1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (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, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

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. New Dietary Ingredient Premarket Notification—21 CFR 190.6 (OMB Control No. 0910–0330)—Extension

Section 413(a) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 350b(a)) provides that a manufacturer or distributor of dietary supplements or of a new dietary ingredient is to submit information to FDA (as delegate for the Secretary of Health and Human Services) upon which it has based its conclusion that a dietary supplement containing a new dietary ingredient will reasonably be expected to be safe at least 75 days before the introduction or delivery for introduction into interstate commerce of a dietary supplement that contains a new dietary ingredient. FDA's regulations at part 190, subpart B (21 CFR part 190, subpart B) implement these statutory provisions. Section 190.6(a) requires each manufacturer or distributor of a dietary supplement containing a new dietary ingredient, or of a new dietary ingredient, to submit to the Office of Nutritional Products, Labeling, and Dietary Supplements notification of the basis for their conclusion that said supplement or ingredient will reasonably be expected to be safe. Section 190.6(b) requires that the notification include: (1) The complete name and address of the manufacturer or distributor, (2) the name of the new dietary ingredient, (3) a description of the dietary supplements that contain the new dietary ingredient, and (4) the history of use or other evidence of safety establishing that the dietary ingredient will reasonably be expected to be safe.

The notification requirements described previously are designed to enable FDA to monitor the introduction into the food supply of new dietary ingredients and dietary supplements that contain new dietary ingredients, in order to protect consumers from unsafe dietary supplements. FDA uses the information collected under these regulations to help ensure that a manufacturer or distributor of a dietary supplement containing a new dietary ingredient is in full compliance with the act.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN [1]

21 CFR Section	No. of respondents	Annual frequency per response	Total annual responses	Hours per response	Total hours
190.6	35	1	35	20	700

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

The agency believes that there will be minimal burden on the industry to generate data to meet the requirements of the premarket notification program because the agency is requesting only that information that the manufacturer or distributor should already have developed to satisfy itself that a dietary supplement containing a new dietary ingredient is in full compliance with the act. However, the agency estimates that extracting and summarizing the relevant information from the company's files, and presenting it in a format that will meet the requirements of section 413 of the act will require a burden of approximately 20 hours of work per submission.

This estimate is based on the annual average number of premarket notifications FDA received during the last 3 years (*i.e.,* 1999–2001), which was 23. Twenty-three represents 12 more notifications than the agency received as an annual average during the previous 3-year period (*i.e.,* 1996–1998). Therefore, FDA anticipates a similar upward trend will be seen in the annual average number of notifications it receives during 2002–2004, which is estimated to be 35 (23 + 12 = 35).

Dated: March 5, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–6493 Filed 3–18–02; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Food 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). The meeting will be open to the public.

Name of Committee: Food Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on scientific issues related to FDA's regulatory responsibilities.

Date and Time: The meeting will be held on April 4, 2002, from 9 a.m. to 6 p.m. and April 5, 2002, from 8:30 a.m. to 2 p.m.

Location: Marriott Hotel, Grand Ballroom, 6400 Ivy Lane, Greenbelt, MD, 301–441–3700.

Contact Person: Catherine M. DeRoever, Center for Food Safety and Applied Nutrition (HFS–6), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–2397, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 10564. Please call the Information Line for up-to-date information on this meeting.

Agenda: The purpose of this meeting is to discuss general scientific principles related to quality factors for infant formula. The committee will also be asked to discuss the scientific issues related to the generalization of findings from a clinical study using preterm infant formula consumed by preterm infants to a term infant formula intended for use by term infants.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by April 1, 2002. Oral presentations from the public will be scheduled on April 4, 2002, between approximately 1 p.m. and 6 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before April 1, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Catherine M. DeRoever at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: March 14, 2002.

Linda A. Suydam,

Senior Associate Commissioner for Communications and Constituent Relations.

[FR Doc. 02–6620 Filed 3–14–02; 4:50 pm]

BILLING CODE 4160–01–S

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4739–N–05]

Notice of Proposed Information Collection: Comment Request; Requirements for Single Family Mortgage Instruments

AGENCY: Office of the Assistant Secretary for Housing—Federal Housing Commissioner, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

DATES: *Comments Due Date:* May 20, 2002.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB Control Number and should be sent to: Wayne Eddins, Reports Management Officer, Department of Housing and Urban Development, 451 7th Street, SW, L'Enfant Plaza Building, Room 8001, Washington, DC 20410.

FOR FURTHER INFORMATION CONTACT: Vance T. Morris, Director, Officer of Single Family Program Development, Department of Housing and Urban Development, 451 7th Street SW, Washington, DC 20410, telephone (202) 708–2121 (this is not a toll free number) for copies of the proposed forms and other available information.

SUPPLEMENTARY INFORMATION: The Department is submitting the proposed information collection to OMB for review, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35, as amended).

This Notice is soliciting comments from members of the public and affected agencies concerning the proposed collection of information to: (1) Evaluate whether the proposed collection is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond; including the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

This Notice also lists the following information:

Title of Proposal: Requirements for Single Family Mortgage Instruments.

OMB Control Number, if applicable: 2502–0404.

Description of the need for the information and proposed use: The single-family mortgage instruments are the documents used to record the mortgage (or deed of trust) and the mortgage note (or deed of trust note). These are public documents used to protect the interests of the mortgagor and mortgagee.

Agency form number, if applicable: None.

Estimation of the total numbers of hours needed to prepare the information collection including number of respondents, frequency of response, and hours of response: The total number hours needed to prepare the information collection is 250,000, the estimated number of respondents is 9,000, the frequency of response varies according to business activity, but generates an estimated 1,000,000 responses per year, and the amount of time needed is 0.25 hours per response.

Status of the proposed information collection: Extension of a currently approved collection.

Authority: The Paperwork Reduction Act of 1995, 44 U.S.C., Chapter 35, as amended.

Dated: March 12, 2002.

John C. Weicher,

Assistant Secretary for Housing—Federal Housing Commissioner.

[FR Doc. 02–6553 Filed 3–18–02; 8:45 am]

BILLING CODE 4210–27–M

DEPARTMENT OF JUSTICE

Notice of Lodging of Partial Consent Decree Under the Comprehensive Environmental Response, Compensation, and Liability Act of 1980, as Amended

Consistent with Departmental policy, 28 CFR 50.7, 38 FR 19029, and 42 U.S.C. 9622(d), notice is hereby given that on February 7, 2002, a proposed Partial Consent Decree (''Decree'') in *United States of America* v. *AlliedSignal Inc., et al.*, Civil Action No. 95–CV–0950–C(Sc), and *United States of America* v. *Niagara Frontier Transportation Authority, Inc., et al.*, Civil Action No. 96–CV–0219C(Sc), was lodged with the United States District Court for the Western District of New York.

In these consolidated actions, the United States sought reimbursement of response costs incurred by the United States in connection with clean up activities at the Bern Metals and Universal Iron and Metals Superfund Sites located in the City of Buffalo, Erie County, New York. The proposed Decree will resolve the United States' claims under the Comprehensive Environmental Response, Compensation, and Liability Act (''CERCLA''), 42 U.S.C. 9601 et seq., on behalf of the United States Environmental Protection Agency (''EPA'') against defendants AlliedSignal Inc. (now Honeywell International,