Assets and a trustee to divest any divestiture assets that SCI fails to timely divest. The Commission also may seek civil penalties from SCI for non-compliance with the Consent Agreement.

The proposed Consent Agreement prohibits SCI from acquiring any interest or assets engaged in the provision of cemetery services in the Las Vegas metropolitan area for ten (10) years without providing prior written notice to the Commission. In addition, SCI is required to file periodic reports of compliance with the proposed orders.

The purpose of this analysis is to facilitate public comment on the proposed Consent Agreement, and it is not intended to constitute an official interpretation of the proposed Consent Agreement or to modify its terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. E9–29332 Filed 12–8–09; 9:02 am]

BILLING CODE: 6750–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–10–0021]

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) 639–5960 or send an e-mail 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

National Coal Workers' Autopsy Study (NCWAS)—Extension—(0920–0021 Exp. 1/31/2010) National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Under the Federal Coal Mine Health and Safety Act of 1977, Public Law 91–173 (amended the Federal Coal Mine and Safety Act of 1969); the Public Health Service has developed a nationwide autopsy program (NCWAS) for underground coal miners. The Consent Release and History Form are primarily used to obtain written authorization from the next-of-kin to perform an autopsy on the deceased miner. Because a basic reason for the post-mortem examination is research (both epidemiological and clinical), a minimum of essential information is collected regarding the deceased miners, including occupational history and smoking history. The data collected will be used by the staff at NIOSH for research purposes in defining the diagnostic criteria for coal workers' pneumoconiosis (black lung) and pathologic changes and will be correlated with x-ray findings.

It is estimated that only 5 minutes is required for the pathologist to put a statement on the invoice affirming that no other compensation is received for the autopsy. From past experience, it is estimated that 15 minutes is required for the next-of-kin to complete the Consent Release and History Form. Since an autopsy report is routinely completed by a pathologist, the only additional burden is the specific request of abstract of terminal illness and final diagnosis relating to pneumoconiosis. Therefore, only 5 minutes of additional burden is estimated for the autopsy report.

There are no costs to respondents other than their time. The total estimated burden hours are 21.

ESTIMATED ANNUALIZED BURDEN

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Pathologist	Invoice	50	1	5/60
Pathologist	NCWAS Checklist	50	1	5/60
Next-of-Kin	Consent Release History	50	1	15/60

Dated: December 3, 2009.

Maryam Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E9–29385 Filed 12–8–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0569]

Approved Tobacco Retailer Training Program; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is establishing a public docket to obtain information on suggested elements for approved tobacco retailer training programs. FDA is establishing this docket in order to provide an opportunity for interested parties to provide information and share views on elements that should be included in an effective retailer training program as provided for in the Family Smoking Prevention and Tobacco Control Act (the Tobacco Control Act).

DATES: Submit electronic or written comments by January 8, 2010.

ADDRESSES: Submit electronic comments to *http://www.regulations.gov*. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Anne Kirchner, Center for Tobacco Products, Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850–3229, 301–796–4800, *Anne.Kirchner@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

Tobacco products are responsible for more than 440,000 deaths each year. The Centers for Disease Control and Prevention (CDC) report an estimated 60 million adults smoke cigarettes in the