The Secretary's Advisory Committee on National Health Promotion and Disease Prevention Objectives for 2020 will address efforts to develop the nation's health promotion and disease prevention objectives and strategies to improve the health status and reduce health risks for Americans by the year 2020. The Committee will provide to the Secretary of Health and Human Services advice and consultation for developing and implementing the next iteration of national health promotion and disease prevention goals and objectives and provide recommendations for initiatives to occur during the initial implementation phase of the goals and objectives. HHS will use the recommendations to inform the development of the national health promotion and disease prevention objectives for 2020 and the process for implementing the objectives. The intent is to develop and launch objectives designed to improve the health status and reduce health risks for Americans by the year 2020.

DATES: The Committee will meet on July 10, 2009 from 2 p.m. to 4 p.m. Eastern Daylight Time (EDT).

ADDRESSES: The meeting will be held online, via WebEx software. For detailed instructions about how to make sure that your windows computer and browser is set up for WebEx, please visit the ''Secretary's Advisory Committee'' Web page of the Healthy People Web site at: *http://www.healthypeople.gov/hp2020/advisory/default.asp*

FOR FURTHER INFORMATION CONTACT: Emmeline Ochiai, Designated Federal Officer, Secretary's Advisory Committee on National Health Promotion and Disease Prevention Objectives for 2020, U.S. Department of Health and Human Services, Office of Public Health and Science, Office of Disease Prevention and Health Promotion, 1101 Wootton Parkway, Room LL–100, Rockville, MD 20852, (240) 453–8259 (telephone), (240) 453–8281 (fax). Additional information is available on the Internet at *http://www.healthypeople.gov.*

SUPPLEMENTARY INFORMATION:

Purpose of Meeting: Every 10 years, through the Healthy People initiative, HHS leverages scientific insights and lessons from the past decade, along with the new knowledge of current data, trends, and innovations to develop the next iteration of national health promotion and disease prevention objectives. Healthy People provides science-based, 10-year national objectives for promoting health and preventing disease. Since 1979, Healthy People has set and monitored national health objectives to meet a broad range of health needs, encourage collaborations across sectors, guide individuals toward making informed health decisions, and measure the impact of our prevention and health promotion activities. Healthy People 2020 will reflect assessments of major risks to health and wellness, changing public health priorities, and emerging issues related to our nation's health preparedness and prevention.

Public Participation at Meeting: Members of the public are invited to listen to the online Committee meeting. There will be no opportunity for oral public comments during the online Committee meeting. Written comments, however, are welcome throughout the development process of the national health promotion and disease prevention objectives for 2020. They can be submitted through the Healthy People Web site at: *http://www.healthypeople.gov/hp2020/comments/* or they can be e-mailed to *HP2020@hhs.gov.* Please note that the public comment Web site will be updated throughout the Healthy People development process, so people should return to the site frequently and provide their input.

To listen to the Committee meeting, individuals must pre-register to attend at the Healthy People Web site located at *http://www.healthypeople.gov.* Participation in the meeting is limited. Registrations will be accepted until maximum WebEx capacity is reached and must be completed by 9 a.m. EDT on July 10, 2009. A waiting list will be maintained should registrations exceed WebEx capacity. Individuals on the waiting list will be contacted as additional space for the meeting becomes available.

Registration questions may be directed to Hilary Scherer at *HP2020@norc.org* (e-mail), (301) 634–9374 (phone) or (301) 634–9301 (fax).

Dated: June 5, 2009.

Penelope Slade-Sawyer,

RADM, USPHS, Deputy Assistant Secretary for Health (Disease Prevention and Health Promotion).

[FR Doc. E9–13635 Filed 6–10–09; 8:45 am]

BILLING CODE 4150–32–P

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 Report Requirements.

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 provides instructions for preparing 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
NEW program plan guidance	26	1	29	754
NEW program report	48	1	15	720

Estimated Total Annual Burden Hours: 1,474.

In compliance with the requirements of Section 506(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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, *Attn:* ACF Reports Clearance Officer. *E-mail address: infocollection@acf.hhs.gov.* 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: June 8, 2009.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E9–13684 Filed 6–10–09; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0572]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Implementation of the Animal Generic Drug User Fee Act of 2008; User Fee Cover Sheet Form FDA 3728

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 (the PRA).

DATES: Fax written comments on the collection of information by July 13, 2009.

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 0910–0632. 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 (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 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.

Implementation of the Animal Generic Drug User Fee Act of 2008; User Fee Cover Sheet Form FDA 3728—21 U.S.C. 379j-21 (OMB Control Number 0910–0632)—Extension

This collection of information is currently approved under the emergency processing provisions of the PRA for 180 days. FDA is now seeking a 3-year clearance.

Section 741 of the act (21 U.S.C. 379j-21), establishes three different kinds of user fees: (1) Fees for certain types of abbreviated applications for generic new animal drugs, (2) annual fees for certain generic new animal drug products, and (3) annual fees for certain sponsors of abbreviated applications for generic new animal drugs and/or investigational submissions for generic new animal drugs. Because the submission of user fees concurrently with applications is required, the review of an application cannot begin until the fee is submitted. Form FDA 3728, the Animal Generic Drug User Fee Cover Sheet, is designed to provide the minimum necessary information in order to do the following: (1) Determine whether a fee is required for review of an application, (2) determine the amount of fee required, and (3) account for and track user fees.

In the **Federal Register** of March 11, 2009 (74 FR 10596), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 U.S.C. 379j-21	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Form FDA 3728	20	2	40	.08	3.2

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

Respondents to this collection of information are generic animal drug applicants. Based on FDA's database system, there are an estimated 20 sponsors of new animal drugs potentially subject to the Animal Generic Drug User Fee Act of 2008.

Dated: June 3, 2009.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E9–13716 Filed 6–10–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Performance Progress Report.

OMB No.: 0970–0334.