in tracing the disposal of a certified batch or a batch that has been refused certification for noncompliance with the color additive regulations. The manufacturer's batch weight is used for assessing the certification fee. The batch weight also is used to account for the disposal of a batch of certified or certification-rejected color additive. The batch weight can be used in a recall to determine whether all unused color additive in the batch has been recalled. The manufacturer's name and address and the name and address of the person requesting certification are used to contact the person responsible should a question arise concerning compliance with the color additive regulations. Information on storage conditions pending certification is used to evaluate whether a batch of certified color additive is inadvertently or intentionally altered in a manner that would make the sample submitted for certification analysis unrepresentative of the batch. FDA checks storage information during inspections. Information on intended uses for a batch of color additive is used to assure that a batch of certified color additive will be used in accordance with the requirements of its listing regulation. The statement of the fee on a certification request is used for accounting purposes so that a person requesting certification can be notified promptly of any discrepancies.

In the **Federal Register** of July 24, 2007 (72 FR 40310), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

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
80.21	32	174	5,568	0.20	1,114
80.22	32	174	5,568	0.05	278
Total				0.25	1,392

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

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

21 CFR Section	No. of Recordkeepers	Annual Frequency of Recordkeeping	Total Annual Records	Hours per Record	Total Hours
80.39	32	174	5,568	0.25	1,392
TOTAL					1,392

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

FDA bases its estimate on its review of the certification requests received over the past 3 fiscal years (FY). The annual burden estimate for this information collection is 2,784 hours. The estimated reporting burden for this information collection is 1,392 hours and the estimated recordkeeping burden for this information collection is 1,392 hours. From FY 2004 to FY 2006, FDA processed an average of 5,568 responses (requests for certification of batches of color additives) per year. There were 32 different respondents, corresponding to an average of approximately 174 responses from each respondent per year. Using information from industry personnel, FDA estimates that an average of 0.25 hour per response is required for reporting (preparing certification requests and accompanying sample labels) and an average of 0.25 hour per response is required for recordkeeping.

On February 13, 2006, FDA introduced a Web-based Color Certification information system. The system was fully operational for FY 2007. This system allows certifiers to request color certification on-line, follow their submissions through the process, and obtain information on account status. The system sends back the certification results electronically, allowing certifiers to sell their certified color before receiving hard copy certificates. Any delays in the system result only from shipment of color additive samples to FDA's Office of Cosmetics and Colors for analysis. FDA expects future reductions in the hour burdens for reporting and recordkeeping from use of the Web-based system.

Dated: November 27, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–23352 Filed 11–30–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007F–0454]

General Mills, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that General Mills, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of ultraviolet radiation for the reduction of pathogens and other microorganisms in aqueous sugar solutions and potable water intended for use in food production.

FOR FURTHER INFORMATION CONTACT: Laura A. Dye, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–1275.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 7M4770) has been filed by General Mills, Inc., One General Mills Blvd., Minneapolis, MN 55426. The petition proposes to amend the food additive regulations in § 179.39 *Ultraviolet radiation for the processing and treatment of food* (21 CFR 170.39) to provide for the safe use of ultraviolet radiation for the reduction of pathogens

and other microorganisms in aqueous sugar solutions and potable water intended for use in food production.

The agency has determined under 21 CFR 25.32(j) that this action is of a 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: November 27, 2007.

Laura M. Tarantino,

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

[FR Doc. E7–23400 Filed 11–30–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Heritable Disorders and Genetic Diseases in Newborns and Children; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: Advisory Committee on Heritable Disorders and Genetic Diseases in Newborns and Children (ACHDGDNC).

Dates and Times: Jan. 14, 2008, 9 a.m. to 5 p.m. Jan. 15, 2008, 8:30 a.m. to 3:00 p.m.

Place: Bethesda North Marriott Hotel and Conference Center, 5701 Marinelli Road, Bethesda, MD 20852.

Status: The meeting will be open to the public with attendance limited to space availability.

Purpose: The ACHDGDNC was established to advise and guide the Secretary regarding the most appropriate application of universal newborn screening tests, technologies, policies, guidelines and programs for effectively reducing morbidity and mortality in newborns and children having or at risk for heritable disorders. The ACHDGDNC also provides advice and recommendations concerning the grants and projects authorized under the Heritable Disorders Program.

Agenda: The meeting will include a presentation and continued discussions on the nomination/evaluation process for newborn screening candidate conditions. There will be presentations on utilizing partnerships for follow-up in newborn screening systems, a presentation from the Secretary's Advisory Committee on Genetics, Health and Society on an assessment of the impact of patients on access to tests in both clinical practice and public health settings, as well as presentations on the continued work and reports of the ACHDGDNC's subcommittees on laboratory standards and procedures, follow-up and treatment, and education and training, and the workgroup on research. Proposed agenda items are subject to change.

Time will be provided for public comment. Individuals who wish to provide public comment or who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the ACHDGDNC Staff, Jill F. Shuger, M.S. (contact information provided below).

Contact Person: Anyone interested in obtaining a roster of members or other relevant information should write or contact Jill F. Shuger, M.S., Maternal and Child Health Bureau, Health Resources and Services Administration, Room 18A–19, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443–1080, jshuger@hrsa.gov. Information on the Advisory Committee is available at *http://mchb.hrsa.gov/programs/genetics/committee.*

Dated: November 27, 2007.

Alexandra Huttinger,

Acting Director, Division of Policy Review and Coordination.

[FR Doc. E7–23334 Filed 11–30–07; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HOMELAND SECURITY

Transportation Security Administration

New Agency Information Collection Activity Under OMB Review: Pipeline Security Awareness (CD–1) Effectiveness Assessment

AGENCY: Transportation Security Administration, DHS.

ACTION: Notice.

SUMMARY: This notice announces that the Transportation Security Administration (TSA) has forwarded the new Information Collection Request (ICR) abstracted below to the Office of Management and Budget (OMB) for review and approval under the Paperwork Reduction Act. The ICR describes the nature of the information collection and its expected burden. TSA published a **Federal Register** notice, with a 60-day comment period soliciting comments, of the following collection of information on January 31, 2007, 72 FR 4526.

DATES: Send your comments by January 2, 2008. A comment to OMB is most effective if OMB receives it within 30 days of publication.

ADDRESSES: Interested persons are invited to submit written comments on the proposed information collection to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to Nathan Lesser, Desk Officer, Department of Homeland Security/TSA, and sent via electronic mail to *oira_submission@omb.eop.gov* or faxed to (202) 395–6974.

FOR FURTHER INFORMATION CONTACT: Joanna Johnson, Communications Branch, Business Management Office, Operational Process and Technology, TSA–11, Transportation Security Administration, 601 South 12th Street, Arlington, VA 22202–4220; telephone (571) 227–3651; facsimile (571) 227–3885.

SUPPLEMENTARY INFORMATION:

Comments Invited

In accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 et seq.), an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. Therefore, in preparation for OMB review and approval of the following information collection, TSA is soliciting comments to—

(1) Evaluate whether the proposed information requirement 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;

(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 using appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Information Collection Requirement

Title: Pipeline Security Awareness (CD–1) Effectiveness Assessment.

Type of Request: New collection.

OMB Control Number: Not yet assigned.

Form(s): NA.

Affected Public: Pipeline companies.

Abstract: As prescribed by the President in Homeland Security Presidential Directive 7 (HSPD–7), the Department of Homeland Security (DHS) was tasked to protect our nation's critical infrastructure and key resources (CI/KR). Through the National Infrastructure Protection Plan (NIPP), DHS gives guidance and direction as to how the Nation will secure its infrastructure. Furthermore, HSPD–7 and the NIPP assigned the responsibility for infrastructure security in the transportation sector to TSA. To this effect, the NIPP further tasks each sector to build security partnerships, set security goals and to measure their