsidiary bodies
4.2 Encourage Codex contributions to the work of other international bodies
Description: Encourage other relevant international bodies, when elaborating food standards and related texts, to take due account of Codex standards, related texts and any relevant ongoing work. Where appropriate, propose inclusion of appropriate cross-reference to Codex standards and relevant texts.
Timeline: Continuing
Responsible parties: Codex Secretariat
4.3 Encourage contributions from other international bodies in Codex work
Description: Invite international bodies concerned with food safety and food quality to participate in the standards development process of Codex
Timeline: Ongoing
Responsible parties: Observers, Codex Secretariat
4.4 Consider cooperation with other relevant international intergovernmental organizations
Description: While recognizing the needs to further improve interaction with OIE and IPPC, where appropriate, explore possibilities for cooperation to ensure effective collaboration and coordination, and that such cooperation should be in line with the Guidelines on Cooperation between Codex and International Intergovernmental Organizations.
Timeline: Continuing
Responsible parties: Legal Counsels of FAO and WHO, Codex Secretariat
4.5 Promote interdisciplinary coordination at the national and regional level
Description: Encourage Codex member countries to establish effective mechanisms within their own countries so that horizontal coordination and communication occurs among national delegates to various food-standards-related international organizations. Invite members to develop evaluation criteria to assess the success of the mechanisms that they have established and report progress in this activity through their respective Codex Regional Coordinating Committees to the CAC.
Timeline: Completion by 2009
Responsible parties: Codex Members, FAO/WHO Coordinating Committees

Goal 5: Promoting Maximum and Effective Participation of Members

<p>5.1 Promote Enhanced Participation of developing countries in Codex</p> <p>Description: Request FAO/WHO to encourage current donors to continue to provide funds to the FAO/WHO Trust Fund and invite other donors to contribute to the Fund to ensure sustainability. Request FAO/WHO to analyse the impact of the Codex Trust Fund on the capacity of beneficiary countries and report its findings to the CCEXEC and the Commission. Provide recommendations to FAO / WHO with a view to improved operation of the Trust Fund based on the outcome of the Trust Fund mid-term evaluation.</p> <p>Timeline: Continuing</p> <p>Responsible parties: CAC, CCEXEC</p>
<p>5.2 Promote effective use of written comments in the Codex process</p> <p>Description: Encourage members and observers to make maximum use of opportunities to submit written comments in response to Circular Letters (CLs) while respecting the deadlines for such submissions to allow all members and observers to study the positions of other members and observers in a timely manner.</p> <p>Codex Secretariat and Chairs of the Committees will examine how best to ensure that written comments of members that are not present at the meetings are taken into consideration, and how to handle the late submission of comments in response to CLs, from the view points of transparency and inclusiveness.</p> <p>Timeline: Continuing</p> <p>Responsible parties: Codex Members, Observers, CCEXEC, Codex Secretariat</p>
<p>5.3 Evaluate effectiveness of Codex Committee sessions held in developing countries</p> <p>Description: Evaluate the effectiveness of holding Codex sessions in developing countries in terms of enhanced participation. Analyse the effectiveness of co-hosting arrangements, and continue to explore possibilities of convening Codex sessions outside the host countries.</p> <p>Timeline: Completion by 2009</p> <p>Responsible parties: Host countries, CCEXEC</p>
<p>5.4 Strengthen Codex Contact Points and National Codex Committees</p> <p>Description: Request FAO and WHO to provide technical assistance for the strengthening of national Codex structures; provide improved support by the Codex Secretariat to Codex Contact Points through the effective use of Internet facilities.</p> <p>Timeline: Continuing</p> <p>Responsible parties: CAC, CCEXEC, Codex Secretariat</p>
<p>5.5 Enhance participation of non-governmental organizations at international, regional and national levels</p> <p>Description: Encourage non-governmental organizations to participate in Codex work at national, regional and international levels. Encourage members to establish sound structures and processes for consultation on Codex matters to ensure effective involvement and participation of all interested parties.</p> <p>Timeline: Continuing</p> <p>Responsible parties: CAC, Codex Members, subsidiary bodies</p>

5.6 Enhance communication about Codex work at international and national levels

Description: Develop new communication approaches to promote the work of Codex at national and international levels. Develop direct and easily understandable messages on Codex to interested parties including consumers and especially emphasizing high level policy makers. .

Timeline: Continuing

Responsible parties: Codex Secretariat, WHO and FAO, Codex Contact Points, subsidiary bodies

PART 3
IMPLEMENTATION OF STRATEGIC PLAN

This Part contains two tables:

- Table 1: Implementation of Strategic Plan

(This table is a checklist of the Strategic Plan activities to monitor the progress and achievement of the activities listed in Part 2. This table will be regularly updated for review by the Executive Committee.)

- Table 2: Critical Review of Proposals for New Work and Monitoring Progress of Standards Development

(This table is a checklist of ongoing work, to manage current and future work undertaken by the subsidiary bodies of the Commission. This Part will regularly be presented for critical review by the Executive Committee to monitor the progress of the ongoing work of standards setting mentioned in 1.1, 1.2, 1.3 and 1.4 of Part 2.)

Table 1: Implementation of Strategic Plan

Goal	Activities	Responsible parties	Timeframe	Output/Measurable indicators	Current Status	Notes *	Advice by EXEC	Decision by CAC
Goal 1: Promoting Sound Regulatory Frameworks	1.1 Review and develop Codex standards and related texts for food safety	CCFH, CCFA, CCCF, CCPR, CCRVDF, CCNFSDU, relevant Task Forces and Commodity Committees	Continuing	Standards related texts adopted at respective steps	See Table 2			
	1.2 Review and develop Codex standards and related texts for food quality	Relevant Task Forces, Commodity Committees and FAO/WHO Coordinating Committees	Continuing	Standards and related texts adopted at respective steps	See Table 2			
	1.3 Review and develop Codex standards and related texts for food labelling and nutrition	CCFL, CCNFSDU	Continuing	Standards and related texts adopted at respective steps	See Table 2			

	1.4 Review and develop Codex standards and related texts for food inspection and certification, and methods of sampling and analysis	CCMAS, CCFICS	Continuing	Standards and related texts adopted at respective steps	See Table 2			
	1.5 Develop guidance for safe and prudent non-human antimicrobial usage for containment of resistance	Existing relevant Codex Committees, <i>ad hoc</i> Intergovernmental Task Force on Antimicrobial Resistance	Completion by 2011	Guidance for safe and prudent non-human antimicrobial usage for containment of resistance	See Table 2			
	1.6 Explore innovative risk management frameworks	CCRVDF, CCPR	Completion by 2009	Reports by CCRVDF and CCPR respectively to CCEXEC and CAC				
	1.7 Encourage FAO/WHO to expand capacity building programmes	CAC, CCEXEC and FAO/WHO Coordinating Committees	Continuing	Report from FAO/WHO to CAC, CCEXEC and Coordinating Committees				
	1.8 Publish and disseminate the Codex Alimentarius	Codex Secretariat, Codex Contact Points	Continuing	Publication and dissemination of Codex Alimentarius				

* (Notes from Committees, Secretariat, whichever necessary)

Goal	Activities	Responsible parties	Timeframe	Output/Measurable indicators	Current Status	Notes *	Advice by EXEC	Decision by CAC
Goal 2: Promoting Widest and Consistent Application of Scientific Principles and Risk Analysis	2.1 Review the consistency of risk analysis principles elaborated by the relevant Codex Committees	CCGP	Completion by 2011	Report by CCGP of completed review to CAC.	See Table 2			
	2.2 Review risk analysis principles developed by relevant Codex Committees	CAC, CCEXEC, CCFA, CCCF, CCPR, CCRVDF, CCFH, CCFNSDU	Completion by 2013	Report by relevant Committees of completed review, taking into account the review of the activities in 2.1 and 2.3.	See Table 2			
	2.3 Enhance communication among relevant Codex subsidiary bodies and the FAO/WHO scientific expert bodies	CCFA, CCCF, CCPR, CCRVDF, CCFH, CCFNSDU	Ongoing	Incorporated into report as required in 2.2.				
	2.4 Review the set of criteria recommended to FAO and WHO for prioritization of requests from Codex for scientific advice.	CCEXEC	Completion by 2009	Report of review with recommendation to better match of priorities and resources by CCEXEC to CAC				

	2.5 Encourage countries to channel their requests for scientific advice to FAO / WHO through the CAC	CAC, CCEXEC, all subsidiary bodies	Continuing	Reports by FAO and WHO of requests for scientific advice received directly from countries vs. requests received through CAC.				
	2.6 Encourage FAO/WHO to provide training and capacity building on risk analysis for food safety	CAC, CCEXEC, FAO/WHO Coordinating Committees and Codex Members	Continuing	Report from FAO/WHO to CAC, CCEXEC and Coordinating Committees	See Table 2			

* (Notes from Committees, Secretariat, whichever necessary)

Goal	Activities	Responsible parties	Timeframe	Output/Measurable indicators	Current Status	Notes *	Advice by EXEC	Decision by CAC
Goal 3: Strengthening Codex Work-Management Capabilities	3.1 Review the Criteria for the Establishment of Work Priorities and procedures of the critical review carried out by the CCEXEC	CCEXEC, CCGP	Completion by 2009 Completion by 2011	Report by CCEXEC on analysis of the critical review process If there is need for revision, revise the Criteria for the Establishment of Work Priorities by CCGP				
	3.2 Ensure effective standards management	CCEXEC	Continuing	Report on compliance of standards development against timelines (associated with critical review process)	See Table 2			

	3.3 Develop committee-specific decision making and priority setting criteria	All General Subject Committees and some other subsidiary bodies as appropriate	Completion by 2008 Continuing	Committee-specific decision-making and priority-setting criteria Confirmed review of criteria beginning in 2008				
	3.4 Analyse work-management approaches that facilitate advancement of texts in the Codex step process	Either by Codex Secretariat or by consultant for next step (3.5)	Completion by 2009	Report to the CCEXEC and CAC on analysis of work-management approaches				
	3.5 Adopt approaches proven to facilitate advancement of texts in the Codex step procedure by subsidiary bodies not currently using such approaches	CCEXEC and CAC	Completion by 2011	Adoption by CAC on work - management approaches.				
	3.6 Implement priority-ranked comprehensive accounting of all requests for scientific advice	CCEXEC, CAC, FAO and WHO	Continuing	Comprehensive report by FAO/WHO to CAC on accounting of all requests for scientific advice				
	3.7 Evaluate the capacity of the Codex Secretariat to perform its function effectively	Codex Secretariat, CCEXEC, CAC	Completion by 2009	Report by Secretariat to CAC on the staff and other key resources				

	3.8 Streamline Codex Commodity work	CCEXEC, CAC	Completion by 2010	Accomplished Implementation of the decisions of the Commission on how to streamline Codex work on commodities through an improved structure of Codex subsidiary bodies				
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* (Notes from Committees, Secretariat, whichever necessary)

Goal	Activities	Responsible parties	Timeframe	Output/Measurable indicators	Current Status	Notes *	Advice by EXEC	Decision by CAC
Goal 4: Promoting Cooperation between Codex and other relevant international organizations	4.1 Track the activities of other international standard-setting bodies	CAC, CCEXEC, Codex Secretariat, subsidiary bodies	Continuing	Report to the CCEXEC and CAC indicating potential complementarities, gaps, duplication, or conflict with the work of other international organizations				
	4.2 Encourage Codex contributions to the work of other international bodies	Codex Secretariat	Continuing	Number of standards elaborated by other international organisations with a cross reference to Codex standards and relevant texts				
	4.3 Encourage contributions from other international bodies in Codex work	Observers, Codex Secretariat	Continuing	Number of Codex standards elaborated with identifiable input from other international organisations				

	4.4 Consider cooperation with other relevant international intergovernmental organizations	Legal Counsels of FAO and WHO, Codex Secretariat	Continuing	Devise the means by which Codex can strengthen cooperation with OIE and IPPC				
	4.5 Promote interdisciplinary coordination at the national and regional level	Codex Members, FAO/WHO Coordinating Committees	Completion by 2009	Reports from members to Regional Coordinating Committees on mechanisms and evaluation criteria.				

* (Notes from Committees, Secretariat, whichever necessary)

Goal	Activities	Responsible parties	Timeframe	Output/Measurable indicators	Current Status	Notes *	Advice by EXEC	Decision by CAC
Goal 5: Promoting Maximum and Effective Participation of Members	5.1 Promote enhanced participation of developing countries in Codex	CAC, CCEXEC	Continuing	Reports from FAO / WHO presenting analysis of measures of enhanced participation achieved through the Codex Trust Fund.				
	5.2 Promote effective use of written comments in the Codex process	Codex Members, Observers, CCEXEC, Codex Secretariat	Continuing	Reports by host countries on patterns of submission of written comments in response to CLs, and adherence by chairs to the guidelines for the conduct of the meetings.				
	5.3 Evaluate effectiveness of Codex Committee sessions held in developing countries	Host countries, CCEXEC	Completion by 2009	Reports from host and co-hosting countries documenting experience co-hosting experience.				

	5.4 Strengthen Codex Contact Points and National Codex Committees	CAC, CCEXEC, Codex Secretariat	Continuing	Reports from FAO/WHO on countries whose National structure and codex Contact Points have been supported.				
	5.5 Enhance participation of non-governmental organizations at international, regional and national levels	CAC, Codex Members, subsidiary bodies	Continuing	Reports from member countries under relevant agenda items of the Regional Committees, on participations of non-governmental organizations at National level.				
	5.6 Enhance communication about Codex work at international and national levels	Codex Secretariat, WHO and FAO, Codex Contact Points, subsidiary bodies	Continuing	Reports by Secretariat to CAC on increased use of audio / webcasting, enhancement of webpages, increase use of electronic distribution of codex materials, etc.				

* (Notes from Committees, Secretariat, whichever necessary)

Table 2: Critical Review of Proposals for New Work and Monitoring Progress of Standards Development**Sample**

<i>Codex Committee on xx</i> ¹ Document title	Timeframe		Current Status ²	Relevant Output Codes ³	Provision of scientific advice	Explanatory notes	Specific Comments from the Chairperson of the Committee	Advice given by the Executive Committee
	Job ID ⁴	Target Year ⁵						
Draft Guidelines for A	N03-2005	2009	6/7	1.2	Not required			
Draft Standard for B	N04-2006	2011	5	1.3	Not required			
Proposed Draft Guidelines for C	N05-2006	2011	3/4	1.2	FAO/WHO Expert Consultation planned in 2009			
Proposed Draft Standard for D	N04-2008	2013	2	1.3	FAO/WHO Expert Consultation requested by 2011			
General comments by the Chairperson / host countries of the Committee:								

¹ Name of the subsidiary body

² Step in the Elaboration Procedure

³ Reference made to Part 2 of the Strategic Plan

⁴ Job IDs are assigned by the Commission upon approval as new work

⁵ Year by which the draft text is to be adopted at Step 8, as agreed by the Commission on the basis of the Project Document

APPENDIX X

CHAIRMANSHIP OF CODEX SUBSIDIARY BODIES

Subsidiary Bodies established under Rule XI.1(b)(i)

Code	Subsidiary Body	Member Responsible	Status
CX 703	Codex Committee on Milk and Milk Products	New Zealand	Active
CX 708	Codex Committee on Cocoa Products and Chocolate	Switzerland	<i>Sine die</i>
CX 709	Codex Committee on Fats and Oils	Malaysia	Active
CX 710	Codex Committee on Sugars	United Kingdom	<i>Sine die</i>
CX 711	Codex Committee on Food Additives	China	Active
CX 735	Codex Committee on Contaminants in Foods	The Netherlands	Active
CX 712	Codex Committee on Food Hygiene	United States of America	Active
CX 713	Codex Committee on Processed Fruits and Vegetables	United States of America	Active
CX 714	Codex Committee on Food Labelling	Canada	Active
CX 715	Codex Committee on Methods of Analysis and Sampling	Hungary	Active
CX 716	Codex Committee on General Principles	France	Active
CX 718	Codex Committee on Pesticide Residues	China	Active
CX 719	Codex Committee on Natural Mineral Waters	Switzerland	Active
CX 720	Codex Committee on Nutrition and Foods for Special Dietary Uses	Germany	Active
CX 722	Codex Committee on Fish and Fishery Products	Norway	Active
CX 723	Codex Committee on Meat Hygiene	New Zealand	<i>Sine die</i>
CX 728	Codex Committee on Vegetable Proteins	Canada	<i>Sine die</i>
CX 729	Codex Committee on Cereals, Pulses and Legumes	United States of America	<i>Sine die</i>
CX 730	Codex Committee on Residues of Veterinary Drugs in Foods	United States of America	Active
CX 731	Codex Committee on Fresh Fruits and Vegetables	Mexico	Active
CX 733	Codex Committee on Food Import and Export Certification and Inspection Systems	Australia	Active

Ad hoc Intergovernmental Task Force established by the 27th Session of the Commission

CX 802	<i>Ad hoc</i> Codex Intergovernmental Task Force on Foods derived from Biotechnology	Japan	Active
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Ad hoc Intergovernmental Task Force established by the 29th Session of the Commission

CX 804	<i>Ad hoc</i> Codex Intergovernmental Task Force on Antimicrobial Resistance	Republic of Korea	Active
CX 805	<i>Ad hoc</i> Codex Intergovernmental Task Force on the Processing and Handling of Quick Frozen Foods	Thailand	Active

Subsidiary Bodies established under Rule XI.1(b)(i)

Code	Subsidiary Body	Member Responsible
CX 706	FAO/WHO Coordinating Committee for Europe	Coordinator for Europe
CX 707	FAO/WHO Coordinating Committee for Africa	Coordinator for Africa
CX 725	FAO/WHO Coordinating Committee for Latin America and the Caribbean	Coordinator for Latin America and the Caribbean
CX 727	FAO/WHO Coordinating Committee for Asia	Coordinator for Asia
CX 732	FAO/WHO Coordinating Committee for North America and the South West Pacific	Coordinator for North America and the South West Pacific
CX 734	FAO/WHO Coordinating Committee for the Near East	Coordinator for the Near East
