

jarring/jolting-related injuries. Questions will be asked of survey participants regarding health and work history with respect to farm equipment operation. The specific data will include risk factors (both on and off the job) and outcome (the prevalence of symptoms in various body parts). The National Education Center for Agricultural Safety

(NECAS) will assist in administering the questionnaire to survey participants at the American Farm Bureau Federation Annual Meeting in Tampa, FL, in January 19–20, 2003. Since the conference is well attended, researchers expect 10–12 percent of the meeting attendees to participate in the study. Respondents will complete the survey

questionnaire that includes 80 questions. Based on prior experience with a similar questionnaire, the anticipated time for a participant to complete the questionnaire is 20 minutes or less. There will be no cost to respondents.

Respondents	Number of respondents	Number of responses/respondent	Avg. burden/response (in hours)	Total burden (in hours)
Attendees at the American Farm Bureau Federation's Annual Meeting	600	1	20/60	200
Total				200

Dated: September 17, 2002.
Nancy E. Cheal,
Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0393]

Assessing Acrylamide in the U.S. Food Supply; Public Meeting; Draft Action Plan on Acrylamide; Availability; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting and availability; correction.

SUMMARY: The Food and Drug Administration is correcting a notice that appeared in the *Federal Register* of September 12, 2002. The document announced a public meeting entitled “Assessing Acrylamide in the U.S. Food Supply.” The document was published with an incorrect Internet address for an analytical test methodology to measure acrylamide levels. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Doris B. Tucker, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7626.

SUPPLEMENTARY INFORMATION: In FR Doc. 02–23193, appearing on page 57827 in the *Federal Register* of Thursday, September 12, 2002, the following correction is made:

1. On page 57827, in the second column, in the last paragraph, beginning on line 7, the Internet address is corrected to read “<http://www.cfsan.fda.gov/~dms/acrylami.html>”.

Dated: September 18, 2002.
Margaret M. Dotzel,
Associate Commissioner for Policy.
 [FR Doc. 02–24201 Filed 9–23–02; 8:45 am]
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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: Uncompensated Services Assurance Report (OMB No. 0915–0077)—Revision

Under the Hill-Burton Act, the Government provides grants and loans for construction or renovation of health care facilities. As a condition of receiving this construction assistance, facilities are required to provide services to persons unable to pay. A condition of receiving this assistance requires facilities to provide assurances periodically that the required level of uncompensated care is being provided, and that certain notification and record keeping procedures are being followed. These requirements are referred to as the uncompensated services assurance.

Estimate of Information Collection Burden

Type of requirement and regulatory citation	No. of respondents	Responses per respondent	Total responses	Hours per response	Total hour burden
Disclosure Burden (42 CFR)					
Published Notices (124.504(c))	216	1	216	0.75	162
Individual Notices (124.504(c))	216	1	216	43.6	9,418