enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: The Food Advisory Committee will meet to discuss whether available relevant data demonstrate a link between children's consumption of synthetic color additives in food and adverse effects on behavior.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm*. Scroll down to the appropriate advisory committee link.

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 on or before March 23, 2011. Oral presentations from the public will be scheduled between approximately 10 a.m. and 11 a.m. on March 31, 2011. Those individuals interested in making formal oral presentations should notify the contact person 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 on or before March 15, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by March 16, 2011.

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 Carolyn Jeletic at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

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

Dated: November 24, 2010.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010–30187 Filed 11–30–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Food Labeling Workshop; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA), Office of Regulatory Affairs, Southwest Regional Small Business Representative (SWR SBR) Program, in collaboration with Iowa State University, is announcing a public workshop entitled "Food Labeling Workshop." This public workshop is intended to provide information about FDA food labeling regulations and other related subjects to the regulated industry, particularly small businesses and startups.

Date and Time: This public workshop will be held on March 3 and 4, 2011, from 8 a.m. to 5 p.m.

Location: The public workshop will be held at the Scheman Conference Center, Lincoln Way and University Avenue, Iowa State Center, Ames, IA.

Contact: David Arvelo, Food and Drug Administration, Southwest Regional Office, 4040 North Central Expressway, suite 900, Dallas, TX 75204, 214–253–4952, FAX: 214–253–4970, or email: *david.arvelo@fda.hhs.gov.*

For information on accommodation options, visit *http://www.fshn.hs.iastate.edu/foodlabel/register.php* or contact Dr. Ruth MacDonald, 2312 Food Sciences Building, Iowa State University, Ames, IA 50011, 515–294–5991, FAX: 515–294–8181, email: *ruthmacd@iastate.edu.*

Registration: You are encouraged to register by February 21, 2011. The workshop has a $250 registration fee to cover the cost of facilities, materials, lunch on day 1, and breaks. There is no registration fee for FDA employees. Seats are limited; please submit your registration as soon as possible. Workshop space will be filled in order of receipt of registration. Those accepted into the workshop will receive confirmation. Registration will close after the workshop is filled. Registration at the site is not guaranteed but may be possible on a space available basis on the day of the public workshop beginning at 8 a.m. The cost of registration at the site is $350 payable to: "Iowa State University." If you need special accommodations due to a disability, please contact Dr. Ruth MacDonald (*see Contact*) at least 14 days in advance.

Registration Form Instructions: To register, please complete the online registration form at *http://www.fshn.hs.iastate.edu/foodlabel/register.php,* or submit your full name, business or organization name, complete mailing address, telephone number, email address, optional fax number, and any special accommodations required due to disability, along with a check or money order for $250 payable to "Iowa State University." Mail to: Dr. Ruth MacDonald, Food Science and Human Nutrition, 2312 Food Sciences Building, Ames, IA 50011.

Transcripts: Transcripts of the public workshop will not be available due to the format of this workshop. Requests for workshop handouts may be obtained through David Arvelo (see *Contact*).

SUPPLEMENTARY INFORMATION: This public workshop is being held in response to the large volume of food labeling inquiries from small food manufacturers and startups originating from the area covered by FDA's Kansas City District Office. The SWR SBR presents these workshops to help achieve objectives set forth in section 406 of the Food and Drug Administration Modernization Act of 1997 (21 U.S.C. 393), which include working closely with stakeholders and maximizing the availability and clarity of information to stakeholders and the public. This is consistent with the purposes of the SBR Program, which are in part to respond to industry inquiries, develop educational materials, and sponsor workshops and conferences to provide firms, particularly small businesses, with firsthand working knowledge of FDA's requirements and compliance policies. This workshop is

also consistent with the Small Business Regulatory Enforcement Fairness Act of 1996 (Pub. L. 104–121), as outreach activities by government agencies to small businesses.

The goal of this public workshop is to present information that will enable manufacturers and regulated industry to better comply with labeling requirements, especially in light of growing concerns about obesity and food allergens. Information presented will be based on Agency position as articulated through regulation, compliance policy guides, and information previously made available to the public. This is a hands-on workshop. Topics to be discussed at the workshop include: (1) Mandatory label elements, (2) nutrition labeling requirements, (3) the Food Allergen Labeling and Consumer Protection Act of 2004, (4) health and nutrient content claims, (5) special labeling issues such as exemptions, and (6) current topics on food labeling and nutrition. FDA expects that participation in this public workshop will provide regulated industry with greater understanding of the Agency's regulatory and policy perspectives on food labeling and increase voluntary compliance with labeling requirements.

Dated: November 24, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–30191 Filed 11–30–10; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERIVCES

Health Resources and Services Administration

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

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443–1129. The following request has been submitted to OMB for review under the Paperwork Reduction Act of 1995:

Proposed Project: Ryan White HIV/AIDS Program Annual Data Report Form: Data Report Form: (OMB No. 0915–0253)—Extension

The Ryan White HIV/AIDS Program Annual Data Report was first implemented in 2002 by HRSA's HIV/AIDS Bureau (HAB) as the CARE Act Data Report (CADR). Grantees and their subcontracted service providers who are funded under Parts A, B, C, and D of Title XXVI of the Public Health Service Act, as amended by the Ryan White HIV/AIDS Treatment Extension Act of 2009 (Ryan White HIV/AIDS Program), complete the report. All Parts of the Ryan White HIV/AIDS Program specify HRSA's responsibilities in the administration of grant funds, the allocation of funds, the evaluation of programs for the population served, and the improvement of the quantity and quality of care. Accurate records of the providers receiving Ryan White HIV/AIDS Program funding, the services provided, and the clients served continue to be critical to the implementation of the legislation and thus are necessary for HRSA to fulfill its responsibilities. Ryan White HIV/AIDS Program Grantees are required to report aggregate data to HRSA annually. The Ryan White Data Report (RDR) is completed by grantees and their subcontracted service providers. The Report has seven different sections requesting: (1) Characteristics of the service providers; (2) demographic information about the clients served; (3) information about the type of core and support services provided and the number of clients served; (4) information about HIV counseling and testing services; (5) clinical information about the clients who receive medical care; (6) demographic tables for Parts C and D; and (7) information about the Health Insurance Program. The primary purposes of the Data Report are to: (1) Characterize the organizations where clients receive services; (2) provide information on the number and characteristics of clients who receive Ryan White HIV/AIDS Program Services; and (3) enable HAB to describe the type and amount of services a client receives. In addition to meeting the goal of accountability to the Congress, clients, advocacy groups, and the general public, information collected on the RDR is critical for HRSA, state and local grantees, and individual providers to assess the status of existing HIV-related service delivery systems.

The estimated burden is as follows:

Program under which grantee is funded	Number of grantee respondents	Responses per grantee	Hours to coordinate receipt of data	Total hour burden
Part A	56	1	40	2,240
Part B	59	1	40	2,360
Part C	354	1	20	7,080
Part D	98	1	20	1,960
Subtotal	567	.....	.....	13,640

Program under which provider is funded	Number of provider respondents	Responses per provider	Hours per response	Total hour burden
Part A only	685	1	26	17,810
Part B only	558	1	26	14,508
Part C only	95	1	44	4,180
Part D only	59	1	42	2,478
Multiply funded	683	1	50	34,150
Subtotal	2,080	.....	.....	73,126