of 2008," title IV–E was expanded, effective with FY 2009, to include a third program, Guardianship Assistance, and was further expanded, effective with FY 2010, to include Tribes, tribal organizations and consortia as additional grantees.

Ultimately, the combined effect of these changes will be to significantly increase the number of grantees, the number of grant awards and the required amount of financial reporting. In recognition of these substantial program revisions and to accommodate these changes, the quarterly financial report has been revised, redesigned and re-designated as Form CB–496, the "Title IV–E Programs Quarterly Financial Report."

The Administration for Children and Families (ACF) provides Federal funding at the rate of 50 percent for most administrative and other related costs and at enhanced rates ranging from 55 to 75 percent for training costs as detailed in Federal statute and regulations. This form is submitted quarterly by each State and Tribe to estimate the funding needs for the upcoming fiscal quarter and to report expenditures for the fiscal quarter just ended. The information collected in this report is used by this agency to calculate quarterly Federal grant awards and to enable oversight of the financial management of the programs.

Comments concerning these revisions were received from both Federal and grantee staffs by the ACF Office of Grants Management, both directly and in response to an earlier **Federal Register** Notice (74 FR 22749, May 14, 2009) that provided many useful recommendations and suggestions, many of which were incorporated into the final draft product.

Respondents: State and Tribal title IV–E agencies administering the Foster Care, Adoption Assistance and Guardianship Assistance Programs.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Form CB–496	62	4	17	4,216

Estimated Total Annual Burden Hours: 4,216

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Dated: May 12, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–11814 Filed 5–17–10; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Application for the Pharmacology Research Associate Program

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Institute of General Medical Sciences (NIGMS), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: Application for the Pharmacology Research Associate Program. *Type of Information Collection Request:* Extension of a currently approved collection, OMB No. 0925–0378, expiration date December 31, 2010. *Form Numbers:* NIH 2721–1, NIH 2721–2. *Need and Use of Information Collection:* The Pharmacology Research Associate (PRAT) Program will use the applicant and referee information to award opportunities for training and experience in laboratory or clinical investigation to individuals with a Ph.D. degree in pharmacology or a related science, M.D., or other professional degree through appointments as PRAT Fellows at the National Institutes of Health or the Food and Drug Administration. The goal of the program is to develop leaders in pharmacological research for key positions in academic, industrial, and Federal research laboratories. *Frequency of Response:* Once a year. *Affected Public:* Individuals or households; Businesses or other for-profit. *Type of Respondents:* Applicants and Referees.

The annual reporting burden is as follows:

Type and number of respondents	Estimated number of responses per respondent	Estimated total responses	Average burden hours per responses	Estimated total annual burden hours requested
Applicants, 25	1	25	8.00	200
Referees, 75	1	75	1.75	131.25

Total Number of Respondents: 100.
Total Number of Responses: 100.
Total Hours: 331.25.

The annualized cost to respondents is estimated at:

Applicants: $10,250.00.

Referees: $6,562.50.

There are no Capital Costs, Operating costs, and/or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Ms. Liz Elliott, NIGMS, NIH, Natcher Building, Room 2AN–18H, 45 Center Drive, MSC 6200, Bethesda, MD 20892–6200, or call non-toll-free number 301–594–2755 or e-mail your request, including your address to: *elliotte@nigms.nih.gov*.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: May 10, 2010.

Sally Lee,

Executive Officer, National Institute of General Medical Sciences, National Institutes of Health.

[FR Doc. 2010–11857 Filed 5–17–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0057]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Submit Information in Electronic Format to the Center for Veterinary Medicine Using the FDA Electronic Submission Gateway

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by June 17, 2010.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0454. Also include the FDA docket number found in brackets in the heading of this document

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Guidance for Industry on How to Submit Information in Electronic Format to the Center for Veterinary Medicine Using the FDA Electronic Submission Gateway (OMB Control Number 0910–0454)—Extension

The Center for Veterinary Medicine (CVM), accepts certain types of submissions electronically with no requirement for a paper copy. These types of documents are listed in public docket 97S–0251 as required by 21 CFR 11.2. CVM's ability to receive and process information submitted electronically is limited by its current information technology capabilities and the requirements of the Electronic Records; Electronic Signatures final regulation. CVM's guidance entitled "Guidance for Industry: How to Submit Information in Electronic Format to CVM Using the FDA Electronic Submission Gateway" outlines general standards to be used for the submission of any information by e-mail. The likely respondents are sponsors for new animal drug applications.

In the **Federal Register** of February 5, 2010 (75 FR 6038), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section/Form 3538	No. of Respondents	Annual Frequency per Response	Total Annual Responses[2]	Hours per Respondent	Total Hours
11.2	40	1.3	52	.08	4.2

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] Electronic submissions received between January 1, 2008, and December 31, 2008.

The number of respondents in table 1 of this document is the number of sponsors registered to make electronic submissions (40). The number of total annual responses is based on a review of the actual number of such submissions made between January 1, 2008, and December 31, 2008 (52 x hours per response (.08) = 4.2 total hours).

Dated: May 12, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–11808 Filed 5–17–10; 8:45 am]

BILLING CODE 4160–01–S