is used to locate noncustodial parents and to verify income and employment.

Respondents: State IV–D programs.

ANNUAL BURDEN ESTIMATES

Reporting	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
States	12	12	2	288

Estimated Total Annual Burden: 288.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Washington, DC 20503, Attn: Ms. Laura Oliven.

Dated: October 25, 2001.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 01–27351 Filed 10–30–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Announcement of Financial Assistance to Expand Head Start Enrollment to Unserved Communities.

AGENCY: Administration on Children, Youth and Families, ACF, DHHS.

ACTION: Correction.

SUMMARY: This document contains a correction to the Notice that was published in the **Federal Register** on Monday, September 24, 2001.

In Appendix A, add Bloomfield County, Colorado to the list of geographic areas currently unserved by Head Start grantees, and delete Lemhi County, Idaho because it is a served community.

FOR FURTHER INFORMATION CONTACT: The ACYF Operations Center at 1815 N. Fort Myer Drive, Suite 300, Arlington, VA 22209 or telephone: 1–(800)–351–2293, or E. Mail to: *ehs@lognet.com.*

Dated: October 25, 2001.

James A. Harrell,

Acting Commissioner, Administration on Children, Youth, and Families.

[FR Doc. 01–27375 Filed 10–30–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00D–1305]

Compliance Policy Guide: ''Apple Juice, Apple Juice Concentrates, and Apple Juice Products—Adulteration with Patulin,'' Availability; and ''Patulin in Apple Juice, Apple Juice Concentrates and Apple Juice Products,'' Supporting Document; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a compliance policy guide (CPG) entitled ''Apple Juice, Apple Juice Concentrates, and Apple Juice Products—Adulteration with Patulin.'' This document is intended to make FDA offices and the industry aware of FDA's guidance for enforcement concerning apple juice, apple juice concentrates, and apple juice products that contain patulin, a toxic substance produced by molds that may grow on apples and that has been found to occur at high levels in some apple juice products offered for sale in the United States. The agency also is announcing the availability of a document entitled ''Patulin in Apple Juice, Apple Juice Concentrates and Apple Juice Products'' (final supporting document).

DATES: Submit written or electronic comments on the CPG or the final supporting document at any time.

ADDRESSES: Submit written requests for single copies of the CPG entitled ''Apple Juice, Apple Juice Concentrates, and Apple Juice Containing Products—Adulteration with Patulin'' and/or the final supporting document entitled ''Patulin in Apple Juice, Apple Juice Concentrates and Apple Juice Products'' to the Office of Plant and Dairy Foods and Beverages (HFS–305), Center for Food Safety and Applied Nutrition, Food and Drug Administration, 200 C St. SW., Washington, DC 20204. Send two self-addressed adhesive labels to assist that office in processing your request. Submit written comments to the Docket Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments. See the **SUPPLEMENTARY INFORMATION** section for electronic access to these documents.

FOR FURTHER INFORMATION CONTACT:

Technical questions concerning patulin in apple juice products: Michael E. Kashtock, Center for Food Safety and Applied Nutrition (HFS–305), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–5321, FAX 202–205–4422, e-mail: mkashtoc@cfsan.fda.gov.

Questions concerning regulatory actions: MaryLynn Datoc, Office of Enforcement (HFC–230), Office of Regulatory Affairs, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–0413, FAX 301–827–0482, e-mail: mdatoc@ora.fda.gov.

SUPPLEMENTARY INFORMATION:

I. Background

In a notice published in the **Federal Register** of June 16, 2000 (65 FR 37791), FDA announced the availability of a draft CPG entitled ''Apple Juice, Apple Juice Concentrates, and Apple Juice Products—Adulteration with Patulin'' and a draft supporting document entitled ''Patulin in Apple Juice, Apple Juice Concentrates, and Apple Juice Products.'' The agency has finalized the draft CPG and the draft supporting document after considering the

comments we received on these documents.

FDA received four comments on these two documents, three from food industry associations and one from an organization representing several foreign governments. Three of the comments supported the draft CPG and draft supporting document and did not raise any questions.

The fourth comment posed three questions about the action level and the safety assessment described in the draft supporting document. FDA has considered these questions and has responded to them in the revised supporting document.

II. Significance of Guidance

This CPG is being issued as guidance consistent with FDA's good guidance practices regulation in 21 CFR 10.115. The CPG represents the agency's current thinking on its enforcement process concerning the adulteration of apple juice, apple juice concentrates, and apple juice products with patulin. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute and regulations.

III. Comments

Interested persons may, at any time, submit to the Docket Management Branch (address above) written or electronic comments concerning the CPG entitled ''Apple Juice, Apple Juice Concentrates, and Apple Juice Products—Adulteration with Patulin'' or the final supporting document. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Comments also may be submitted electronically. A copy of the CPG and the final supporting document and received comments are available for public examination in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

IV. Electronic Access

Persons with access to the Internet may obtain the CPG at http://www.fda.gov/ora under ''Compliance References.'' The supporting document may be accessed at http://www.cfsan.fda.gov under the heading ''How to Obtain FDA Food & Cosmetic Guidance Documents.''

Dated: October 22, 2001.

Dennis E. Baker,,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 01–27319 Filed 10–30–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources And Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443–1129.

Comments are invited 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) ways to enhance 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.

Proposed Project: The National Health Service Corps Uniform Data System (OMB No. 0915–0232): Extension

The National Health Service Corps (NHSC) of the Bureau of Health Professions (BHPr), Health Resources and Services Administration (HRSA), is committed to improving the health of the Nation's underserved by uniting communities in need with caring health professionals and by supporting communities' efforts to build better systems of care.

The NHSC needs to collect data on its programs to ensure compliance with legislative mandates and to report to Congress and policymakers on program accomplishments. To meet these objectives, the NHSC requires a core set of information collected annually that is appropriate for monitoring and evaluating performance and reporting on annual trends. The following information will be collected from each site: services offered and delivery method; users by various characteristics; staffing and utilization; charges and collections; receivables, income, and expenses; and managed care.

The estimated burden is as follows:

Type of report	Number of respondents	Responses per respondent	Hours per response	Total burden hours
Universal Report	620	1	27	16,740

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 14–22, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: October 25, 2001.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 01–27366 Filed 10–30–01; 8:45 am]

BILLING CODE 4515–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA)