This section of the FEDERAL REGISTER contains documents other than rules or proposed rules that are applicable to the public. Notices of hearings and investigations, committee meetings, agency decisions and rulings, delegations of authority, filing of petitions and applications and agency statements of organization and functions are examples of documents appearing in this section.

DEPARTMENT OF AGRICULTURE

Food Safety and Inspection Service

[FDMS Docket No. FSIS–2007–0006]

International Standard-Setting Activities

AGENCY: Food Safety and Inspection Service, USDA.

ACTION: Notice.

SUMMARY: This notice informs the public of the sanitary and phytosanitary standard-setting activities of the Codex Alimentarius Commission (Codex), in accordance with section 491 of the Trade Agreements Act of 1979, as amended, and the Uruguay Round Agreements Act, Public Law 103–465, 108 Stat. 4809. This notice also provides a list of other standard-setting activities of Codex, including commodity standards, guidelines, codes of practice, and revised texts. This notice, which covers the time periods from June 1, 2006, to May 31, 2007, and June 1, 2007, to May 31, 2008, seeks comments on standards under consideration and revised texts. This notice, which contains documents other than rules or regulations, also will be available for public inspection in the FSIS Docket Room at the address listed above between 8:30 a.m. and 4:30 p.m., Monday through Friday. The comments also will be posted on the Agency’s Web site at http://www.fsis.usda.gov/regulations_policies/2007_Notices_Index/index.asp.

FOR FURTHER INFORMATION CONTACT: F. Edward Scarbrough, PhD, United States Manager for Codex, U.S. Department of Agriculture, Office of the Under Secretary for Food Safety, Room 4861, South Agriculture Building, 1400 Independence Avenue, SW., Washington, DC 20250–3700; (202) 205–7760. For information pertaining to Codex are accessible via the World Wide Web at the following address: http://www.codexalimentarius.net/current.asp. The U.S. Codex Office also maintains a Web site at http://www.fsis.usda.gov/Regulations_Policies/Codex_Alimentarius/index.asp.

SUPPLEMENTARY INFORMATION:

Background

The World Trade Organization (WTO) was established on January 1, 1995, as the common international institutional framework for the conduct of trade relations among its members in matters related to the Uruguay Round Trade Agreements. The WTO is the successor organization to the General Agreement on Tariffs and Trade (GATT). U.S. membership in the WTO was approved and the Uruguay Round Agreements Act was signed into law by the President on December 8, 1994. The Uruguay Round Agreements became effective, with respect to the United States, on January 1, 1995. Pursuant to section 491 of the Trade Agreements Act of 1979, as amended, the President is required to designate an agency to be “responsible for informing the public of the sanitary and phytosanitary (SPS) standard-setting activities of each international standard-setting organization.” The main organizations are Codex, the World Organisation for Animal Health, and the International Plant Protection Convention. The President, pursuant to Proclamation No. 6780 of March 23, 1995 (60 FR 15845), designated the U.S. Department of Agriculture as the agency responsible for informing the public of SPS standard-setting activities of each international standard-setting organization. The Secretary of Agriculture has delegated to the Administrator, Food Safety and Inspection Service (FSIS), the responsibility to inform the public of the SPS standard-setting activities of Codex. The FSIS Administrator has, in turn, assigned the responsibility for informing the public of the SPS standard-setting activities of Codex to the U.S. Codex Office, FSIS.

Codex was created in 1962 by two U.N. organizations, the Food and Agriculture Organization (FAO) and the World Health Organization (WHO). Codex is the principal international organization for encouraging fair international trade in food and protecting the health and economic interests of consumers. Through adoption of food standards, codes of practice, and other guidelines developed by its committees and by promoting their adoption and implementation by governments, Codex seeks to protect the health of consumers, ensure fair trade practices in the food trade, and promote coordination of food standards work undertaken by international governmental and non-governmental organizations. In the United States, the United States Department of Agriculture (USDA); the Food and Drug Administration (FDA), Department of Health and Human...
would like to access or receive information about specific committees. The information provided in Attachment 1 describes the status of Codex standard-setting activities by the Codex Committees for the time periods from June 1, 2006, to May 31, 2007, and June 1, 2007, to May 31, 2008. Attachment 2 provides the list of U.S. Codex Officials (includes U.S. delegates and alternate delegates). A list of forthcoming Codex sessions may be found at: http://www.codexalimentarius.net/web/current.jsp?lang=en.

Additional Public Notification
Public awareness of all segments of rulemaking and policy development is important. Consequently, in an effort to ensure that the public and in particular minorities, women, and persons with disabilities, are aware of this notice, FSIS will announce it on-line through the FSIS Web page located at http://www.fsis.usda.gov/regulations&_policies/2007_Notices_Index/index.asp.

The Regulations.gov Web site is the central online rulemaking portal of the United States government. It is being offered as a public service to increase participation in the Federal government’s regulatory activities. FSIS participates in Regulations.gov and will accept comments on documents published on the site. The site allows visitors to search by keyword or Department or Agency for rulemakings that allow for public comment. Each entry provides a quick link to a comment form so that visitors can type in their comments and submit them to FSIS. The Web site is located at http://www.regulations.gov.

FSIS also will make copies of this Federal Register publication available through the FSIS Constituent Update, which is used to provide information regarding FSIS policies, procedures, regulations, Federal Register notices, FSIS public meetings, recalls, and other types of information that could affect or would be of interest to our constituents and stakeholders. The update is communicated via Listserv, a free e-mail subscription service consisting of industry, trade, and farm groups, consumer interest groups, allied health professionals, scientific professionals, and other individuals who have requested to be included. The update also is available on the FSIS Web page. Through Listserv and the web page, FSIS is able to provide information to a much broader, more diverse audience.

In addition, FSIS offers an e-mail subscription service which provides automatic and customized access to selected food safety news and information. This service is available at http://www.fsis.usda.gov/news_and_events/email_subscription/. Options range from recalls to export information to regulations, directives and notices. Customers can add or delete subscriptions themselves and have the option to password protect their account.

Done at Washington, DC on: May 23, 2007.

F. Edward Scarbrough,
United States Manager for Codex.

Attachment 1
Sanitary and Phytosanitary Activities of Codex Alimentarius Commission and Executive Committee

The Codex Alimentarius Commission will hold its Thirtieth Session July 2–7, 2007, in Rome, Italy. At that time, it will consider procedural matters, and the standards, codes of practice, and related matters brought to its attention by the general subject committees, commodity committees, ad hoc Task Forces and member delegations. It will also consider options to implement recommendations from the review of Codex committee structure and mandates of Codex committees and task forces, as well as budgetary and strategic planning issues. At this Session, the Commission will elect a Chair and three Vice Chairs.

Prior to the Commission meeting, the Executive Committee will have met at its Fifty-ninth Session on June 26–30, 2007. It is composed of the chairperson, vice-chairpersons, and seven members elected from the Commission, one from each of the following geographic regions: Africa, Asia, Europe, Latin America and the Caribbean, Near East, North America, and South-West Pacific. Additionally, regional coordinators from the six regional committees serve as members of the Executive Committee. It will consider the Codex Strategic Plan 2006–1013; review the Codex committee structure and mandate of Codex committees and task forces; review matters arising from reports of Codex Committees, proposals for new work, and standards management issues; and review the Trust Fund for the Participation of Developing Countries and Countries in Transition in the Work of the Codex Alimentarius.

Responsible Agency: USDA/FSIS.
U.S. Participation: Yes.

Codex Committee on Residues of Veterinary Drugs in Foods

The Codex Committee on Residues of Veterinary Drugs in Foods determines priorities for the consideration of residues of veterinary drugs in foods...
and recommends Maximum Residue Limits (MRLs) for veterinary drugs. A veterinary drug is defined as any substance applied or administered to a food producing animal, such as meat or dairy animals, poultry, fish or bees, for therapeutic, prophylactic or diagnostic purposes or for modification of physiological functions or behavior.

A Codex Maximum Limit for Veterinary Drugs (MRLVD) is the maximum concentration of residue resulting from the use of a veterinary drug (expressed in mg/kg or ug/kg on a fresh weight basis) that is adopted by the Codex Alimentarius Commission to be permitted or recognized as acceptable in or on a food. An MRLVD is based on the Acceptable Daily Intake (ADI) and indicates the amount of residue in food that is considered to be without appreciable toxicological hazard. An MRLVD also takes into account other relevant public health risks as well as food technological aspects.

When establishing an MRLVD, consideration is also given to residues that occur in food of plant origin and/ or the environment. Furthermore, the MRLVD may be reduced to be consistent with good practices in the use of veterinary drugs and to the extent that practical analytical methods are available.

Acceptable Daily Intake (ADI): An estimate by the Joint FAO/WHO Expert Committee on Food Additives (JECFA) of the amount of a veterinary drug expressed on a body weight basis, that can be ingested daily over a lifetime without appreciable health risk (standard man = 60 kg).

The Committee will meet in the United States on September 3–7, 2007. The Committee will continue work on the following:

The Committee worked on:
- Draft MRLs for Flumequine, Melengestrol acetate, Colistin, Ractopamine, Erithromycin, Triclabendazole.
- Proposed Draft Guidelines for the Design and Implementation of National Regulatory Food Safety Assurance Programmes Associated with the Use of Veterinary Drugs in Food Producing Animals.
- Risk Analysis Principles Applied by the Codex Committee on Residues of Veterinary Drugs in Foods.
- Risk Assessment Policy for the Setting of MRLs in Food.
- Priority List of Veterinary Drugs Requiring Evaluation or Reevaluation.
- Compendium of Methods of Analysis Identified as Suitable to Support Codex MRLs.
- Discussion Paper on Risk Management Topics and Options for the CRVRDF.
- Report of the Working Group on Residues of Veterinary Drugs without ADI/MRL.

Responsible Agencies: HHS/FDA; USDA/FSIS.

U.S. Participation: Yes.

Codex Committee on Contaminants in Foods

The Codex Committee on Contaminants in Foods (CCCF) was established by the 29th Session of the Commission when it decided to split the former Codex Committee on Additives and Contaminants into two committees. The CCCF establishes or endorses permitted maximum levels for contaminants and naturally occurring toxicants in food and feed, prepares priority lists of contaminants and naturally occurring toxicants for risk assessment by the Joint FAO/WHO Expert Committee on Food Additives (JECFA), considers methods of analysis and sampling for the determination of contaminants and naturally occurring toxicants in food and feed, considers and elaborates standards or codes of practice for related subjects, and considers other matters assigned to it by the Commission in relation to contaminants and naturally occurring toxicants in food and feed.

The Committee held its first session in Beijing, China, on April 16–20, 2007. The relevant document is ALINORM 07/30/41. The following items will be considered at the 30th Session of the Commission on July 2–7, 2007.

To be considered at Step 5:
- Proposed Draft Maximum Levels for Tin in Canned Foods (other than beverages) and in Canned Beverages.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
- Proposed Draft Code of Practice on the Reduction of Chloropropanols During the Production of Acid-HVPs.
- Proposed Draft Code of Practice on Risk Assessment by JECFA.
Committees and General Committees:
• Food Additives
  • Endorsement and/or Revision of
    Maximum Levels for Food Additives
    and Processing Aids in Codex
    Standards.
  • Inclusion of Food Additive
    Provisions of Commodity Standards into
    the Codex General Standard for Food
    Additives.
  • General Standard for Food
    Additives: Draft Food Additive
    Provisions (in Tables 1, 2 and 3).
• Revisions to the General Standard
  for Food Additives’ Food Category
  System: Project Document.
• Guidelines for the Use of
  Flavourings.
• Inventory of Processing Aids.
• International Numbering System
  and Harmonization of Terms Used by
  Codex and JECFA.
• Revision of the Class Names and
  International Numbering System for
  Food Additives.
• Specifications for the Identity and
  Purity of Food Additives.
• Priority List of Food Additives
  Proposed for Evaluation by JECFA.
U.S. Participation: Yes.

Codex Committee on Pesticide Residues

The Codex Committee on Pesticide
Residues recommends to the Codex
Alimentarius Commission
establishment of maximum limits for
pesticide residues for specific food
items or in groups of food. A Codex
Maximum Residue Limit for Pesticide
(MRLP) is the maximum concentration
of a pesticide residue (expressed as mg/
kg), recommended by the Codex
Alimentarius Commission to be legally
permitted in or on food commodities
and animal feeds. Foods derived from
commodities that comply with the
respective MRLPs are intended to be
toxicologically acceptable, that is,
consideration of the various dietary
residue intake estimates and
determinations both at the national and
international level in comparison with
the ADI*, should indicate that foods
complying with Codex MRLPs are safe
for human consumption.

Codex MRLPs are primarily intended
to apply in international trade and are
derived from reviews conducted by the
Joint Meeting on Pesticide Residues
(JMPR).
(a) Review of residue data from
    supervised trials and supervised uses
    including those reflecting national good
    agricultural practices (GAP). Data from
    supervised trials conducted at the
    highest nationally recommended,
    authorized, or registered uses are
    included in the review. In order to
    accommodate variations in national pest
    control requirements, Codex MRLPs
take into account the higher levels
    shown to arise in such supervised trials,
    which are considered to represent
    effective pest control practices.
(b) Toxicological assessments of the
    pesticide and its residue.
    The following items will be
    considered by the Commission at its
    30th Session in July 2007. The relevant
    document is ALINORM 07/30/24.
    To be considered at Step 8:
    • Draft and Draft Revised Maximum
      Residue Limits.
    • Proposed Draft Maximum Residue
      Limits.
    To be considered at Step 5:
    • Proposed Draft and Proposed Draft
      Revised Maximum Residue Limits.
    To be considered for Revocation:
    • Codex CLX–D9.
    To be considered for New Work:
    • Priority List of Pesticides for review
      by JMPR.
    • The committee is continuing work on:
      • Draft and Proposed Draft MRLs.
      • Revision of the List of
        Recommended Methods on Analysis for
        Pesticide Residues.
      • Revision of the Codex Priority List
        of Pesticides for review by JMPR.
      • Discussion paper on the how Codex
        MRLs are used at the national level.
      • Discussion paper on the
        establishment of MRLs for Processed or
        Ready-to-Eat Foods.
      • Extended Revision of the Codex
        Classification of foods and animal feeds.
      *Acceptable Daily Intake (ADI) of a
        chemical is the daily intake which,
during an entire lifetime, appears to be
        without appreciable risk to the health of
        the consumer on the basis of all the
        known facts at the time of the
        evaluation of the chemical by the Joint
        FAO/WHO Meeting on Pesticide
        Residues. It is expressed in milligrams
        of the chemical per kilogram of body
        weight.

Responsible Agencies: EPA; USDA/AMS.
U.S. Participation: Yes.

Codex Committee on Methods of
Analysis and Sampling

The Codex Committee on Methods of
Analysis and Sampling:
(a) Defines the criteria appropriate to
    Codex Methods of Analysis and
    Sampling;
(b) Serves as a coordinating body for
    Codex with other international groups
    working in methods of analysis and
    sampling and quality assurance systems
    for laboratories;
(c) Specifies, on the basis of final
    recommendations submitted to it by the
    other bodies referred to in (b) above,
    Reference Methods of Analysis and
    Sampling appropriate to Codex
    Standards which are generally
    applicable to a number of foods;
(d) Considers, amends, if necessary,
    and endorses, as appropriate, methods
    of analysis and sampling proposed by
    Codex (Commodity) Committees, except
    that methods of analysis and sampling
    for residues of pesticides or veterinary
    drugs in food, the assessment of
    microbiological quality and safety in
    food, and the assessment of
    specifications for food additives do not
    fall within the terms of reference of this
    Committee;
(e) Elaborates sampling plans and
    procedures, as may be required;
(f) Considers specific sampling and
    analysis problems submitted to it by the
    Commission or any of its Committees;
    and
(g) Defines procedures, protocols,
    guidelines or related texts for the
    assessment of food laboratory
    proficiency, as well as quality assurance
    systems for laboratories.

The 28th Session of the Committee
met in Budapest, Hungary, on March
5–9, 2007. The relevant document is
ALINORM 07/30/23. For endorsement
at the 30th Commission in 2007:
• Proposed Amendment to the
  Principles for the Establishment of
  Codex Sampling Procedures (Procedural
  Manual).
• Endorsement of methods of analysis
  in Draft Standards and existing
  standards.
• Reference to IUPA/ISO/AOAC
  Protocols (amendment to references).
The Committee will continue to work
on:
• Draft Guidelines for Evaluating
  Acceptable Methods of Analysis.
• Draft Guidelines for Settling of
  Disputes on Analytical (Test) Results.
• Proposed Draft Guideline on
  Analytical Terminology.
• Conversion of methods for trace
  elements into criteria.
• Criteria for methods of analysis for
  foods derived from biotechnology.
• Guidance on measurement
  uncertainty and uncertainty of
  sampling.
• Discussion paper on role and terms
  of reference of CCMAS.
• Discussion paper on the reliability of
  analytical data.

Responsible Agencies: HHS/FDA;
USDA/GIPSA.
U.S. Participation: Yes.

Codex Committee on Food Import and
Export Inspection and Certification
Systems

The Codex Committee on Food Import
and Export Inspection and Certification
systems is charged with developing principles and guidelines for food import and export inspection and certification systems to protect consumers and to facilitate trade. Additionally, the Committee develops principles and guidelines for the application of measures by competent authorities to provide assurance that foods comply with essential requirements, especially statutory health requirements. This encompasses work on equivalence of food inspection systems, including equivalence agreements, processes and procedures to ensure that sanitary measures are implemented; guidelines on food import control systems; and guidelines on food product certification and information exchange. The development of guidelines for the appropriate utilization of quality assurance systems to ensure that foodstuffs conform to requirements and to facilitate trade also are included in the Committee’s terms of reference. The Committee met November 6–10, 2006. The reference document is ALINORM 07/30/30. The following will be considered for adoption by the Commission at its 30th Session in July 2007:

- Discussion paper on the reply to the question raised by the 22nd Session of the Codex Committee on General Principles regarding the revision of the Codex Code of Ethics for International Trade of Foods.
- Discussion Paper identifying areas for guidance for national food inspection systems.
- Discussion Paper on the development of Guidelines for the Conduct of Foreign Audit Team Inspections.
- Discussion Paper on the need of guidance on traceability/product tracing.

**Codex Committee on General Principles**

The Codex Committee on General Principles deals with procedure and general matters as are referred to it by the Codex Alimentarius Commission. The 24th Session was held on April 2–6, 2006, in Paris, France. The relevant document is ALINORM 07/30/33. Matters to be considered for adoption by the 29th Commission in July 2007:

- Proposed Draft Working Principles for Risk Analysis for Food Safety (Guidance to National Governments) for adoption at Step 5.
- Amendments to the Codex Procedural Manual clarifying the roles of Members elected to the Codex Executive Committee on a geographic basis and Regional Coordinators as members of the Executive Committee.
- Amendments to the Codex Procedural Manual dealing with the revision and amendment of Codex standards.
- Amendments to the General Principles of the Codex Alimentarius.
- Amendments to the Principles Concerning the Participation of International Non-Governmental Organizations in the Work of Codex.
- Risk Management Methodologies, including Risk Assessment Policies in the Codex Committee on Residues of Veterinary Drugs in Foods for inclusion in the Procedural Manual.
- Amendment to the Principles for the Establishment or Selection of Codex Sampling Procedures (Codex Procedural Manual).

The Committee continued work on:

- Code of Ethics for International Trade in Food (returned to Step 3).
- Consideration of the structure, content and presentation of the Procedural Manual.
- New definitions of risk analysis terms related to food safety.

**Responsible Agencies:** USDA/FSIS.

**U.S. Participation:** Yes.

**Codex Committee on Food Hygiene**

The Codex Committee on Food Hygiene has four primary responsibilities. First, to draft basic provisions on food hygiene applicable to all food. These provisions normally take the form of Codes of Hygienic Practice for a specific commodity (e.g., bottled water) or group of commodities (e.g., milk and milk products). Second, to suggest and prioritize areas where there is a need for microbiological risk assessment at the international level and to consider microbiological risk management matters in relation to food hygiene and in relation to the risk assessment activities of FAO and WHO. Third, to consider, amend if necessary, and endorse food hygiene provisions that are incorporated into specific Codex commodity standards by the Codex commodity committees. Fourth, to provide such other general guidance to the Commission on matters relating to food hygiene as may be necessary. The 38th Session of the Committee met in Houston, TX, on December 4–8, 2006. The relevant document is ALINORM 07/30/13. The following items will be considered by the Commission at its 30th Session in July 2007:

- Draft Amendment to the General Standard (Draft Recommendations for the Labelling of Foods obtained through certain techniques of GM/GE): Definitions.
- Proposed Draft Amendment to the General Standard for the Labelling of Prepackaged Foods: Quantitative Declaration of Ingredients.
- Proposed Draft Amendment to the Guidelines for Organically Produced Foods (Addition of Ethylene).
- Proposed Draft Definition of Advertising in relation to nutrition and health claims.

**Responsible Agencies:** HHS/FDA; USDA/FSIS.

**U.S. Participation:** Yes.

**Codex Committee on Food Labelling**

The Codex Committee on Food Labelling is responsible for drafting provisions on labelling issues assigned by the Codex Alimentarius Commission. The reference document is ALINORM 07/30/22. The Committee held its 35th Session at Ottawa, Canada, on April 30–May 4, 2007. It considered the following items:
Hygiene to the Control of Listeria monocytogenes in Ready-to-Eat Foods.

- Draft Code of Hygienic Practice for Eggs and Egg Products.
- Draft Principles and Guidelines for the Conduct of Microbiological Risk Management.

New Work:
- Proposed Draft Guidelines for the Control of Campylobacter and Salmonella spp. in Broiler (Young Bird) Chicken Meat.
- CCHF Risk Analysis Policies.

The committee will continue to work on:
- Proposed Draft Guidelines for Validation of Food Hygienic Control Measures.
- Proposed Draft Code of Hygienic Practice for Powdered Formulae for Infants and Children.
- Endorsement of Hygiene Provisions in Codex Standards and Codes of Practice.
- Microbiological Criteria for Listeria monocytogenes in Ready-to-Eat Foods.

**Responsible Agencies:** USDA/AMS; USDA/FSIS.

**U.S. Participation:** Yes.

### Codex Committee on Fresh Fruits and Vegetables

The Codex Committee on Fresh Fruits and Vegetables is responsible for elaborating world-wide standards and codes of practice for fresh fruits and vegetables. The Committee met in Mexico City, Mexico, on September 25–29, 2006. The relevant document is ALINORM 07/30/35. The following items will be considered by the Commission at its 30th Session in July 2007.

To be considered at Step 8:
- Draft Revised Advisory List of Nutrient Compounds for Use in Foods for Special Dietary Uses Intended for Infants and Young Children.
- Application of Risk Analysis to the Work of the CCNFSDU.

The Committee continues work on:
- Discussion Paper on Proposals for Additional or Revised Nutrient Reference Values (NRVs).

**Responsible Agencies:** HHS/FDA; USDA/ARS.

**U.S. Participation:** Yes.

### Codex Committee on Milk and Milk Products

The Codex Committee on Milk and Milk Products is responsible for establishing international codes and standards for milk and milk products. The Committee will hold its 8th Session in 2008 in New Zealand. The Committee is working on:

- Proposed Draft Model Export Certificate for Milk and Milk Products.
- Proposed Draft Amendment to the Codex Standard for Fermented Milks pertaining to Fermented Milk Drinks.
- Proposed Draft Standard for Processed Cheese.
- Amendment to the List of Additives of the Codex Standard for Creams and Prepared Creams.
- Food Additive Listings for the Codex Standard for Fermented Milks (flavoured fermented milks).
- Methods of Analysis and Sampling for Milk and Milk Products Standards.
- Discussion paper on sampling plans for milk products in presence of significant measurement error.

**Responsible Agencies:** USDA/AMS; USDA/FDA.

**U.S. Participation:** Yes.
Codex Committee on Fats and Oils

The Codex Committee on Fats and Oils is responsible for elaborating standards for fats and oils of animal, vegetable, and marine origin. The Committee met February 19–23, 2007. The relevant document is ALINORM 07/30/17. To be considered by the Commission at Step 8:

- Draft Standard for Fat Spreads and Blended Spreads.
- New Work:
- The Committee continues work on:
  - Draft List of Acceptable Previous Cargoes.
  - Unbleached palm oil: total carotenoids.
- Responsible Agencies: HHS/FDA; USDA/ARS.
- U.S. Participation: Yes.

Codex Committee on Processed Fruits and Vegetables

The Codex Committee on Processed Fruits and Vegetables is responsible for elaborating standards for Processed Fruits and Vegetables. The Committee met on October 16–21, 2006. The relevant document is ALINORM 07/30/27. The following items will be considered by the Commission at its 30th Session in July 2007.

- To be considered at Step 8:
  - Draft Codex Standard for Pickled Fruits and Vegetables.
  - Draft Codex Standard for Processed Tomato Concentrates.
  - Draft Codex Standard for Preserved (Canned) Tomatoes.
  - Draft Codex Standards for Certain Canned Citrus Fruits.
- To be considered at Step 5:
- The Committee continues to work on:
  - Standard Layout for Processed Fruits and Vegetables, Methods of Analysis for Processed Fruits and Vegetables.

- Priority List for the Standardization of Processed Fruits and Vegetables.
- Responsible Agencies: USDA/AMS; HHS/FDA.
- U.S. Participation: Yes.

Certain Codex Commodity Committees

Several Codex Alimentarius Commodity Committees have adjourned sine die. The following Committees fall into this category:

- Sugars. Responsible Agencies: USDA/ARS; HHS/FDA.
- U.S. Participation: Yes.

Codex Alimentarius Commission for a four year period of time, completed its work, but was re-established at the 27th Session of the Commission. The relevant document is ALINORM 07/30/34. The Committee will hold its 7th Session in Japan on November 26–30, 2007. The Task Force will discuss the following items:

- Proposed Draft Guideline for the Conduct of Food Safety Assessment of Foods Derived from Recombinant-DNA Animals.
- Proposed Draft Annex to the Guideline for the Conduct of Food Safety Assessment of Foods Derived from Recombinant-DNA Plants: Food Safety Assessment of Foods Derived from Recombinant DNA-Plants Modified for Nutritional or Health Benefits.
- Responsible Agencies: HHS/FDA; USDA/APHIS.
- U.S. Participation: Yes.

Ad Hoc Intergovernmental Task Force on the Processing and Handling of Quick Frozen Foods

The Ad hoc Intergovernmental Task Force on the Processing and Handling of Quick Frozen Foods was created by the 29th Session of the Commission to resolve all outstanding issues including the quality and safety provisions of the Code of Practice for the Processing and Handling of Quick Frozen Foods. The Task Force, hosted by Thailand, was given two years to finalize the Code. Thailand and the United States prepared a Circular Letter requesting comments on a revised Code. The resulting document prepared from these comments will serve as the basis for discussion at the Session of the Task Force that will take place in early 2008.

FAO/WHO Regional Coordinating Committees

The Codex Alimentarius Commission is made up of an Executive Committee, as well as approximately 30 subsidiary bodies. Included in these subsidiary bodies are coordinating committees for groups of countries located in proximity to each other who share common concerns. There are currently six Regional Coordinating Committees:

- Coordinating Committee for Africa.
- Coordinating Committee for Asia.
- Coordinating Committee for Europe.
- Coordinating Committee for Latin America and the Caribbean.
Coordinating Committee for the Near East.
- Coordinating Committee for North America and the South-West Pacific.

The United States participates as an active member of the Coordinating Committee for North America and the South-West Pacific, and is informed of the other coordinating committees through meeting documents, final reports, and representation at meetings. Each regional committee:
- Defines the problems and needs of the region concerning food standards and food control;
- Promotes within the committee contacts for the mutual exchange of information on proposed regulatory initiatives and problems arising from food control and stimulates the strengthening of food control infrastructures;
- Recommends to the Commission the development of world-wide standards for products of interest to the region, including products considered by the committee to have an international market potential in the future; and
- Serves a general coordinating role for the region and performs such other functions as may be entrusted to it by the Commission.

Codex Coordinating Committee for North America and the South-West Pacific

The Coordinating Committee is responsible for defining problems and needs concerning food standards and food control of all Codex member countries of the region. Items on the agenda for the next meeting may include:
- Draft new Strategic Plan for NASWP.
- Report of the Electronic Working Group on Objective 6 of the Strategic Plan for CCNASWP.
- Progress Report: Joint FAO/WHO Evaluation of the Codex Alimentarius and other FAO and WHO Work on Food Standards.
- Evaluation of the effectiveness of the Trust Fund for the participation of developing countries in Codex.
- Nomination of regional coordinator.

Responsible agency: USDA/FSIS.

U.S. Participation: Yes.

Attachment 2

U.S. Codex Alimentarius Officials

Codex Committee Chairpersons

Codex Committee on Food Hygiene
Dr. Karen Hulebak, Chief Scientist, Office of Public Health Science, Food Safety and Inspection Service, U.S. Department of Agriculture, 1400 Independence Avenue, SW., Room 3130, South Building, Washington, DC 20250–3700, Phone: (202) 720–5735, Fax: (202) 720–2980, E-mail: karen.hulebak@fsis.usda.gov.

Codex Committee on Residues of Veterinary Drugs in Foods
Dr. Stephen F. Sundlof, Director, Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Place (HFV–1), Rockville, MD 20855, Phone: (301) 827–2950, Fax: (301) 827–8401, E-mail: ssundlof@cvm.fda.gov.

Codex Committee on Cereals, Pulses and Legumes (adjourned sine die)
Mr. Steven N. Tanner, Director, Technical Services Division, Grain Inspection, Packers & Stockyards Administration, U.S. Department of Agriculture, 10383 N. Executive Hills Boulevard, Kansas City, MO 64153–1394, Phone: (816) 891–0401, Fax: (816) 891–0478, E-mail: Stephen.n.tanner@gsipa.usda.gov.

Codex Committee on Processed Fruits and Vegetables
Mr. Terry Bane, Branch Chief, Processed Products Branch, Fruit and Vegetable Programs, AMS, Room 0709, South Building, Stop 9247, 1400 Independence Avenue, SW., Washington, DC 20250–0247, Phone: (202) 720–4693, Fax: (202) 690–1087, E-mail: terry.bane@usda.gov.

Codex Committee on Residues of Veterinary Drugs in Foods
Dr. Stephen F. Sundlof, Director, Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Place (HFV–1), Rockville, MD 20855, Phone: (301) 827–2950, Fax: (301) 827–8401, E-mail: ssundlof@cvm.fda.gov.

Codex Committee on Contaminants in Foods
Mr. Steve J. Davis, Acting Director, Hazard Analysis and Critical Control Points (HACCP) Program, U.S. Department of Agriculture, 620 Central Avenue, Building 2–A, Alameda, CA 95501, Phone: (510) 337–5031, ext. 3004, Fax: (510) 337–5036, Emilio.Esteban@fsis.usda.gov.

Codex Committee on Food Additives (Host Government—China)

U.S. Delegate

Dennis M. Keefe, PhD, Office of Premarket Approval, Center for Food Safety and Applied Nutrition, FDA (HFS–200), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (202) 418–3113, Fax: (202) 418–3131, E-mail: dennis.keefe@fda.hhs.gov.

Alternate Delegate

Susan E. Carberry, PhD, Supervisory Chemist, Division of Petition Review, Office of Food Additive Safety (HFS–265), Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5100 Paint Branch Parkway, College Park, MD 20740, Phone: (301) 436–1269, Fax: (301) 436–2972, E-mail: Susan.Carberry@fda.hhs.gov.

Codex Committee on Pesticide Residues (Host Government—the Netherlands)

U.S. Delegate

Nega Beru, PhD, Director, Office of Plant and Dairy Foods, (HFS–300), Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5100 Paint Branch Parkway, College Park, MD 20740, Phone: (301) 436–1700, Fax: (301) 436–2651, E-mail: Nega.Beru@fda.hhs.gov.

Alternate Delegate

Kerry Dearfield, PhD, Scientific Advisor for Risk Assessment, Office of Public Health Science, Food Safety and Inspection Service, U.S. Department of Agriculture, 1400 Independence Ave. SW., Room 380, Aerospace Center, Washington, DC 20250, Phone: (202) 690–6451, Fax: (202) 690–6337, E-mail: Kerry.Dearfield@fsis.usda.gov.

Codex Committee on Pesticide Residues (Host Government—China)

U.S. Delegate

Lois Rossi, Director of Registration Division, Office of Pesticide Programs, U.S. Environmental Protection Agency, Ariel Rios Building, 1200 Pennsylvania Avenue, N.W., Washington, DC 20460, Phone: (703) 305–5035, Fax: (703) 305–5147, E-mail: rossi.lois@epa.gov.
Alternate Delegate

Robert Epstein, PhD, Associate Deputy Administrator, Science and Technology, Agricultural Marketing Service, USDA, P.O. Box 96456, Room 35225, Mail Stop 0222, 1400 Independence Avenue, SW., Washington, DC 20250, Phone: (202) 720–2138, Fax: (202) 720–1484, E-mail: robert.epstein@usda.gov.

Codex Committee on Methods of Analysis and Sampling (Host Government—Hungary)

U.S. Delegate

Gregory Diachenko, PhD, Director, Division of Product Manufacture and Use, Office of Premarket Approval, Center for Food Safety and Applied Nutrition (CFSAN), FDA (HFS–300), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2387, Fax: (301) 436–2364, E-mail: gregory.diachenko@fda.hhs.gov.

Alternate Delegate

Donald C. Kendall, Technical Services Division, Grain, Inspection, Packers & Stockyards Administration, U.S. Department of Agriculture, 10383 N. Ambassador Drive, Kansas City, MO 64153–1394, Phone: (816) 891–0463, Fax: (816) 891–0478, E-mail: donnald.c.kendall@usda.gov.

Codex Committee on Food Import and Export Inspection and Certification Systems (Host Government—Australia)

U.S. Delegate

Catherine Carnevale, D.V.M, Director, International Affairs Staff, Center for Food Safety and Applied Nutrition, FDA (HFS–300), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2380, Fax: (301) 436–2612, E-mail: catherine.carnevale@fda.hhs.gov.

Alternate Delegate

Mary Stanley, Director, Office of International Affairs, Food Safety and Inspection Service, U.S. Department of Agriculture, Room 2147–South Building, 1400 Independence Ave., SW., Washington, DC 20250, Phone: (202) 720–0287, Fax: (202) 720–6050, E-mail: Mary.Stanley@fsis.usda.gov.

Codex Committee on General Principles (Host Government—France)

U.S. Delegate

Note: A member of the Steering Committee heads the delegation to meetings of the General Principles Committee.

Codex Committee on Food Labeling (Host Government—Canada)

U.S. Delegate

Barbara O. Schneeman, PhD, Director, Office of Nutritional Products, Labelling and Dietary Uses, Center for Food Safety and Applied Nutrition, FDA, 5100 Paint Branch Parkway (HFS–800), College Park, MD 20740, Phone: (301) 436–2373, Fax: (301) 436–2636, E-mail: barbara.schneeman@fda.hhs.gov.

Alternate Delegate

Robert Post, PhD, Director, Labeling and Consumer Protection Staff, Food Safety and Inspection Service, USDA, 1400 Independence Avenue, SW. (602 Annex), Washington, DC 20250, Phone: (202) 205–0279, Fax: (202) 205–3625, E-mail: Robert.post@fsis.usda.gov.

Codex Committee on Food Hygiene (Host Government—United States)

U.S. Delegate

Robert L. Buchanan, PhD, Lead Scientist, Food Safety Initiative, Center for Food Safety and Applied Nutrition, FDA (HFS–006), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2309, Fax: (301) 436–2360, E-mail: robert.buchanan@fda.hhs.gov.

Alternate Delegates

Daniel Engeljohn, PhD, Deputy Assistant Administrator, Office of Policy, Program, and Employee Development, Food Safety and Inspection Service, USDA, Room 350–E, Jamie L. Whitten Building, 1400 Independence Avenue, SW., Washington, DC 20250, Phone: (202) 205–0405, Fax: (202) 401–1760, E-mail: daniel.engeljohn@fsis.usda.gov.

Rebecca Buckner, PhD, Consumer Safety Officer, Center for Food Safety and Applied Nutrition, FDA, Room 3B–0033, Harvey Wiley Building, 5100 Paint Branch Parkway, College Park, MD 10740, Phone: (301) 436–1486, Fax: (301) 436–2632, E-mail: rebecca.buckner@fda.hhs.gov.

Codex Committee on Nutrition and Food for Special Dietary Uses (Host Government—Germany)

U.S. Delegate

Barbara O. Schneeman, PhD, Director, Office of Nutritional Products, Labelling and Dietary Uses, Center for Food Safety and Applied Nutrition, FDA, 5100 Paint Branch Highway (HFS–800), College Park, MD 20740, Phone: (301) 436–2373, Fax: (301) 436–2636, E-mail: barbara.schneeman@fda.hhs.gov.

Alternate Delegate

Allison Yates, PhD, Director, Beltsville Human Nutrition Research Center, Agricultural Research Service, U.S. Department of Agriculture, 10300 Baltimore Avenue, Bldg 307C, Room 117, Beltsville, MD 20705, Phone: (301) 504–8157, Fax: (301) 504–9381, E-mail: Allison.Yates@ars.usda.gov.

Worldwide Commodity Codex Committees Codex Committee on Fresh Fruits and Vegetables (Host Government—Mexico)

U.S. Delegate

Dorian LaFond, International Standards Coordinator, Fruit and Vegetables Program, Agricultural Marketing Service, USDA, Room 2086, South Building, 1400 Independence Avenue, SW., Washington, DC 20250, Phone: (202) 690–4944, Fax: (202) 720–4722, E-mail: dorian.lafond@usda.gov.

Alternate Delegate

Michelle Smith, PhD, Interdisciplinary Scientist, Office of Plant and Dairy Foods, Center for Food Safety and Applied Nutrition, Food and Drug Administration (HFS–306), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2024, Fax: (301) 436–2651, E-mail: Michelle.Smith@fda.hhs.gov.

Codex Committee on Fish and Fishery Products (Host Government—Norway)

U.S. Delegate

Donald Kraemer, Acting Director, Office of Seafood, Center for Food Safety and Applied Nutrition, Food and Drug Administration, Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2300, Fax: (301) 436–2599, E-mail: donald.kraemer@fda.hhs.gov.

Alternate Delegate

Timothy Hansen, Director, Seafood Inspection Program, National Oceanic and Atmospheric Administration, Department of Commerce, Room 10837, 1315 East West Highway, Silver Spring, MD 20910, Phone: (301) 713–2355, Fax: (301) 713–1081 E-mail: Timothy.Hansen@noaa.gov.

Codex Committee on Cereals, Pulses and Legumes (Host Government—United States)

U.S. Delegate

Henry Kim, PhD, Supervisory Chemist, Division of Plant Product Safety,
Office of Plant and Dairy Foods, Center for Food Safety and Applied Nutrition, FDA, 5100 Paint Branch Parkway, College Park, MD 20740, Phone: (301) 436–2023, Fax: (301) 436–2651, E-mail: henry.kim@fda.hhs.gov.

Codex Committee on Milk and Milk Products (Host Government—New Zealand)

U.S. Delegate

Duane Spomer, Food Defense Advisor, Agricultural Marketing Service, U.S. Department of Agriculture, Room 2750, South Building, 1400 Independence Ave., SW., Washington, DC 20250, Phone: (202) 720–1861, Fax: (202) 205–5772, E-mail: duane.spomer@usda.gov.

Alternate Delegate

John F. Sheehan, Director, Division of Dairy and Egg Safety, Office of Plant and Dairy Foods and Beverages, Center for Food Safety and Applied Nutrition, FDA (HFS–306), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740, Phone: (301) 436–1488, Fax: (301) 436–2632, E-mail: john.shee@fda.hhs.gov.

Codex Committee on Fats and Oils (Host Government—United Kingdom)

U.S. Delegate

Dennis M. Keefe, PhD, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition, FDA (HFS–200), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–1284, Fax: (301) 436–2972, E-mail: dennis.keefe@fda.hhs.gov.

Alternate Delegate

Kathleen Warner, Agricultural Research Service, USDA, 1815 N. University Street, Peoria, IL 61604, Phone: (309) 681–6584, Fax: (309) 681–6688, E-mail: warnerk@ncaur.usda.gov.

Codex Committee on Cocoa Products and Chocolate (Host Government—Switzerland)

U.S. Delegate

Michelle Smith, PhD, Food Technologist, Office of Plant and Dairy Foods and Beverages, Center for Food Safety and Applied Nutrition, FDA (HFS–306), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–2024, Fax: (301) 436–2651, E-mail: michelle.smith@fda.hhs.gov.

Codex Committee on Sugars (Host Government—United Kingdom)

U.S. Delegate

Martin Stutsmann, J.D., Office of Plant and Dairy Foods and Beverages, Center for Food Safety and Applied Nutrition, FDA (HFS–306), Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–1642, Fax: (301) 436–2651, E-mail: martin.stutsmann@fda.hhs.gov.

Codex Committee on Processed Fruits and Vegetables (Host Government—United States)

U.S. Delegate

Dorian LaFond, International Standards Coordinator, Fruit and Vegetable Division, Agricultural Marketing Service, USDA, Room 2086, South Building, 1400 Independence Avenue, SW., Washington, DC 20250, Phone: (202) 690–4944, Fax: (202) 720–0016, E-mail: dorian.lafond@usda.gov.

Alternate Delegate

Paul South, PhD, Division of Plant Product Safety, Office of Plant and Dairy Foods, Center for Food Safety and Applied Nutrition, FDA, 5100 Paint Branch Parkway, College Park, MD 20740, Phone: (301) 436–1640, Fax: (301) 436–2561, E-mail: paul.south@fda.hhs.gov.

Codex Committee on Vegetable Proteins (Host Government—Canada)

U.S. Delegate

Dr. Wilda H. Martinez, Area Director, ARS North Atlantic Area, Agricultural Research Service, USDA, 600 E. Mermaid Lane, Wyndmoor, PA 19038, Phone: (215) 233–6593, Fax: (215) 233–6719, E-mail: wmartinez@ars.usda.gov.

Codex Committee on Meat Hygiene (Host Government—New Zealand)

U.S. Delegate

Perfeito Santiago, D.V.M., Deputy Assistant Administrator, Office of Food Security and Emergency Preparedness, Room 3130, South Building, Food Safety and Inspection Service, USDA, 1400 Independence Avenue, SW., Washington, DC 20250, Phone: (202) 205–0452, Fax: (202) 690–5634, E-mail: perfecto.santiago@fsis.usda.gov.

Codex Committee on Natural Mineral Waters (Host Government—Switzerland)

U.S. Delegate

Lauren Robin, PhD, Review Chemist, Office of Plant and Dairy Foods, Center for Food Safety and Applied Nutrition, Food and Drug Administration, Harvey W. Wiley Federal Building, 5100 Paint Branch Parkway, College Park, MD 20740–3835, Phone: (301) 436–1639, Fax: (301) 436–2651, E-mail: Lauren.Robin@fda.hhs.gov.

Ad Hoc Intergovernmental Task Forces

Ad Hoc Intergovernmental Task Force on Foods Derived From Modern Biotechnology (Host Government—Japan)

U.S. Delegate

Eric Flamn, PhD, Senior Advisor, Office of the Commissioner, Food and Drug Administration, Room 1561, Parklawn Building, Rockville, MD 20857, Phone: (301) 827–0591, Fax: (301) 827–4774, E-mail: EFLAMM@OC.FDA.GOV.

Alternate Delegate

Cindy Smith, Deputy Administrator, Biotechnology Regulatory Services, Animal and Plant Health Inspection Service, U.S. Department of Agriculture, Unit 98 Ste. SB05, 4700 River Road, Riverdale, MD 20737, Phone: (301) 734–7324, Fax: (301) 734–6352, E-mail: Cindy.J.Smith@aphis.usda.gov.

Ad Hoc Intergovernmental Task Force on Antimicrobial Resistance (Host Government—Republic of Korea)

Delegate

David G. White, D.V.M., Director, National Antimicrobial Resistance Monitoring System (NARMS), U.S. Food and Drug Administration, Center for Veterinary Medicine, Office of Research, 8401 Muirkirk Rd., Laurel, MD 20708, Phone: (301) 210–4181, Fax: (301) 210–4683, E-mail: David.White@fda.hhs.gov.

Alternate Delegate

Neena Anandaraman, D.V.M., Veterinary Medical Officer, Zoonotic Diseases & Residue Surveillance Division, Office of Public Health Science, Food Safety and Inspection Service, U.S. Department of Agriculture, Room 343, Aerospace Center, Washington, DC 20250, Phone: (202) 690–6429, Fax: (202) 690–6355, E-mail: neena.anandaraman@fsis.usda.gov.

Ad Hoc Intergovernmental Task Force on Quick Frozen Foods (Host Government—Thailand)

Delegate

Donald Zink, Ph.D., Senior Scientist, Office of Plant and Dairy Foods,
Coated Free Sheet Paper from Indonesia: Notice of Preliminary Determination of Sales at Less Than Fair Value and Postponement of Final Determination

AGENCY: Import Administration, International Trade Administration, Department of Commerce.

SUMMARY: The U.S. Department of Commerce (the Department) preliminarily determines that coated free sheet paper (CFS) from Indonesia is being, or is likely to be, sold in the United States at less than fair value (LTFV), as provided in section 733(b) of the Tariff Act of 1930, as amended (the Act). The estimated margins of sales at LTFV are listed in the “Suspension of Liquidation” section of this notice. Interested parties are invited to comment on this preliminary determination. Pursuant to requests from interested parties, we are postponing for 60 days the final determination and extending provisional measures from a four-month period to not more than six months. Accordingly, we will make our final determination not later than 135 days after publication of the preliminary determination.

EFFECTIVE DATE: June 4, 2007.

FOR FURTHER INFORMATION CONTACT: Brian Smith, AD/CVD Operations, Office 2, Import Administration, International Trade Administration, U.S. Department of Commerce, 14th Street and Constitution Avenue, NW, Washington, DC 20230; telephone (202) 482–1766.

SUPPLEMENTARY INFORMATION:

Background


The Department set aside a period of time for parties to raise issues regarding product coverage and encouraged all parties to submit comments within 20 calendar days of publication of the Initiation Notice. See Initiation Notice, 71 FR at 68538; see also Antidumping Duties; Countervailing Duties; Final Rule, 62 FR 27296, 27323 (May 19, 1997). On December 18, 2006, the two largest known producers exporters of CFS from Indonesia, PT. Fabrik Kertas Tjiwi Kimia Tbk. (TK) and PT. Pindo Deli Pulp and Paper Mills (PD), submitted timely comments, in which they requested that the Department exclude cast-coated CFS from the scope of the investigation.

On December 22, 2006, the United States International Trade Commission (ITC) preliminarily determined that there is a reasonable indication that imports of CFS from Indonesia, the People’s Republic of China (PRC), and the Republic of Korea (Korea) are materially injuring the U.S. industry and notified the Department of its findings. See Coated Free Sheet Paper from China, Indonesia, and Korea Investigation Nos. 701–TA–444–448 and 731–TA–1107–1109 (Preliminary), 71 FR 78464 (Dec. 29, 2006).

Also on December 22, 2006, we selected PD and TK as the mandatory respondents in this proceeding. See Memorandum from James Maeder, Office Director, to Stephen J. Claeys, Deputy Assistant Secretary, entitled: “Antidumping Duty Investigation of Coated Free Sheet Paper from Indonesia - Selection of Respondents,” dated December 22, 2006. We subsequently issued the antidumping questionnaire to these companies on December 22, 2006.

On January 12, 2007, the Department requested that PD and TK file their December 18, 2006, scope comments on the administrative record of the companion LTFV and countervailing duty (CVD) investigations of CFS from the PRC and Korea. See Memorandum from Alice Gibbons to The File, dated January 12, 2007. PD and TK did so on the same date.

On January 17, 2007, the petitioner made a country-wide allegation that sales of CFS in the home market were made below the cost of production (COP) during the period of investigation (POI).

On January 19, 2007, the petitioner objected to the respondents’ request to exclude cast-coated paper from the scope of the investigation. For further discussion, see the “Scope Comments” section of this notice, below.

On January 26, 2007, PD and TK submitted a consolidated response to section A of the questionnaire (i.e., the section involving general information). In this submission, PD and TK indicated that, not only are they affiliated with each other, but they are also affiliated with a third company that produces CFS in Indonesia, PT. Indah Kiat Pulp and Paper Tbk (IK). Based on an analysis of this information, as well as additional information obtained during the course of this proceeding (see below), we find that it is appropriate to treat these three companies as a single entity, hereinafter referred to as PD/TK. Nonetheless, we did not require PD/TK to report sales and cost data related to IK’s POI sales of CFS because: 1) these sales were made only in the home market; 2) the quantity of the sales was insignificant; and 3) these sales would not be the most similar matches to products sold in the United States by PD or TK. For further discussion, see the “Collapsing IK, PD, and TK” section of this notice, below.

On February 2, 2007, the Department initiated a country-wide sales-below-cost investigation to determine whether PD/TK’s sales of CFS in the home market were made at prices below the COP during the POI. See the Memorandum from The Team to James Maeder, Office Director, Office 2, Office of AD/CVD Operations, entitled, “The Petitioner’s Allegation of Country-Wide Sales Below the Cost of Production” (Below–Cost Allegation), dated February 2, 2007. On February 5, 2007, the Department instructed PD/TK to respond to section D of the questionnaire with respect to its home market sales of CFS in order to acquire the necessary information to determine whether such sales were made at prices below the companies’ COP.