comments and suggestions submitted within 60 days of this publication.

Dated: October 28, 1996.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 96–28139 Filed 11–1–96; 8:45 am]

BILLING CODE 4184–01–M

Submission for OMB Review; Comment Request

Title: Federal Parent Locator Service.
OMB No.: 0970–0142.

Description: The Office of Child support Enforcement (OCSE) operates the Federal Parent Locator Services (FPLS), a computerized national location network which provides address and social security number information to State and local child support enforcement agencies upon request to locate parents in order to establish or enforce a child support order and to assist authorized persons in resolving parental kidnapping and child custody cases.

State and local agency requests to the FPLS can be made by tape, cartridge, electronic file transfer or by dialing-up using a personal computer. The FPLS serves as a conduit between child support enforcement offices and Federal and State agencies by conducting weekly, biweekly, or monthly matches of the collected information with various agencies and distributing the information back to the requesting State or local child support office.

Respondents: State, Local, Tribal or Federal Govt. Governments.

ANNUAL BURDEN ESTIMATE

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Standard Forms	200	24	1	4,800
Estimated Total Annual Burden Hours:				4,800

Explanation

*The specific number of annual burden hours per respondent will vary depending on individual circumstance including a States' frequency in submitting requests and their mode of submission.

*Burden hour for initial collection of information included in the submission are not considered as part of their day-to-day operation of the child support enforcement program.

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, S.W., Washington, D.C. 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, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

Dated: October 28, 1996.

Douglas J. Godesky,

Reports Clearance Officer.

[FR Doc. 96–28140 Filed 11–1–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0298]

Agency Information Collection Activities: Proposed Collection; Comment Request; Extension

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995, Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the voluntary collection of information for the Medical Devices Standards Activities Report, a comprehensive listing of current national and international standards for medical devices.

DATES: Submit written comments on the collection of information by January 3, 1997.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c). To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Medical Devices Standards Activities Report (OMB Control Number 0910–0219—Extension)

FDA is collecting information necessary to update a comprehensive listing of current national and international standards activities in the field of medical devices. The collection of this information is authorized by section 514(a)(4)(B) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360d(a)(4)(B)), which requires FDA to consult with other nationally or internationally recognized standard-setting entities, including other Federal agencies concerned with standard-setting, in carrying out its responsibility to establish special controls for medical devices. This report is used by approximately 39 standards-developing organizations to coordinate their standards activities. This coordination prevents duplication of effort and insures efficient and expeditious management of standards development. Over 700 copies of this report are used by government, hospitals, libraries, industry, private citizens, and State and local government agencies, including FDA, to keep abreast of standards development activities and current technology concerning the safety of medical devices. Without the report, there would be duplication of standards efforts by voluntary standards organizations since there is no other publication that can be easily referenced to ascertain if a certain medical device standard is being or has been developed.

FDA estimates the burden of this collection of information as follows:

ESTIMATED ANNUAL REPORTING BURDEN

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
39	.5	19.5	3	58.5

There are no capital costs or operating and maintenance costs associated with this collection of information.

This collection occurs biennially and is voluntary. There are 39 national and international organizations with one report each reporting period.

Dated: October 29, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–28209 Filed 11–1–96; 8:45 am]

BILLING CODE 4160–01–F

[Docket No. 93F–0273]

Lonza, Inc.; Withdrawal of Food Additive Petition; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the Federal Register of August 12, 1996 (61 FR 41793). The document announced the withdrawal of a food additive petition (FAP 3B4392) proposing that the food additive regulations be amended to provide for the safe use of didecyldimethylammonium chloride as a slimicide used in the manufacture of paper and paperboard intended to contact food. The document was published with an error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Andrew J. Zajac, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3095.

In FR Doc. 96–20437, appearing on page 41793 in the Federal Register of Monday, August 12, 1996, the following correction is made: On page 41793, in the first column, in the first line, ''[Docket No. 93F–0269]'' is corrected to read ''[Docket No. 93F–0273]''.

Dated: October 16, 1996.

Alan M. Rulis,

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

[FR Doc. 96–28210 Filed 11–1–96; 8:45 am]

BILLING CODE 4160–01–F

Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). This notice also summarizes the procedures for the meeting and methods by which interested persons may participate in open public hearings before FDA's advisory committees.

FDA has established an Advisory Committee Information Hotline (the hotline) using a voice-mail telephone system. The hotline provides the public with access to the most current information on FDA advisory committee meetings. The advisory committee hotline, which will disseminate current information and information updates, can be accessed by dialing 1–800–741–8138 or 301–443–0572. Each advisory committee is assigned a 5-digit number. This 5-digit number will appear in each individual notice of meeting. The hotline will enable the public to obtain information about a particular advisory committee by using the committee's 5-digit number. Information in the hotline is preliminary and may change before a meeting is actually held. The hotline will be updated when such changes are made.

MEETING: The following advisory committee meeting is announced:

Orthopedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee

Date, time, and place. November 20, 1996, 10 a.m., and November 21, 1996, 8 a.m., Gaithersburg Hilton, Ballroom Salons A, B, and C, 620 Perry Pkwy., Gaithersburg, MD. A limited number of overnight accommodations have been reserved at the Holiday Inn—Gaithersburg, Two Montgomery Village Ave., Gaithersburg, MD. Attendees requiring overnight accommodations may contact the hotel at 301–948–8900 and reference FDA's Panel meeting block. Reservations will be confirmed at the group rate based on availability. Attendees with a disability requiring special accommodations should contact Sue Bae, KRA Corp., 301–495–1591, ext. 227. The availability of appropriate accommodations cannot be assured unless prior notification is received.

Type of meeting and contact person. Closed committee deliberations, November 20, 1996, 10 a.m. to 11:30 a.m.; open public hearing, 11:30 a.m. to 12:30 p.m., unless public participation does not last that long; open committee discussion, 12:30 p.m. to 6 p.m.; open committee discussion, November 21, 1996, 8 a.m. to 1:30 p.m.; Jodi H. Nashman, Center for Devices and Radiological Health (HFZ–410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2036, or FDA Advisory Committee Information Hotline, 1–800–