TRANSACTION GRANTED EARLY TERMINATION—Continued

ET date	Trans No.	ET req status	Party name
	.....	G	Hartmarx Corporation.
25–JUN–09	20090505	G	Valline S.r.l.
	.....	G	Cornerstone Therapeutics Inc.
	.....	G	Cornerstone Therapeutics Inc.
	20090512	G	Sumitomo Metal Mining Co., Ltd.
	.....	G	Teck Resources Limited.
	.....	G	Teck-Pogo, Inc.
	20090526	G	Golden Gate Capital Opportunity Fund, L.P.
	.....	G	Aeon Co. Ltd.
	.....	G	J. Jill, LLC.
	.....	G	Birch Pond Realty Corporation.
26–JUN–09	20090537	G	Communications Infrastructure Investments, LLC.
	.....	G	FiberNet Telecom Group, Inc.
	.....	G	FiberNet Telecom Group, Inc.
	20090541	G	Aquiline Financial Services Fund L.P.
	.....	G	Swiss Reinsurance Company Ltd.
	.....	G	Conning & Company.
	.....	G	Conning Asset Management Limited.
	.....	G	Conning Asset Management (Europe) Limited.
29–JUN–09	20090492	G	John C. Malone.
	.....	G	Liberty Entertainment, Inc.
	.....	G	Liberty Entertainment, Inc.
02–JUL–09	20090502	G	EMC Corporation.
	.....	G	Data Domain, Inc.
	.....	G	Data Domain, Inc.
	20090506	G	NetApp, Inc.
	.....	G	Data Domain, Inc.
	.....	G	Data Domain, Inc.
06–JUL–09 20090543	.....	G	Golden Gate Capital Opportunity Fund, L.P.
	.....	G	SoftBrands, Inc.
	.....	G	SoftBrands, Inc.
	20090544	G	Nokia Corporation.
	.....	G	Nortel Networks Corporation.
	.....	G	Nortel Networks Corporation.
	20090545	G	SAIC, Inc.
	.....	G	R.W. Beck Group, Inc.
	.....	G	R.W. Beck Group, Inc.
	20090553	G	CCMP Capital Investors II, L.P.
	.....	G	Eddie Bauer Holdings, Inc.
	.....	G	Eddie Bauer Holdings, Inc.
07–JUL–09	20090539	G	Sageview Capital Master, L.P.
	.....	G	Gerresheimer AG.
	.....	G	Gerresheimer AG.

For Further Information Contact: Sandra M. Peay, Contact Representative; or Renee Hallman, Contact Representative, Federal Trade Commission Premerger, Notification Office, Bureau Of Competition, Room H–303, Washington, DC 20580, (202) 326–3100.

By Direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2010–20814 Filed 8–23–10; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: Office of Community Services (OCS) Community Economic Development (CED) and Job Opportunities for Low-Income Individuals (JOLI) Standard Reporting Format.

OMB No.: New Collection.

Description: The Office of Community Services (OCS) is collecting key information about projects funded through the Community Economic Development (CED) and Job Opportunities for Low-Income Individuals (JOLI) programs. The legislative requirement for these two programs is in Title IV of the Community Opportunities, Accountability and Training and Educational Services Act (COATS Human Services Reauthorization Act) of October 27, 1998, Public Law 105–285, section 680(b) as amended. The Performance Progress Report (PPR) is a new proposed reporting format that will collect information concerning the outcomes and management of CED and JOLI projects. OCS will use the data to critically review the overall design and effectiveness of each program.

The PPR will be administered to all active grantees of the CED and JOLI programs. Grantees will be required to use this reporting tool for their semiannual reports. The majority of the questions in this tool were adapted from

a previously approved questionnaire, Office of Management and Budget (OMB) Control Number: 0970–0317. Questions were also adapted to the OMB-approved reporting format of the PPR, specifically forms SF–PPR, SF–PPR–A, SF–PPR–B, and SF–PPR–E. Additional changes were made to improve the clarity and quality of the data and to eliminate unnecessary questions.

Respondents: Current CED and JOLI grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Questionnaire for current OCS–JOLI grantees	40	2	1.50	120
Questionnaire for current OCS–CED grantees	170	2	1.50	510
Estimated Total Annual Burden Hours				630

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: August 18, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–20892 Filed 8–23–10; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Menthol Report Subcommittee of the Tobacco Products Scientific 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: Menthol Report Subcommittee of the Tobacco Products Scientific 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 September 27, 2010, from 9 a.m. to 1 p.m.

Location: Center for Tobacco Products, Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 1–877–287–1373.

Contact Person: Cristi Stark, Center for Tobacco Products, Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 1–877–287–1373 (choose option 4), email: *TPSAC@fda.hhs.gov* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 8732110002. 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 September 27, 2010, the subcommittee will receive a presentation and discuss the timelines and structure of the Tobacco Products Scientific Advisory Committee's required report to the Secretary of Health and Human Services regarding the impact of use of menthol in cigarettes on the public health.

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 September 17, 2010. Oral presentations from the public will be scheduled between approximately 10 a.m. and 11 a.m. on September 27, 2010. Those desiring to make 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 September 9, 2010. 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