pesticide manufacturer. The following list of North American Industrial Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

B. What should I consider as I prepare my comments for EPA?

1. Submitting CBI. Do not submit this information to EPA through regulations.gov or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI information in a disk or CD-ROM that you mail to EPA, mark the outside of the disk or CD-ROM as CBI and then identify electronically within the disk or CD-ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. Tips for preparing your comments. When submitting comments, remember to:

- Identify the document by docket ID number and other identifying information (subject heading, Federal Register date and page number).
- Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.
- Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.
- Describe any assumptions and provide any technical information and/or data that you used.
- If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.
- Provide specific examples to illustrate your concerns and suggest alternatives.
- Explain your views as clearly as possible, avoiding the use of profanity or personal threats.
- Make sure to submit your comments by the comment period deadline identified.

II. Registration Applications

EPA has received applications to register new uses for pesticide products containing currently registered active ingredients. Pursuant to the provisions of FIFRA section 3(c)(4), EPA is hereby providing notice of receipt and opportunity to comment on these applications. Notice of receipt of these applications does not imply a decision by the Agency on these applications. For actions being evaluated under the Agency’s public participation process for registration actions, there will be an additional opportunity for a 30-day public comment period on the proposed decision. Please see the Agency’s public participation Web site for additional information on this process (http://www.epa.gov/pesticides/regulating/registration-public-involvement.html). EPA received the following applications to register new uses for pesticide products containing currently registered active ingredients:

1. EPA Registration Numbers: 100–889 and 100–963. Docket ID number: EPA–HQ–OPP–2013–0268. Applicant: Syngenta Crop Protection, LLC, P.O. Box 18300, Greensboro, NC 27419–8300. Active ingredient: Thiabendazole, 2-(4-thiazolyl) benzimidazole. Product type: Fungicide. Proposed uses: Vegetable, root (except sugar beet), subgroup 1B; onion, bulb, subgroup 3–07A; Brassica, head and stem, subgroup 5–A; vegetable, cucurbit, group 9; alfalfa, spinach (including baby); and small grain cereals (wheat, barley, oats, rye, and triticale).


List of Subjects

Environmental protection, Pesticides and pest.

Dated: August 1, 2013.

Daniel J. Rosenblatt,
Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2013–19358 Filed 8–8–13; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Sunshine Act Meeting

Pursuant to the provisions of the “Government in the Sunshine Act” (5 U.S.C. 552b), notice is hereby given that at 10:00 a.m. on Wednesday, August 7, 2013, the Board of Directors of the Federal Deposit Insurance Corporation met in closed session to consider matters related to the Corporation’s supervision, corporate, and resolution activities.

In calling the meeting, the Board determined, on motion of Vice Chairman Thomas M. Hoenig, seconded by Director Jeremiah O. Norton (Appointive), concurred in by Director Thomas J. Curry (Comptroller of the Currency), Director Richard Cordray (Director, Consumer Financial Protection Bureau), and Chairman Martin J. Gruenberg, that Corporation business required its consideration of the matters which were to be the subject of this meeting on less than seven days’ notice to the public; that no earlier notice of the meeting was practicable; that the public interest did not require consideration of the matters in a meeting open to public observation; and that the matters could be considered in a closed meeting by authority of subsections (c)(4), (c)(6), (c)(8), (c)(9)(A)(ii), (c)(9)(B), and (c)(10) of the...
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–19606–30D]

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, will submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for renewal of the approved information collection assigned OMB control number 0990–0221, scheduled to expire on January 31, 2014. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Deadline: Comments on the ICR must be received on or before September 9, 2013.

ADDRESS: Submit your comments to OIRA_submission@omb.eop.gov or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance Staff, Information.CollectionClearance@hhs.gov or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the OMB control number 0990–0221 and document identifier HHS–OS–19606–30D for reference.

Information Collection Request Title: Family Planning Annual Report: Forms and Instructions.

OMB No.: 0990–0221.

Abstract: The Office of Population Affairs (OPA), Office of the Assistant Secretary for Health (OASH), U.S. Department of Health and Human Services (HHS) administers and oversees the Title X Family Planning Program. The Family Planning Annual Report (FPAR) is an annual reporting requirement for family planning services delivery projects ("Title X service grantees") authorized and funded by the Title X Family Planning Program ["Population Research and Voluntary Family Planning Programs" (Pub. L. 91–572), which was enacted in 1970 as Title X of the Public Health Service Act (Section 1001 of Title X of the Public Health Service Act, 42 United States Code 300C). The Title X Family Planning Program is the only Federal grant program dedicated solely to providing individuals with comprehensive family planning and related preventive health services.

The FPAR, the only source of annual, uniform reporting by all Title X service grantees, provides consistent, national-, regional-, state-, and grantee-level data on the services provided and the characteristics of the individuals served. OPA uses FPAR data to monitor compliance with statutory requirements and accountability and federal performance requirements for Title X family planning funds as required by the 1993 Government Performance and Results Act (GPRA) and HHS, to guide financial and program planning and evaluation, and to respond to inquiries about the program from policymakers and Congress. Note that there are no changes to the FPAR except minor corrections or clarifications to submission and reporting instructions or definitions. The estimated average hour burden has been reduced to 36 hours, which is 4 hours lower than the 40-hour estimate of the previous OMB submission.

Need and Proposed Use of the Information: The program’s purpose is to assist individuals in determining the number and spacing of their children. It is designed to provide access to contraceptive services, supplies, and information to all who want and need them. By law, priority is given to persons from low-income families