public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility; (2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The Protection and Advocacy Voting Access Annual Narrative Report from the Protection and Advocacy Systems is required by federal statute and regulation, the Help America Vote Act (HAVA), Public Law 107–252, title II, subtitle D, section 291, Payments for Protection and Advocacy to Assure Access for Individuals with Disabilities (42 U.S.C. 15461). The report is provided in writing to the Administration for Community Living, Administration on Intellectual and Developmental Disabilities (AIDD). Each eligible Protection and Advocacy System (P&As) must prepare and submit an annual report at the end of every fiscal year by the 31st of December. The report addresses the activities conducted with the funds provided during the year. The information collected from the annual report will be aggregated into an annual profile of how the P&As have utilized the funds and review the P&As activities carried out for each of the seven mandated area. These areas include full participation in the electoral process; education, training and assistance; advocacy and education around HAVA implementation efforts; training and education of election officials, poll workers and election volunteers regarding the rights of voters with disabilities and best practices; assistance in filing complaints; assistance to State and other governmental entities regarding the physical accessibility of polling places; and obtaining training and technical assistance on voting issues. The PAVA annual narrative report will also provide an overview of the goals and accomplishments for each P&A as well as permit the Administration on Intellectual and Developmental Disabilities (AIDD) to track voting progress to monitor grant activities and create the bi-annual report to Congress.

ACL estimates the burden of this collection of information as follows: 55 Protection and Advocacy Systems (P&A) respond annually which should be an average burden of 20 hours per State per year or a total of 1,100 hours for all states annually.

Dated: March 25, 2015.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2015–07313 Filed 3–30–15; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Treatment; Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given that the Substance Abuse and Mental Health Services Administration's (SAMHSA's) Center for Substance Abuse Treatment (CSAT) National Advisory Council will meet on April 15, 2015, from 9:30 a.m.–5:00 p.m. (EDT) and will include a session that is closed to the public.

The closed meeting will include the review of grant applications, which contain budget information, including the description of how an agency prices its services, information on proposed business relationships and subcontracts. Grant applications also contain personal information and contact information on agency principles. Discussion of proposed funding and awardees would be made public prior to the required congressional notification of grant award. Since the closed meeting will include discussion and evaluation of grant applications reviewed by Initial Review Groups and involve an examination of confidential financial and business information as well as personal information concerning the applicants, it will be closed to the public from 9:30 a.m. to 11:00 a.m. as determined by the SAMHSA Administrator, in accordance with Title 5 U.S.C. 552b(c)(4) and (6) and (c)(9)(B) and 5 U.S.C. App. 2, section 10(d).

The open session of the meeting will be held from 11:00 a.m.–5:00 p.m. and will include consideration of minutes from the SAMHSA CSAT NAC meeting of August 27, 2014, Director's report, discussion of SAMHSA's role regarding treatment of mental illness and substance use disorders, budget update, Pregnant and Postpartum Women and Medication Assisted Treatment panel discussions, and a recovery presentation and discussion.

The meeting will be held at the SAMHSA building, 1 Choke Cherry Road, Great Falls Conference Room, Rockville, MD 20850. Attendance by the public will be limited to space available and will be limited to the open sessions of the meeting. Interested persons may present data, information, or views, orally or in writing, on issues pending before the Council. Written submissions should be forwarded to the contact person on or before April 5, 2015. Oral presentations from the public will be scheduled at the conclusion of the meeting. Individuals interested in making oral presentations are encouraged to notify the contact on or before April 5, 2015. Five minutes will be allotted for each presentation.

The open meeting session may be accessed via telephone. To attend on site, obtain the call-in number and access code, submit written or brief oral comments, or request special accommodations for persons with disabilities, please register on-line at *http://nac.samhsa.gov/Registration/meetingsRegistration.aspx,* or communicate with SAMHSA's Committee Management Officer, LCDR Holly Berilla (see contact information below).

Substantive meeting information and a roster of Council members may be obtained either by accessing the SAMHSA Council Web site at: *http://www.samhsa.gov/about-us/advisory-councils/csat-national-advisory-council* or by contacting LCDR Berilla.

Substantive program information may be obtained after the meeting by accessing the SAMHSA Council Web site, *http://nac.samhsa.gov/,* or by contacting LCDR Berilla.

Council Name: SAMHSA's Center for Substance Abuse Treatment National Advisory Council.

Date/Time/Type:

April 15, 2015, 9:30 a.m.–11:00 a.m. EDT, CLOSED

April 15, 2015, 11:00 a.m.–5:00 p.m. EDT, OPEN

Place: SAMHSA Building, 1 Choke Cherry Road, Great Falls Conference Room, Rockville, Maryland 20850.

Contact: LCDR Holly Berilla, Committee Management Officer and Acting Designated Federal Official, CSAT National Advisory Council, 1 Choke Cherry Road, Rockville, Maryland 20857 (mail). Telephone: (240) 276–1252. Fax: (240) 276–2252. Email: *holly.berilla@samhsa.hhs.gov.*

Summer King,

Statistician, SAMHSA.

[FR Doc. 2015–07284 Filed 3–30–15; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0473]

Agency Information Collection Activities; Proposed Collection; Comment Request; Irradiation in the Production, Processing, and Handling of Food

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on our proposed collection of certain information. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies must publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and allow 60 days for public comment. This notice invites comments on the information collection provisions of our requirements for food irradiation processors.

DATES: Submit either electronic or written comments on the collection of information by June 1, 2015.

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

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, we invite comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of our functions, including whether the information will have practical utility; (2) the accuracy of our estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Irradiation in the Production, Processing, and Handling of Food—21 CFR Part 179 (OMB Control Number 0910–0186)—Extension

Under sections 201(s) and 409 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 321(s) and 348), food irradiation is subject to regulation under the food additive premarket approval provisions of the FD&C Act. The regulations providing for uses of irradiation in the production, processing, and handling of food are found in part 179 (21 CFR part 179). To ensure safe use of a radiation source, § 179.21(b)(1) requires that the label of sources bear appropriate and accurate information identifying the source of radiation and the maximum (or minimum and maximum) energy of the emitted radiation. Section 179.21(b)(2) requires that the label or accompanying labeling bear adequate directions for installation and use and a statement supplied by us that indicates maximum dose of radiation allowed. Section 179.26(c) requires that the label or accompanying labeling bear a logo and a radiation disclosure statement. Section 179.25(e) requires that food processors who treat food with radiation make and retain, for 1 year past the expected shelf life of the products up to a maximum of 3 years, specified records relating to the irradiation process (*e.g.,* the food treated, lot identification, scheduled process, etc.). The records required by § 179.25(e) are used by our inspectors to assess compliance with the regulation that establishes limits within which radiation may be safely used to treat food. We cannot ensure safe use without a method to assess compliance with the dose limits, and there are no practicable methods for analyzing most foods to determine whether they have been treated with ionizing radiation and are within the limitations set forth in part 179. Records inspection is the only way to determine whether firms are complying with the regulations for treatment of foods with ionizing radiation.

Description of respondents: Respondents are businesses engaged in the irradiation of food.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]

21 CFR section	Number of record-keepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeping	Total hours
179.25(e), large processors	4	300	1,200	1	1,200
179.25(e), small processors	4	30	120	1	120