

specifically at ex-offenders, an additional supplementary module will be administered by Audio-CASI. Similarly, an additional supplementary module will be administered by Audio-CASI in the site operating a program aimed at survey respondents with

mental health problems. Finally, in the two-generation sites (two of the six sites), survey respondents will complete a two-generation survey administered by a Computer Assisted Personal Interview (CAPI). Approximately 12,000 respondents will complete the core

survey, 2,000 will complete the criminal justice module, 2,000 will complete the mental health module, and 4,000 will complete the two-generation CAPI survey.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Audio-CASI Core	12,000	1	10 minutes or .17 hrs	2,000
Criminal Justice Module	2,000	1	10 minutes or .17 hrs	333.33
Mental Health Module	2,000	1	10 minutes or .17 hrs	333.33
Two Generation	4,000	1	30 minutes or .5 hrs	2,000
Estimated Total Annual Burden Hours	4,666.66

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: February 6, 2003.

Gerald L. Fralick,

Director, Office of Information Systems.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 03F-0023]

Kerry, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Kerry, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of gum arabic as a thickener, emulsifier, or stabilizer in the manufacture of creamers for use in alcoholic beverages at a maximum level of use of 20 percent.

FOR FURTHER INFORMATION CONTACT:

Andrew D. Laumbach, Center for Food Safety and Applied Nutrition (S-265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 202-418-3071.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 1A4730) has been filed by Kerry, Inc., c/o Bell, Boyd, and Lloyd LLC, Three First National Plaza, 70 West Madison St., suite 3300, Chicago, IL 60602-4207. The petition proposes to amend the food additive regulations in part 172 *Food Additives Permitted for Direct Addition to Food for Human Consumption* (21 CFR part 172) to provide for the safe use of gum arabic as a thickener, emulsifier, or stabilizer in the manufacture of creamers for use in alcoholic beverages at a maximum level of use of 20 percent.

The food additive petition filed as FAP 1A4730 was initially filed as a generally recognized as safe (GRAS) affirmation petition GRP 3G0287 as announced in a notice that was published in the **Federal Register** of October 13, 1983 (48 FR 46626) (The GRAS affirmation petition was filed by Beatrice Foods Co., now Kerry, Inc.). Kerry, Inc., requested in a letter dated September 6, 2001, that FDA convert the GRAS affirmation petition (GRP 3G0287) to a food additive petition (FAP 1A4730).

The agency has determined under 21 CFR 25.32(r) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment.

Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: December 19, 2002.

Alan M. Rulis,

Director, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition.

[FR Doc. 03-3557 Filed 2-12-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00E-1249]

Determination of Regulatory Review Period for Purposes of Patent Extension; Avandia

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for