All of the current McCaffrey's supermarkets are located outside the relevant geographic area. Its Yardley, Pennsylvania, store is approximately six miles, and approximately 15 minutes driving time, from the Genuardi's in Newtown. The Newtown Genuardi's is outside McCaffrey's primary service area and vice versa.

The proposed Order requires Respondents Ahold and Safeway to divest the assets of the Genuardi's to McCaffrey's no later than ten days following Ahold's acquisition of the 16 Genuardi's stores that are subject to the Asset Purchase Agreement. If McCaffrey's ultimately is not approved by the Commission to purchase the assets, Respondents must immediately rescind the divestiture and divest the Newtown Genuardi's assets to a buyer that receives the Commission's prior approval. The proposed Order contains additional provisions designed to ensure the adequacy of the proposed relief. For example, for a period of one year, the Order prohibits Respondents from interfering with the hiring of or employment of any employees currently working at the Newtown Genuardi's. Additionally, for a period of ten years, Ahold is required to give the Commission prior notice of plans to acquire a supermarket, or an interest in a supermarket, that has operated or is operating in Newtown, Pennsylvania.

V. Opportunity for Public Comment

The proposed Consent Agreement has been placed on the public record for 30 days to solicit comments from interested persons. Comments received during this period will become part of the public record. After 30 days, the Commission will again review the proposed Consent Agreement, as well as the comments received, and will decide whether to modify the proposed Consent Agreement, withdraw its acceptance of the proposed Consent Agreement, or issue its final Consent Orders.

The sole purpose of this Analysis is to facilitate public comment on the proposed Consent Agreement. This Analysis does not constitute an official interpretation of the proposed Consent Agreement, nor does it modify its terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2012–15308 Filed 6–21–12; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–12–0210]

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

List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products—Extension—Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC), Office on Smoking and Health (OSH) has the primary responsibility for the Department of Health and Human Services (HHS) smoking and health program. HHS's overall goal is to reduce death and disability resulting from cigarette smoking and other forms of tobacco use through programs of information, education and research.

Since 1986, as required by the Comprehensive Smoking Education Act of 1984 (CSEA, 15 U.S.C. 1336 or Pub. L. 98–474), CDC has collected information about the ingredients used in cigarette products. Respondents are commercial cigarette manufacturers, packagers, or importers (or their representatives), who are required by the CSEA to submit ingredient reports to HHS on an annual basis.

Respondents are not required to submit specific forms, however, they are required to submit a list of all ingredients used in their products. CDC requires the ingredient report to be submitted by chemical name and Chemical Abstract Service (CAS) Registration Number, consistent with accepted reporting practices for other companies currently required to report ingredients added to other consumer products. Typically, respondents submit a summary report to CDC with the ingredient information for multiple products, or a statement that there are no changes to their previously submitted ingredient report.

Ingredient reports for new products are due at the time of first importation. Thereafter, ingredient reports are due annually on March 31. Information is submitted to OSH by mailing a written report on the respondent's letterhead, by CD, three-inch floppy disk, or thumb drive. Electronic mail submissions are not accepted. The estimated burden per response is 6.5 hours.

Upon receipt and verification of the annual ingredient report, OSH issues a Certificate of Compliance to the respondent. OSH also uses the information to report to the Congress (as deemed appropriate) discussing the health effects of these ingredients.

There are no costs to respondents other than their time. The total estimated annualized burden hours are 501.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Cigarette Manufacturers, Packagers, and Importers	77	1	6.5

Dated: June 15, 2012.

Ron A. Otten,

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012–15354 Filed 6–21–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry, Department of Health and Human Services, has been renewed for a 2-year period through May 21, 2014.

For information, contact Vikas Kapil, Designated Federal Officer, Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry, Department of Health and Human Services, 4770 Buford Highway Mailstop F61, Chamblee, Georgia 30341, telephone 770/488–8316 or fax 770/488–3385.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: June 13, 2012.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2012–14923 Filed 6–21–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–3262–PN]

Medicare and Medicaid Programs; Application From American Association for Accreditation of Ambulatory Surgery Facilities for Continued Approval of Its Ambulatory Surgery Facilities Accreditation Program

AGENCY: Centers for Medicare and Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: This proposed notice with comment period acknowledges the receipt of an application from the American Association for Accreditation of Ambulatory Surgery Facilities (AAAASF) for continued recognition as a national accrediting organization for ambulatory surgery centers (ASCs) wish to participate in the Medicare or Medicaid programs.

DATES: To be assured consideration, comments must be received at one of the addresses provided below, no later than 5 p.m. on July 23, 2012.

ADDRESSES: In commenting, please refer to file code CMS–3262–PN. Because of staff and resource limitations, we cannot accept comments by facsimile (Fax) transmission.

You may submit comments in one of four ways (please choose only one of the ways listed):

1. *Electronically.* You may submit electronic comments on this regulation to *http://www.regulations.gov*. Follow the ''Submit a comment'' instructions.

2. *By regular mail.* You may mail written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–3262–PN, P.O. Box 8010, Baltimore, MD 21244–8010.

Please allow sufficient time for mailed comments to be received before the close of the comment period.

3. *By express or overnight mail.* You may send written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–3262–PN, Mail Stop C4–26–05, 7500 Security Boulevard, Baltimore, MD 21244–1850.

4. *By hand or courier.* If you prefer, you may deliver (by hand or courier) your written comments before the close of the comment period to either of the following addresses:

a. For delivery in Washington, DC—Centers for Medicare & Medicaid Services, Department of Health and Human Services, Room 445–G, Hubert H. Humphrey Building, 200 Independence Avenue SW., Washington, DC 20201.

(Because access to the interior of the Hubert H. Humphrey Building is not readily available to persons without Federal government identification, commenters are encouraged to leave their comments in the CMS drop slots located in the main lobby of the building. A stamp-in clock is available for persons wishing to retain a proof of filing by stamping in and retaining an extra copy of the comments being filed.)

b. For delivery in Baltimore, MD—Centers for Medicare & Medicaid Services, Department of Health and Human Services, 7500 Security Boulevard, Baltimore, MD 21244–1850.

If you intend to deliver your comments to the Baltimore address, please call telephone number (410) 786–7195 in advance to schedule your arrival with one of our staff members.

Comments mailed to the addresses indicated as appropriate for hand or courier delivery may be delayed and received after the comment period.

For information on viewing public comments, see the beginning of the **SUPPLEMENTARY INFORMATION** section.

FOR FURTHER INFORMATION CONTACT:

Georganne Kuberski, (410) 786–0799.

Patricia Chmielewski, (410) 786–6899.

Cindy Melanson, (410) 786–0310.

SUPPLEMENTARY INFORMATION:

Inspection of Public Comments: All comments received before the close of the comment period are available for viewing by the public, including any personally identifiable or confidential business information that is included in a comment. We post all comments received before the close of the comment period on the following Web site as soon as possible after they have been received: *http://www.regulations.gov*. Follow the search instructions on that Web site to view public comments.

Comments received timely will also be available for public inspection as they are received, generally beginning approximately 3 weeks after publication of a document, at the headquarters of the Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore, Maryland 21244, Monday through Friday of each week from 8:30 a.m. to 4 p.m. To schedule an appointment to view public comments, phone 1–800–743–3951.