Competition Bureau, Federal Communications Commission, The Portals II, 445 12th Street, SW., Suite 5–A420, Washington, DC 20554. Requests to make an oral statement or provide written comments to the NANC should be sent to Deborah Blue.

FOR FURTHER INFORMATION CONTACT: Deborah Blue, Special Assistant to the Designated Federal Officer (DFO) at (202) 418–1466 or *dblue@fcc.gov.* The fax number is: (202) 418–2345. The TTY number is: (202) 418–0484.

SUPPLEMENTARY INFORMATION: Released: June 24, 2003.

The North American Numbering Council (NANC) has scheduled a meeting to be held Tuesday, July 15, 2003, from 9 a.m. until 5 p.m. The meeting will be held at the Federal Communications Commission, Portals II, 445 12th Street, SW., Room TW–C305, Washington, DC. This meeting is open to members of the general public. The FCC will attempt to accommodate as many participants as possible. The public may submit written statements to the NANC, which must be received two business days before the meeting. In addition, oral statements at the meeting by parties or entities not represented on the NANC will be permitted to the extent time permits. Such statements will be limited to five minutes in length by any one party or entity, and requests to make an oral statement must be received two business days before the meeting.

Proposed Agenda—Tuesday, July 15, 2003, 9 AM

1. Announcements and Recent News
2. Approval of Minutes
   —Meeting of May 13, 2003
3. Report of the North American Numbering Plan Administrator (NANPA)
4. Report of National Thousands Block Pooling Administrator
   —Activity report
5. Report of 3-Digit DIG IMG
6. Report of Numbering Oversight Working Group
7. Status of Industry Numbering Committee activities
8. Report of the Local Number Portability Administration (LNPA) Working Group
   —Wireless Number Portability Operations (WNPO) Subcommittee
9. Report from NBANC
10. Report of Cost Recovery Working Group
11. Summary of Action Items
12. Steering Committee
    —Table of NANC Projects
13. Public Comments and Participation (5 minutes per speaker)
14. Other Business

Adjourn no later than 5 p.m.

Federal Communications Commission

Cheryl L. Callahan,

Assistant Chief, Telecommunications Access Policy Division, Wireline Competition Bureau.

[FR Doc. 03–16411 Filed 6–27–03; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: Applied Research on Antimicrobial Resistance Prevention, Program Announcement #03120

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Applied Research on Antimicrobial Resistance, Program Announcement #03120.

Times and Dates: 9 a.m.–10 a.m., July 17, 2003 (Open), 10 a.m.–6 p.m., July 17, 2003 (Closed).

Place: National Center for Infectious Disease, Office of the Director, OAS Conference Room #3555, 35 Executive Park Drive, Atlanta, GA 30342, Telephone 404.498.3240.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement #03120.

Contact Person for More Information: Mary L. Lerchen, Deputy Director, Office of Extramural Research, National Center for Infectious Diseases, CDC, 1600 Clifton Road, MS–C19, Atlanta, GA 30333, Telephone 404.639.0043.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: June 20, 2003.

Alvin Hall,

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

[FR Doc. 03–16407 Filed 6–27–03; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0273]

Agency Information Collection Activities; Proposed Collection; Comment Request; Research Study Complaint Form

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's burden estimates to use an Internet-based complaint form for public complaints concerning misconduct in research studies.

DATES: Submit written or electronic comments on the collection of information by August 29, 2003.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.fda.gov/dockets/ecomments.* 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: Karen Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

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

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: (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.

Currently, FDA's Center for Drug Evaluation and Research, Division of Scientific Investigations (DSI), receives an average of about 150 unsolicited complaints per year about scientific misconduct in clinical research through electronic mail, regular mail, telephone, and personal contacts. DSI will continue to receive and process such complaints. The internet-based complaint form for consumer complaints on research studies will provide an additional convenient and efficient way for the public to submit complaints regarding misconduct in clinical research. The complaint form asks questions about the individual, company, or organization that is the subject of the complaint; the event and the drug product(s) that prompted the complaint; and optional information about the person submitting the complaint. The complaint form is accessible at *http://didit.devis.com/complaints*. The username is ''public'' and the password is ''fdapublic.''

FDA will use the information collected through the complaint form to identify weaknesses in the current services provided to human subjects in clinical research and to improve and maintain a high quality of service to the affected public. The complaint form will be encrypted so that any information of a sensitive nature will not be unnecessarily or prematurely disclosed. The complainants will remain anonymous unless they voluntarily disclose their identity. Participation is fully voluntary and complainants will be able to complete, review, edit, and submit the form to FDA. DSI will acknowledge the receipt of each complaint.

DSI will complete initial analyses of the information from each complaint within 10 working days. Each complaint will be reviewed by a responsible person in DSI and then distributed to the appropriate unit in DSI or FDA for further action. DSI will contact the complainant if the complainant requests a followup contact. If the complainant does not request any followup contact, then no additional contact with the complainant is anticipated.

FDA estimates that approximately 144 persons will voluntarily complete the complaint form each year. The estimated time for completing each complaint form will be 1 hour, resulting in a total burden of 144 hours per year (144 complainants x 1 hour = 144 burden hours per year). The burden of this collection of information is estimated as follows:

TABLE 1.—ESTIMATED ONE-TIME REPORTING BURDEN[1]

No. of Respondents	Annual Frequency per Response	Total Annual Records	Hours per Response	Total Hours
144	1	144	1	144

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

Dated: June 24, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–16494 Filed 6–27–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0268]

Agency Information Collection Activities; Proposed Collection; Comment Request; Biological Products: Reporting of Biological Product Deviations in Manufacturing

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 information collection requirements relating to the reporting of biological product deviations in manufacturing.

DATES: Submit written or electronic comments on the collection of information by August 29, 2003.

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

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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: (1) Whether the proposed