identifier will be assigned, as required, to process the specimen.

Public Burden Statement

Public Burden Statement: An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this project is 0930-0158. Public reporting burden for this collection of information is estimated to average 5 minutes/donor, 4 minutes/collector, 3 minutes/test facility; and 3 minutes/Medical Review Officer. Send comments regarding this burden estimate or any other aspect of this collection information, including suggestions for reducing this burden, to SAMHSA Reports Clearance Officer, 1 Choke Cherry Road, Room 7-1044, Rockville, Maryland 20857.

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Prior to an inspection, a laboratory is required to submit specific information regarding its laboratory procedures. Collecting this information prior to an inspection allows the inspectors to thoroughly review and understand the laboratory's testing procedures before arriving at the laboratory.

The annual total burden estimates for the Federal Drug Testing Custody and Control Form, the NLCP application, the NLCP inspection checklist, and NLCP recordkeeping requirements are shown in the following table.

Form/respondent	Burden/ response (hrs.)	Number of responses	Total annual burden (hrs.)
Custody and Control Form:			
Donor	.08	7,096,000	567,680
Collector	.07	7,096,000	496,720
Laboratory	.05	7,096,000	354,800
Medical Review Officer	.05	7,096,000	354,800
Laboratory Application	3.00	3	9
Laboratory Inspection Checklist	3.00	100	300
Laboratory Recordkeeping	250.00	50	12,500
Total			1,786,809

Written comments and recommendations concerning the proposed information collection should be sent by August 16, 2010 to: SAMHSA Desk Officer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503; due to potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, respondents are encouraged to submit comments by fax to: 202–395–6974.

Dated: July 12, 2010.

Dennis O. Romero,

Deputy Director, Office of Program Services.

[FR Doc. 2010–17400 Filed 7–15–10; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

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

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Project: Pretesting of Substance Abuse Prevention and Treatment and Mental Health Services Communication Messages—(OMB No. 0930–0196)—Extension

As the Federal agency responsible for developing and disseminating authoritative knowledge about substance abuse prevention, addiction treatment, and mental health services and for mobilizing consumer support and increasing public understanding to overcome the stigma attached to addiction and mental illness, the Substance Abuse and Mental Health Services Administration (SAMHSA) is responsible for development and dissemination of a wide range of education and information materials for

both the general public and the professional communities. This submission is for generic approval and will provide for formative and qualitative evaluation activities to (1) assess audience knowledge, attitudes, behavior and other characteristics for the planning and development of messages, communication strategies and public information programs; and (2) test these messages, strategies and program components in developmental form to assess audience comprehension, reactions and perceptions. Information obtained from testing can then be used to improve materials and strategies while revisions are still affordable and possible. The annual burden associated with these activities is summarized below.

Activity	Nunmber of respondents	Responses/ respondent	Hours per response	Total hours
Individual In-depth Interviews:				
General Public	400	1	.75	300
Service Providers	200	1	.75	150
Focus Group Interviews:				
General Public	3,000	1	1.5	4,500
Service Providers	1,500	1	1.5	2,250
Telephone Interviews:				
General Public	335	1	.08	27
Service Providers	165	1	.08	13
Self-Administered Questionnaires:				
General Public	2,680	1	.25	670
Service Providers	1,320	1	.25	330
Gatekeeper Reviews:				
General Public	1,200	1	.50	600
Service Providers	900	1	.50	450
Total	11,700			9,290

Written comments and recommendations concerning the proposed information collection should be sent by August 16, 2010 to: SAMHSA Desk Officer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503; due to potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, respondents are encouraged to submit comments by fax to: 202–395–5806.

Dated: July 9, 2010.

Elaine Parry,

Director, Office of Program Services.

[FR Doc. 2010–17358 Filed 7–15–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0350]

Draft Guidance for Tobacco Retailers on Tobacco Retailer Training Programs; Availability; Agency Information Collection Activities; Proposed Collection; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for tobacco retailers entitled "Tobacco Retailer Training Programs." The Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act) does not require retailers to implement retailer training programs. However, the Tobacco Control Act does provide for lower civil money penalties for violations of access, advertising, and promotion restrictions issued under section 906(d) of the Federal Food, Drug, and Cosmetic Act (the act), as amended by the Tobacco Control Act, for retailers who have implemented a training program that complies with standards developed by FDA for such programs. FDA intends to issue regulations establishing standards for approved retailer training programs. In the interim, this draft guidance document is intended to assist tobacco retailers who wish to implement effective training programs for employees.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit electronic or written comments on the draft guidance and on the proposed collection of information by September 14, 2010.

ADDRESSES: Submit electronic comments on the draft guidance, including comments regarding the proposed collection of information to *http://www.regulations.gov*. Submit written comments on the draft guidance, including comments regarding the proposed collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Identify comments with the docket number found in brackets in the heading of this document.

Submit written requests for single copies of the draft guidance document entitled "Tobacco Retailer Training Programs" to the Center for Tobacco Products, Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850–3229. Send one self-addressed adhesive label to assist that office in processing your request or include a fax number to which the draft guidance document may be sent. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

FOR FURTHER INFORMATION CONTACT:

With regard to the draft guidance: Beth Buckler, Center for Tobacco Products, Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850–3229, 1–877–287–1373, *beth.buckler@fda.hhs.gov*.

With regard to the proposed collection of information: JonnaLynn Capezzuto, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50, Rockville, MD 20850, 301–796–3794.