Dated: November 14, 2002.

Regina B. Schofield,

Director, Office of Intergovernmental Affairs.

[FR Doc. 02–29491 Filed 11–15–02; 2:06 pm]

BILLING CODE 4165–15–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Native Employment Works (NEW) Program Plan Guidance and Program Report.

OMB No.: 0970–0174.

Description: The Native Employment Works (NEW) program plan is the application for NEW program funding. As approved by the Department of Health and Human Services (HHS), it documents how the grantee will carry out its NEW program. The NEW program plan guidance specifies the information needed to complete a NEW program plan and explains the process for plan submission every third year.

The NEW program report provides information on the activities and accomplishments of grantees' NEW programs. The NEW program report and instructions specify the program data that NEW grantees report annually.

Respondents: Federally-recognized Indian tribes and tribal organizations that are NEW program grantees:

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours (annually)
NEW program plan guidance	26	One, every 3 years	30	780
New program report	53	One annually	15	795

Estimated Total Annual Burden Hours: 1575.

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. Copes 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 Administration, 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 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: November 12, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–29225 Filed 11–18–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Refugee Unaccompanied Minor Placement Report, Refugee Unaccompanied Minor Progress Report.

OMB No.: 0970–0034.

Description: The two reports collect information necessary to administer the refugee unaccompanied minor program. The ORR–3 (Placement Report) is submitted to ORR by the service provider agency at initial placement and whenever there is a change in the child's status, including termination from the program. The ORR–4 is submitted annually and records the child's progress toward the goals listed in the child's case plan.

Respondents. State governments.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ORR–3	12	15	.417	75
ORR–4	12	60	.250	180

Estimated Total Annual Burden Hours: 255.

In compliance with the requirements of section 3506(c)(21)(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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20477, 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: November 11, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–29226 Filed 11–18–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0319]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Blood Establishment Registration and Product Listing, Form FDA 2830

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the 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: Submit written comments on the collection of information by December 19, 2002.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Stuart Shapiro, Desk Officer for FDA.

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

Establishment Registration and Product Listing, Form FDA 2830—21 CFR Part 607—(OMB Control Number 0910–0052)—Extension

Under section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360), any person owning or operating an establishment that manufactures, prepares, propagates, compounds, or processes a drug or device must register with the Secretary of Health and Human Services, on or before December 31 of each year, his or her name, place of business and all such establishments, and submit, among other information, a listing of all drug or device products manufactured, prepared, propagated, compounded, or processed by him or her for commercial distribution. In part 607 (21 CFR part 607), FDA has issued regulations implementing these requirements for manufacturers of human and products. Section 607.20(a) requires certain establishments that engage in the manufacture of products to register and to submit a list of products in commercial distribution. Section 607.21 requires the establishments entering into the manufacturing of products to register within 5 days after beginning such operation and to submit a product listing at that time. In addition, establishments are required to register annually between November 15 and December 31 and update their product listing every June and December. Section 607.22 requires the use of Form FDA 2830 for registration and product listing. Section 607.25 indicates the information required for establishment registration and product listing. Section 607.26 requires certain changes to be submitted as an amendment to the establishment registration within 5 days of such changes. Section 607.30 requires establishments to update, as needed, their product listing information every June and at the annual registration. Section 607.31 requires that additional product listing information be provided upon FDA request. Section 607.40 requires foreign product establishments to register and submit the product listing information, the name and address of the establishment, and the name of the individual responsible for submitting product listing information. Among other uses, this information assists FDA in its inspections of facilities, and its collection is essential to the overall regulatory scheme designed to ensure the safety of the nation's supply. Form FDA 2830, Establishment Registration and Product Listing, is used to collect this information. The likely respondents are banks, collection facilities, and component manufacturing facilities. FDA estimates the burden of this collection of information based upon the database and past experience of the Center for Biologics Evaluation and Research, Division of Applications in regulatory establishment registration and product listing. Most banks are familiar with the regulations and registration requirements to fill out this form.

In the **Federal Register** of August 2, 2002 (67 FR 50445), FDA published a 60-day notice requesting public comment on the information collection provisions. One comment was received. The comment agrees that the information collection is necessary and the Form FDA 2830 is helpful with the registration process.

The comment stated that we underestimated the hours per response regarding the initial registration and product listing update. The comment stated that it might take up to 2 hours to complete the initial registration and 0.5 hours to complete the product listing update. We decline to change the estimates based on our review of the activities associated with completing the form. Although it may take some establishments longer to complete the form, others may complete the form more quickly. Since the reporting burden includes an estimated average of the time to complete the various activities associated with the form, we believe that the current burden estimates accurately reflect the range of time to complete the form.

The comment also requested that the annual registration process be automated so that each facility could electronically submit the form, if they desire to do so, and also requested that we continue to send a hard copy of the form and instructions as a reminder to registrants to re-register. We are currently in the process of setting up a program for electronic registration. Use of the electronic system will be voluntary. We intend to continue sending a hard copy of the form and instructions for the foreseeable future.