The official public docket is available for public viewing at the Office of Environmental Information (OEI) Docket in the EPA Docket Center, EPA West, Room 3334, 1301 Constitution Ave. NW., Washington, DC. The EPA Docket Center Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566–1744, and the telephone number for the OEI Docket is (202) 566–1752.

An electronic version of the public docket is available through *www.regulations.gov.* You may use the *www.regulations.gov* to submit or view public comments, access the index listing of the contents of the official public docket, and to access those documents in the public docket that are available electronically. Once in the system, key in the appropriate docket identification number then select ''search.''

It is important to note that EPA's policy is that public comments, whether submitted electronically or in paper, will be made available for public viewing online at *www.regulations.gov* without change, unless the comment contains copyrighted material, CBI, or other information whose disclosure is restricted by statute. Information claimed as CBI and other information whose disclosure is restricted by statute are not included in the official public docket or in the electronic public docket. EPA's policy is that copyrighted material, including copyrighted material contained in a public comment, will not be placed in EPA's electronic public docket but will be available only in printed, paper form in the official public docket. Although not all docket materials may be available electronically, you may still access any of the publicly available docket materials through the EPA Docket Center.

B. How and to whom do I submit comments?

You may submit comments as provided in the **ADDRESSES** section. Please ensure that your comments are submitted within the specified comment period. Comments received after the close of the comment period will be marked ''late.'' EPA is not required to consider these late comments.

If you submit an electronic comment, EPA recommends that you include your name, mailing address, and an email address or other contact information in the body of your comment and with any disk or CD–ROM you submit. This ensures that you can be identified as the submitter of the comment and allows EPA to contact you in case EPA cannot read your comment due to technical difficulties or needs further information on the substance of your comment. Any identifying or contact information provided in the body of a comment will be included as part of the comment that is placed in the official public docket, and made available in EPA's electronic public docket. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment.

Use of the *www.regulations.gov* Web site to submit comments to EPA electronically is EPA's preferred method for receiving comments. The electronic public docket system is an ''anonymous access'' system, which means EPA will not know your identity, email address, or other contact information unless you provide it in the body of your comment. In contrast to EPA's electronic public docket, EPA's electronic mail (email) system is not an ''anonymous access'' system. If you send an email comment directly to the Docket without going through *www.regulations.gov,* your email address is automatically captured and included as part of the comment that is placed in the official public docket, and made available in EPA's electronic public docket.

Dated: January 27, 2012.

Patricia Embrey,

Acting Associate General Counsel.

[FR Doc. 2012–2443 Filed 2–2–12; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

Public Availability of the Federal Communications Commission's FY 2011 Service Contract Inventory and FY 2010 Service Contract Inventory Analysis

AGENCY: Federal Communications Commission.

ACTION: Notice of public availability of service contract inventory and analysis.

SUMMARY: The Federal Communications Commission is publishing this notice to advise the public of the availability of its FY 2011 Service Contract Inventory and FY 2010 Service Contract Inventory Analysis as required by Section 743 of Division C of the Consolidated Appropriations Act of 2010 (Pub. L. 111–117). The FY 2011 inventory provides information on service contract actions over $25,000 that were made in FY 2011. The information is organized by function to show how contracted resources are distributed throughout the agency. The FY 2010 analysis provides additional information about the Federal Communications Commission's FY 2010 inventory. The FY 2011 inventory and analysis of the FY 2010 inventory have been developed in accordance with guidance issued by the Office of Management and Budget's Office of Federal Procurement Policy (OFPP) on November 5, 2010 and December 19, 2011. The guidance is available online at: *http://www.whitehouse.gov/omb/procurement_index_memo.*

The Federal Communications Commission has posted its FY 2011 inventory, a summary of the FY 2011 inventory, and its analysis of its FY 2010 inventory on the Federal Communications Commission's Web site at the following link: *http://www.fcc.gov/encyclopedia/service-contract-inventory.*

FOR FURTHER INFORMATION CONTACT: Questions regarding the service contract inventory or analysis should be directed to Mr. Daniel Daly, Chief of Staff, Office of the Managing Director at (202) 418–1832 or *Daniel.Daly@fcc.gov.*

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary, Office of Managing Director.

[FR Doc. 2012–2386 Filed 2–2–12; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–12–11KS]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Pilot Study of Community-Based Surveillance of Supports for Healthy Eating/Active Living (HE/AL)—New—

National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC proposes to conduct a pilot study to examine the feasibility of establishing a national community-level surveillance system on policy supports for healthful eating and active living. Results of the feasibility study will be used to assess the feasibility of establishing a national surveillance system and the best methods for encouraging a high response rate in a representative sample of communities. The pilot study will be conducted in two states with a representative sample of 400 communities, 200 municipalities in each state. The sample frame will be generated from the U.S. Census of Governments.

The proposed pilot study is designed to address three key methodological objectives. The first objective is to test the feasibility of the proposed sampling frame and to answer sample design issues related to determining sampling criteria for inclusion, as well as the development of weights and estimates.

The second objective is to identify and critically evaluate whether respondents in diverse municipalities of various sizes and organizational structures are able to answer a self-administered survey questionnaire. The survey questionnaire includes 42 items on the following topics: Community-wide planning efforts for healthy eating and active living, the built environment and policies that support physical activity, and policies and practices that support access to healthy food and healthy eating. The estimated burden per response is one hour. Issues to be addressed include critical assessment of the strengths and weaknesses of methods for identifying the best respondents for completing the survey questionnaire; conducting a limited process evaluation that identifies the barriers and challenges respondents may incur in providing reasonable and current data for the questionnaire; and arriving at a data collection instrument with the lowest possible threshold for respondent burden.

The third objective is to identify and critically evaluate different methods of study recruitment and non-response follow-up. A split-sample approach will be used to assign each target respondent to one of two groups: a low-intensity recruitment group or a moderate-intensity recruitment group. All target respondents in the study sample will receive email reminders to encourage participation in the survey. Target respondents in the moderate-intensity recruitment group will also receive up to three telephone contacts to address questions and serve as reminders. The estimated burden per contact is five minutes.

Respondents will be city/town planners and managers, or individuals with similar responsibilities. The majority of survey responses will be collected using a secure, web-based survey data collection system. Respondents who prefer to complete a paper survey will be able to print the survey from the web-based data collection system, complete it, and return it using instructions that will be provided. OMB approval is requested for one year. Participation is voluntary and there are no costs to respondents other than their time. The total estimated burden hours are 450.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
City/Town Manager-Planner	Survey of Community-Based Policy and Environmental Supports for Healthy Eating and Active Living	400	1	1
	Telephone Non-Response Follow-up Contact Script	200	3	5/60

Kimberly S. Lane,

Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2012–2413 Filed 2–2–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0001]

Oncologic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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: Oncologic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on March 20, 2012, from 8 a.m. to 5 p.m.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993–0002. Information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/default.htm;* under the heading ''Resources for You,'' click on ''Public Meetings at the FDA White Oak Campus.'' Please note that visitors to the White Oak Campus must enter through Building 1.

Contact Person: Caleb Briggs, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, (301) 796–9001, Fax: (301) 847–8533, email: *ODAC@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–(800) 741–8138 (301) 443–0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly 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: On March 20, 2012, during the morning session, the committee will discuss supplemental new drug