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)