• Respondent falsified data on the enrollment forms and follow-up forms for participant numbers 153 and 154 by changing their enrollment numbers.

ORI acknowledges that the Respondent was remorseful.

Ms. Robertson has entered into a Voluntary Settlement Agreement in which she has voluntarily agreed, for a period of three (3) years, beginning on October 14, 2009:

(1) To exclude herself from serving in any advisory capacity to PHS, including but not limited to service on any PHS advisory committee, board, and/or peer review committee, or as a consultant;

(2) That any institution that submits an application for PHS support for a research project on which the Respondent's participation is proposed or that uses her in any capacity on PHS-supported research, or that submits a report of PHS-funded research in which she is involved, must concurrently submit a plan for supervision of her duties to the funding agency for approval; the supervisory plan must be designed to ensure the scientific integrity of her research contribution; respondent agreed that she will not participate in any PHS-supported research until such a supervisory plan is submitted to ORI; and

(3) That any institution employing her submits, in conjunction with each application for PHS funds or report, manuscript, or abstract of PHS-funded research in which the Respondent is involved, a certification that the data provided by the Respondent are based on actual experiments or are otherwise legitimately derived and that the data, procedures, analyses, and methodology are accurately reported in the application, report, manuscript, or abstract. The Respondent must ensure that the institution sends a copy of the certification to ORI.

FOR FURTHER INFORMATION CONTACT: Director, Division of Investigative Oversight, Office of Research Integrity, 1101 Wootton Parkway, Suite 750, Rockville, MD 20852, (240) 453–8800.

John Dahlberg,

Director, Division of Investigative Oversight, Office of Research Integrity.

[FR Doc. E9–28814 Filed 12–1–09; 8:45 am]

BILLING CODE 4150–31–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0554]

Agency Information Collection Activities; Proposed Collection; Comment Request; Manufactured Food Regulatory Program Standards

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the Manufactured Food Regulatory Program Standards.

DATES: Submit written or electronic comments on the collection of information by February 1, 2010.

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: Daniel Gittleson, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–5156, *Daniel.Gittleson@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, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA'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.

Manufactured Food Regulatory Program Standards—(OMB Control Number 0910–0601—Extension)

I. Background

In the Federal Register of July 20, 2006 (71 FR 41221), FDA announced the availability of a draft document entitled ''Manufactured Food Regulatory Program Standards.'' These draft program standards are the framework that States should use to design and manage its manufactured food program. The implementation of the program standards will be negotiated as an option for payment under the State food contract. States that are awarded this option will receive up to $25,000 over a period of 5 years to fully implement the program standards.

In the first year of implementing the program standards, the State program conducts a baseline self-assessment to determine if they meet the elements of each standard. The State program should use the worksheets and forms contained herein; however it can use alternate forms that are equivalent. The State program maintains the documents and verifying records required for each standard. The information contained in the documents must be current and fit-for-use. If the State program fails to meet all program elements and documentation requirements of a standard, it develops a strategic plan to fully implement the program standards in 5 years. The strategic plan includes the following: (1) The individual element or documentation requirement

of the standard that was not met, (2) improvements needed to meet the program element or documentation requirement of the standard, and (3) projected completion dates for each task.

II. Electronic Access

Persons with access to the Internet may obtain the draft program standards at *http://www.fda.gov/downloads/RegulatoryInformation/Guidances/UCM125448.pdf*.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
44	1	44	40	1,760

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

TABLE 2.—ESTIMATED FIRST-YEAR BASELINE SELF-ASSESSMENT BURDEN[1]

No. of Respondents	Five-Year Frequency per Response	Total First-Year Responses	Hours per Response	Total Hours
17	1	17	200	3,400

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: November 23, 2009.

David Horowitz,

Assistant Commissioner for Policy.

[FR Doc. E9–28834 Filed 12–1–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[LLCAC069000 L17110000 AL0000]

Notice of Public Meeting of the Carrizo Plain National Monument Advisory Council

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice of Public Meeting.

SUMMARY: In accordance with the Federal Land Policy and Management Act and the Federal Advisory Committee Act of 1972, the U.S. Department of the Interior, Bureau of Land Management (BLM) Carrizo Plain National Monument Advisory Council (MAC) will meet as indicated below.

DATES: The meeting will be held on Saturday, January 23, 2010, at the Carissa Plain Elementary School, located approximately 2 miles northwest of Soda Lake Road on Highway 58. The meeting will begin at 10 a.m. and finish at 3 p.m. The meeting will focus on the Resource Management Plan/Environmental Impact Statement (RMP/EIS) being developed for the Carrizo Plain National Monument. There will be a public comment period from 2 p.m. to 3 p.m.

FOR FURTHER INFORMATION CONTACT: The BLM, attention: Johna Hurl, Monument Manager, 3801 Pegasus Drive, Bakersfield, CA 93308. Phone (661) 391–6093 or e-mail: *jhurl@blm.gov*.

SUPPLEMENTARY INFORMATION: The nine-member MAC advises the Secretary of the Interior, through the BLM, on a variety of public land issues associated with the public land management of the Carrizo Plain National Monument in Central California. At this meeting, monument staff will present updated information on the progress of the RMP/EIS. Draft alternatives being developed by the Carrizo managing partners—the BLM, the California Department of Fish and Game and The Nature Conservancy—will be the focus of this meeting. This meeting is open to the public. Depending on the number of persons wishing to comment, and the time available, the time allotted for individual oral comments may be limited. Individuals who plan to attend and need special assistance such as sign language interpretation or other reasonable accommodations should contact the BLM as indicated below.

Dated: November 25, 2009.

Janet Bedrosian,

Deputy State Director, External Affairs, California State Office.

[FR Doc. E9–28808 Filed 12–1–09; 8:45 am]

BILLING CODE 4310–40–P

JUDICIAL CONFERENCE OF THE UNITED STATES

Meeting of the Judicial Conference Advisory Committee on Rules of Civil Procedure

AGENCY: Judicial Conference of the United States Advisory Committee on Rules of Civil Procedure.

ACTION: Notice of open meeting.

SUMMARY: The Advisory Committee on Rules of Civil Procedure will hold a two-day meeting. The meeting will be open to public observation but not participation.

DATES: March 18–19, 2010.

Time: 8:30 a.m. to 5 p.m.

ADDRESSES: Emory Law School, 1301 Clifton Road, Atlanta, GA 30322.

FOR FURTHER INFORMATION CONTACT: John K. Rabiej, Chief, Rules Committee Support Office, Administrative Office of the United States Courts, Washington, DC 20544, telephone (202) 502–1820.

Dated: November 23, 2009.

John K. Rabiej,

Chief, Rules Committee Support Office.

[FR Doc. E9–28531 Filed 12–1–09; 8:45 am]

BILLING CODE 2210–55–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

This is notice that on July 1, 2009, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the basic classes of controlled substances listed in schedule II:

Drug	Schedule
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II