necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* Medicare Program Integrity Program Organizational Conflict of Interest Disclosure Certificate and Supporting Regulations in 42 CFR 421.300 and 421.318; *Form No.:* CMS–R–232 (OMB #0938–0723); *Use:* This information is used to assess whether contractors who perform, or who seek to perform, Medicare Integrity Program functions, such as medical review, fraud review or cost audits, have organizational conflicts of interest and whether any conflicts have been resolved. The entities providing the information will be organizations that have been awarded, or seek award of, a Medicare Integrity Program contract; *Frequency:* On occasion; *Affected Public:* Businesses or other for profit; *Number of Respondents:* 5; *Total Annual Responses:* 5; *Total Annual Hours:* 1,000.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web site address at *http://www.hcfa.gov/regs/prdact95.htm,* or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov,* or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: July 23, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–19502 Filed 8–1–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services

[Document Identifier: CMS–R–65]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare and Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* Information Collection Requirements in Final Peer Review Organization Sanction Regulations—42 CFR 1004.40, 1004.50, 1004.60, and 1004.70; *Form No.:* CMS–R–65 (OMB# 0938–0444); *Use:* This final rule updates the procedures governing the imposition and adjudication of program sanctions predicated on the recommendations of Peer Review Organizations (PROs). These changes are being made as a result of statutory revisions designed to address health care fraud and abuse issues and the OIG sanction process; *Frequency:* On occasion; *Affected Public:* Not-for-profit institutions and Business or other for-profit; *Number of Respondents:* 53; *Total Annual Responses:* 1,060; *Total Annual Hours:* 22,684.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web site address at *http://www.hcfa.gov/regs/prdact95.htm,* or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov,* or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: July 18, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, CMS, Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–19503 Filed 8–1–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02n–0319]

Agency Information Collection Activities; Proposed Collection; Comment Request; Blood Establishment Registration and Product Listing, Form FDA 2830

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 (the PRA), 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 information collection provisions relating to the blood establishment registration and product listing requirements and Form FDA 2830.

DATES: Submit written or electronic comments on the collection of information by October 1, 2002.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments on the collection of information to *http://www.accessdata.fda.gov/scripts/*

oc/dockets/edockethome.cfm. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 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.

Blood Establishment Registration and Product Listing, Form FDA 2830—21 CFR Part 607—(OMB Control Number 0910–0052)—Extension

Under section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360), any person owning or operating an establishment that manufactures, prepares, propagates, compounds, or processes a drug or device must register with the Secretary of Health and Human Services, on or before December 31 of each year, his or her name, place of business and all such establishments, and submit, among other information, a listing of all drug or device products manufactured, prepared, propagated, compounded, or processed by him or her for commercial distribution. In part 607 (21 CFR part 607), FDA has issued regulations implementing these requirements for manufacturers of human blood and blood products.

Section 607.20(a) requires certain establishments that engage in the manufacture of blood products to register and to submit a list of blood products in commercial distribution. Section 607.21 requires the establishments entering into the manufacturing of blood products to register within 5 days after beginning such operation and to submit a blood product listing at that time. In addition, establishments are required to register annually between November 15 and December 31 and update their blood product listing every June and December. Section 607.22 requires the use of Form FDA 2830 for registration and blood product listing. Section 607.25 indicates the information required for establishment registration and blood product listing. Section 607.26 requires certain changes to be submitted as an amendment to the establishment registration within 5 days of such changes. Section 607.30 requires establishments to update, as needed, their blood product listing information every June and at the annual registration. Section 607.31 requires that additional blood product listing information be provided upon FDA request. Section 607.40 requires foreign blood product establishments to register and submit the blood product listing information, the name and address of the establishment, and the name of the individual responsible for submitting blood product listing information.

Among other uses, this information assists FDA in its inspections of facilities, and its collection is essential to the overall regulatory scheme designed to ensure the safety of the nation's blood supply. Form FDA 2830, Blood Establishment Registration and Product Listing, is used to collect this information. The likely respondents are blood banks, blood collection facilities, and blood component manufacturing facilities.

FDA estimates the burden of this collection of information based upon the database and past experience of the Center for Biologics Evaluation and Research, Division of Blood Applications in regulatory blood establishment registration and product listing. Most blood banks are familiar with the regulations and registration requirements to fill out this form.

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	Form FDA 2830	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
607.20(a), 607.21, 607.22, 607.25, and 607.40	Initial Registration	300	1	300	1	300
607.21, 607.22, 607.25, 607.26, 607.31, and 607.40	Re-registration	2,867	1	2,867	0.5	1,434
607.21, 607.25, 607.30, 607.31, and 607.40	Product Listing Update	75	1	75	0.25	19

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]—Continued

21 CFR Section	Form FDA 2830	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Total						1,753

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

Dated: July 26, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–19493 Filed 8–1–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Request for Nominations for Members of Public Advisory Committee; Food Advisory Committee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting nominations for members to serve on the Food Advisory Committee (the Committee) in FDA's Center for Food Safety and Applied Nutrition (CFSAN) and six subcommittees. Nominations will be accepted for current vacancies and vacancies that will or may occur on the Committee during the next 12 months.

FDA has special interest in ensuring that women, minority groups, and individuals with disabilities are adequately represented on advisory committees and, therefore, encourages nominations from these groups. Final selection from among qualified candidates for each vacancy will be determined by the expertise required to meet specific agency needs and in a manner to ensure appropriate balance of membership.

DATES: September 3, 2002.

ADDRESSES: All nominations for membership should be sent to Catherine M. DeRoever (*see* **FOR FURTHER INFORMATION CONTACT**).

FOR FURTHER INFORMATION CONTACT: *Regarding all nominations for membership*: 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, FAX 301–436–2633, e-mail: *Catherine.DeRoever@cfsan.fda.gov*.

SUPPLEMENTARY INFORMATION: FDA is requesting nominations for members to serve on the advisory committees listed below. Individuals should have expertise in the activity of the Committee. Vacancies will begin June 30, 2002.

Food Advisory Committee

The Committee provides advice primarily to the Director, Center for Food Safety and Applied Nutrition (CFSAN), and as needed, to the Commissioner of Food and Drugs (the Commissioner), and other appropriate officials, on emerging food safety, food science, nutrition, and other food-related issues that FDA considers of primary importance for its food and cosmetics program. The Committee may be charged with reviewing and evaluating available data and making recommendations on matters such as those relating to: (1) Broad scientific and technical food or cosmetic related issues, (2) the safety of new foods and food ingredients, (3) labeling of foods and cosmetics, (4) nutrient needs and nutritional adequacy, and (5) safe exposure limits for food contaminants. The Committee also may be asked to provide advice and make recommendations on ways of communicating to the public the potential risks associated with these issues and on approaches that might be considered for addressing the issues.

The Committee was restructured on July 28, 2000, to consist of a ''parent'' Committee and four standing Subcommittees. The Subcommittees are as follows: (1) Additives and Ingredients, (2) Contaminants and Natural Toxicants, (3) Dietary Supplements, and (4) Food Biotechnology. Two additional Subcommittees are being added to the ''parent'' Committee: (1) Infant Formula, and (2) Nutrition.

The purpose of the new Subcommittees is to provide highly specialized expertise in the review and analysis of assigned topics. Meetings of the subcommittees will be open to the public except as otherwise determined by the Commissioner or designee. The Subcommittee's findings, conclusions, and recommendations will be reported to the ''parent'' Committee. As a general matter, included in this report will be a recommendation(s) from the Subcommittee on the final disposition of an assigned topic. Generally, matters that cross-cut agency program areas will fall under the purview of the ''parent'' Committee. Issues relating to the microbiological safety of food will be addressed by the National Advisory Committee on Microbiological Criteria for Foods.

Criteria for Members

Persons nominated for membership on the Committee shall be knowledgeable in the fields of physical sciences, biological and life sciences, food science, risk assessment and other relevant scientific and technical disciplines. The agency particularly is interested in considering candidates with a comprehensive background in food technology, molecular biology, genetics, biotechnology, and a variety of medical specialties, as many issues brought before the Committee involve medical or epidemiological impact on nutrients, additives, contaminants, or other constituents of the diet, such as dietary supplements. The term of office is up to 4 years.

The Committee includes technically qualified members who are identified with consumer interests and representatives of industry interests.

Nomination Procedures

Interested persons may nominate one or more qualified persons for membership on the Committee. Nominations shall state that the nominee is willing to serve as a member of the Committee and appears to have no conflict of interest that would preclude Committee membership. Additionally, the nominee's mailing address, telephone number, and curriculum vitae must accompany the nomination. The agency cannot guarantee further consideration of nominations that do not include this requested information. Potential candidates will be asked by FDA to provide detailed information concerning such matters as financial holdings, employment, consultancies, and research grants and/or contracts to permit evaluation of possible sources of conflicts of interest.

Industry Representatives

Regarding nominations for members representing industry interests, a letter will be sent to each person or