CODEX ALIMENTARIUS COMMISSION ${f E}$





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TO: Codex Contact Points

Interested International Organizations

FROM: Secretariat, Codex Alimentarius Commission, Joint FAO/WHO Food Standards Programme

SUBJECT: Distribution of the Report of the 33rd Session of the Codex Committee on Nutrition and

Foods for Special Dietary Uses (REP12/NFSDU)

A. MATTERS FOR ADOPTION BY THE 35th SESSION OF THE COMMISSION:

Draft Guidelines Step 5/8 of the Procedure

1. Proposed Draft Nutrient Reference Values (NRVs) (para. 76, Appendix III).

Governments and interested international organizations wishing to comments on the above document, should do so in writing, in conformity with the *Procedure for the Elaboration of Codex Standards and Related Texts* (Procedural Manual of the Codex Alimentarius Commission), to the above address, before **15 March 2012**.

Draft Guidelines Step 5 of the Procedure

2. Proposed Draft Revision of the Guidelines on Formulated Supplementary Foods for Older Infants and Young Children (CAC/GL 8-1991) (para. 126, Appendix IV).

Governments and interested international organizations wishing to comments on the above document, should do so in writing, in conformity with the *Procedure for the Elaboration of Codex Standards and Related Texts* (Procedural Manual of the Codex Alimentarius Commission), to the above address, before 15 March 2012.

B. REQUEST FOR COMMENTS AND INFORMATION

Proposed Draft Guidelines at Step 3 of the Procedure

3. General Principles for Establishing Nutrient Reference Values for Nutrients Associated with Risk of Diet-Related Non-communicable Diseases for General Population (NRVs-NCD) (para. 66, Appendix V)

Governments and interested international organizations wishing to submit comments on the above document, should do so by writing preferably by email to Dr Barbara O. Schneeman, Director, Office of Nutrition Labeling and Dietary Supplements, Center for Food Safety & Applied Nutrition, U.S. Food and Drug Administration (HFS-800), 5100 Paint Branch Parkway, College Park, MD 20740, United States of America, E-Mail:barbara.schneeman@fda.hhs.gov, with a copy to the Secretariat at the address above before 1 February 2012.