activities to collect and analyze optional data consistent with the Program Requirements for Part B, including sampling methods and proposed staffing.

3. Budget and Justification: (Not Scored)

The extent to which the budget is reasonable, clearly justified, and consistent with stated objectives and proposed activities.

H. Other Requirements

Technical Reporting Requirements

Provide CDC with original plus 2 copies of:

1. Semi-annual progress reports.
2. Financial status report, no more than 90 days after the end of the budget period.
3. Final financial and performance reports, no more than 90 days after the end of the project period.

Send all reports to the Grants Management Specialist identified in the ''Where to Obtain Additional Information'' section of this announcement.

For descriptions of the following Other Requirements, see Attachment I in the application package:

AR–7—Executive Order 12372 Review
AR–9—Paperwork Reduction Act Requirements
AR–10—Smoke-Free Workplace Requirements
AR–11—Healthy People 2010
AR–12—Lobbying Restrictions
AR–13—Prohibition on Use of CDC Funds for Certain Gun Control Activities
AR–21—Small, Minority, and Women-Owned Businesses

I. Authority and Catalog of Federal Domestic Assistance Number

This program is authorized under sections 301(a), 317(k)(2), 391, 392, 393A, 394, and 394A [42 U.S.C. 241(a), 247b(k)(2), 280b, 280b–1, 280b–2, 280b–3] of the Public Health Service Act, as amended. The Catalog of Federal Domestic Assistance number is 93.136.

J. Where To Obtain Additional Information

This and other CDC announcements are available through the CDC home page on the Internet at: http://www.cdc.gov. To receive additional written information and to request an application kit, call 1–888–GRANTS4 (1–888 472–6874). You will be asked to leave your name and address, and will be instructed to identify the announcement number of interest.

If you have questions after reviewing the contents of all the documents, business management assistance may be obtained from: Angie Nation, Grants Management Specialist, Announcement #01030, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention, 2920 Brandywine Road, Suite 3000, Atlanta, GA 30341–4146, Telephone number (770) 488–2719, Email address: aen4@cdc.gov.

For program technical assistance, contact: Renee Johnson, MSPH, CDC National Center for Injury Prevention and Control, 4770 Buford Highway, NE, Mailstop F41, Atlanta, GA 30341–3724, Telephone (770) 488–4031, Email address: nba7@cdc.gov.

For a copy of the CDC Guidelines for Central Nervous System Injury Surveillance, contact: Patricia Allen, CDC National Center for Injury Prevention and Control, 4770 Buford Highway, NE, Mailstop F41, Atlanta, GA 30341–3724, Telephone (770) 488–4031, Email address: pca9@cdc.gov.

CDC does not guarantee to accept or justify its nonacceptance of recommendations that are received more than 60 days after the application deadline.

Dated: February 23, 2001.

John L. Williams,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 01–4937 Filed 2–28–01; 8:45 am]

BILLING CODE 4163–18–P

ANNUAL BURDEN ESTIMATES

Instrument	Number of Respondents	Number of Responses per Respondent	Average Burden Hours per Response	Total Burden Hours
18-State Telephone Survey	18	1	3	54
54-State Mail Questionaire	54	1	6	324
Open-ended interview for Case Studies	54	1	1.5	81
Follow-up Telephone Interview with Applicants and Non-applicants	1200	1	.33	396
Case abstractions—pulling case files for contractor review and abstraction	6	1	20	120
Estimated Total Annual Burden Hours:				975

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Study of the TANF Application Process.

OMB No. New Collection.

Description: The Study of the TANF Application Process is designed to provide systematic information about how application policies and processes have changed under TANF, and how States define and count applications and application results. The Study will also explore how application polices are implemented in a sample of local TANF offices and will collect data on individuals' application decisions, experiences, and outcomes. In addition, the Study will also collect information on the availability and quality of State-collected data on the TANF application process. The primary purpose of this Study is to provide useful information to be considered in the upcoming TANF reauthorization process.

Respondents: The respondents for the Mail Questionnaire are the 50 States, the District of Columbia, and the U.S. Territories of Guam, Puerto Rico, and the Virgin Islands. Eighteen States will be respondents to the State Telephone Survey, 54 individuals for the Open-ended Interviews for Case Studies, six States for Case Abstractions, and 1200 individuals for the follow-up Telephone Interviews with Applicants and Non-applicants.

In compliance with the requirements of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF.

Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: February 26, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–5009 Filed 2–28–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–1435]

Agency Information Collection Activities; Announcement of OMB Approval; Substantial Evidence of Effectiveness of New Animal Drugs

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Substantial Evidence of Effectiveness of New Animal Drugs'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 16, 2000 (65 FR 49989), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0356. The approval expires on February 29, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: February 22, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01–4961 Filed 2–28–01; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 95N–0220]

Agency Information Collection Activities; Announcement of OMB Approval; Substances Approved for Use in the Preparation of Meat and Poultry Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Substances Approved for Use in the Preparation of Meat and Poultry Products'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 25, 2000 (65 FR 51758), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0461. The approval expires on February 29, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: February 22, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01–4965 Filed 2–28–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0242]

Agency Information Collection Activities; Announcement of OMB Approval; 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 that a collection of information entitled ''Biological Products: Reporting of Biological Product Deviations in Manufacturing'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

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

SUPPLEMENTARY INFORMATION: In the **Federal Register** of November 7, 2000 (65 FR 66621), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0458. The approval expires on February 29, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.