hearing aid dispenser locating the appropriate User Instructional Brochure for the specific model and mailing the brochure to the requester.

Section 801.421(d) recordkeeping estimate assumes that 9,900 hearing aid dispensers will each retain 162 records. Each record documents the dispensing of a hearing aid to a hearing aid user. Each recordkeeping entry is estimated to require 0.25 staff hours.

Dated: June 19, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–17289 Filed 6–29–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0430]

Nalco Chemical Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Nalco Chemical Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of sodium acrylate/styrene sulfonate copolymer for use as an antiscalant boiler treatment where steam from treated boilers may contact food.

DATES: Written comments on the petitioner's environmental assessment by July 30, 1998.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Parvin M. Yasaei, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3189.

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 8A4598) has been filed by Nalco Chemical Co., One Nalco Center, Naperville, IL 60563. The petition proposes to amend the food additive regulations in § 173.310 *Boiler water additives* (21 CFR 173.310) to provide for the safe use of sodium acrylate/sulfonated styrene copolymer for use as an antiscalant boiler treatment where steam from treated boilers may contact food.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before July 30, 1998, submit to the Dockets Management Branch (address above) written comments. 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. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.51(b)(1).

Dated: June 8, 1998.

Laura M. Tarantino,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–17292 Filed 6–29–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0433]

Servo Delden BV; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Servo Delden BV has filed a petition proposing that the food additive regulations be amended to provide for the safe use of polyethylene glycol mono-isotridecyl ether sulfate, sodium salt as a surfactant in adhesives intended for use in contact with food.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

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 8B4600) has been filed by Servo Delden BV, c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes to amend the food additive regulations to provide for the safe use of polyethylene glycol mono-isotridecyl ether sulfate, sodium salt as a surfactant in adhesives intended for use in contact with food.

The agency has determined under 21 CFR 25.32(i) that this action is of the 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: June 4, 1998.

Laura M. Tarantino,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–17321 Filed 6–29–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Center for Veterinary Medicine; Change of Internet Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing a change in the Internet address for the Center for Veterinary Medicine (CVM) to ensure that users can continue to have uninterrupted access to CVM's Internet site. The CVM Internet site is one of the agency's methods of communicating with the public regarding the ongoing mission of CVM, which is the organization within the agency that regulates the manufacture and distribution of animal drugs, feeds, and related products.

FOR FURTHER INFORMATION CONTACT: Jerome J. McDonald, Center for Veterinary Medicine (HFV–16), Food and Drug Administration, 7500 Standish

Pl., Rockville, MD 20855, 301–827–6505.

SUPPLEMENTARY INFORMATION: On May 17, 1998, CVM changed its Internet address to ''www.fda.gov/cvm''. If users enter the old CVM internet address (''www.cvm.fda.gov''), a message will notify them of the change and will automatically redirect them to the new address. The message redirecting users to the new Internet site (redirect) will be in effect for approximately 3 months. CVM recommends that users who have bookmarked pages within the CVM website (e.g., What's New, the On-line Library, CVM's telephone directory, etc.) update the bookmarks with the new CVM Internet address. If users do not update the bookmarks, an error message will result and no redirect instructions will be provided.

Dated: June 17, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–17320 Filed 6–29–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service

Request for Public Comment: 30-day Proposed Collection: Common Reporting Requirements for Urban Indian Health Programs

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, which requires opportunity for public comment on proposed information collection projects, the Indian Health Service (IHS) has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection project was published in the March 10, 1998, **Federal Register** (63 FR 11688) and allowed 60 days of public comment. No public comment was received in response to the notice. The purpose of this notice is to allow 30 days for public comment to be submitted to OMB.

PROPOSED COLLECTION: *Title:* 09–17–0007, ''Common Reporting Requirements for Urban Indian Health Programs.'' *Type of Information Collection Request:* Revision of currently approved information collection, 0917–0007, ''Common Reporting Requirements for Urban Indian Health Programs,'' which expires July 31, 1998. *Form Number:* Reporting forms contained in IHS instruction manual, ''Urban Indian Health Programs Common Reporting Requirements.''

Need and Use of Information Collection: The requested information is provided by American Indian/Alaska Native (AI/AN) urban health organizations contracting with the IHS to provide health care services to AI/ANs in urban settings. The information is collected annually and is used to monitor and evaluate contractor performance, prepare budget reports, and allocate resources.

Affected Public: Businesses or other for-profit, individuals and households, not-for-profit institutions; and State, Local, or Tribal Governments. *Type of Respondents:* Urban Indian health care organizations.

The table below provides the following: types of data collection instruments, estimated number of respondents, number of responses per respondent, average burden hour per response, and total annual burden hour.

Data collection instrument	Estimated number of respondents	Responses per respondent	Annual number of responses	Average burden hour per response*	Total annual burden hours
Face Sheet	34	1	34	0.50 (30 mins)	17.0
Table 1	34	1	34	2.00 (120 mins)	68.0
Table 2	34	1	34	0.75 (45 mins)	26.0
Table 3	34	1	34	2.25 (135 mins)	77.0
Table 4	**24	1	23	0.50 (30 mins)	12.0
Table 5	34	1	34	2.00 (120 mins)	68.0
Table 6	34	1	34	2.00 (120 mins)	68.0
Table 7	34	1	34	1.00 (60 mins)	34.0
Table 8	34	1	34	1.25 (75 mins)	43.0
Total	261		261		413.0

*For ease of understanding, burden hours are also provided in actual minutes.
**Excludes urban Indian health projects with no medical components.

There are no capital costs, operating costs, or maintenance costs to report.

Request for Comments

Your written comments and/or suggestions are invited on one or more of the following points: (1) Whether the information collection activity is necessary to carry out an agency function; (b) whether the IHS processes the information collected in a useful and timely fashion; (c) the accuracy of the public burden estimate (the estimated amount of time needed for individual respondents to provide the requested information); (d) whether the methodology and assumptions used to determine the estimate are logical; (e) ways to enhance the quality, utility, and clarity of the information being collected; and (f) ways to minimize the public burden through the use of automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Send your written comments and suggestions regarding the proposed information collection contained in this notice, especially regarding the estimated public burden and associated response time, to: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, D.C. 20503, Attention: Desk Officer for IHS.

To request more information on the proposed collection or to obtain a copy of the data collection instrument(s) and/or instruction(s), contact: Mr. Lance Hodahkwen, Sr., M.P.H., IHS Reports Clearance Officer, 12300 Twinbook Parkway, Suite 450, Rockville, MD 20852–1601; call non-Toll free at (301) 443–1116; send via fax to (301) 443–1522; or send your e-mail request, comments, and return address to: Ihodahkw@hqe.ihs.gov.

Comment Due Date: Comments regarding this information are best assured to having their full effect if received on or before July 30, 1998.